

www.pan-biotech.de www.pan-biotech.com Dear valued customer

We think that the current biotech industry favors experienced, mid-sized and flexible producers with a broad product portfolio, strong research and development capabilities, best-in-class quality, strong business partner relationships and a wide geographic reach. During the last 26 years PAN-Biotech demonstrated its strength with innovative, first-class quality products and excellent services for cell culture around the globe.

For you, as users and customers of cell culture products, the supply situation has changed rapidly in first quarter of 2013. Due to several mergers and acquisitions of bigger as well as smaller companies there were frictions and changes in supply chains. The customer problems and frustrations were highlighted with the question(s):

Can you deliver / support us / rescue our production...? YES, WE CAN!

We can supply FBS, media, reagents and all other products presented in this catalogue We can support you

We can develop / optimise your media and much more

Our market prediction for 2013 was more frictions and turbulences. Some of the recent acquisitions went through the same route: logistic changes, lack of customer responsiveness, delivery problems, price increases, intransparency and so on. The overall results for customers was a limited choice of vendors and huge international conglomerates with reduced customer service.

PAN-Biotech still remains as one of a few privately owned, ownership managed companies. We value our customers, we value their needs and requirements, we value product quality, we value openness and transparency, we value service and commitments, we value trust and partnership. We followed along these values for the last 26 years and we intend to keep these values for the next 26 years.

We thank all our existing customers, our dedicated and highly motivated staff and our suppliers and industry partners who made possible

26 years biotech made in Germany by PAN-Biotech

And we cordially welcome all our new customers, OEM partners and interested international distributors as well as new business partners. We guarantee you the values described above and our personal commitment for excellent products and services.

Yours sincerely

Christian Niewolik & Jens Hartmann Owners & Managing Directors

March, 2014

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The function of serum in cell cultures

- Stimulates cell growth, proliferation and differentiation through hormonal factors
- Adhesion factors facilitate and enhance cell attachment on culture dishes (bio-matrix)
- Transport and binding proteins provide hormones, minerals and lipids
- Inhibition of toxic substances by binding to serum proteins

Animal serum

The dose of serum added to a cell culture as nutrient source depends on factors such as cell type, primary culture or cell line, adherent or suspension culture, and usually is in the range of 5 % to 15 % of the total liquid volume and most times used at 10 %.

Serum is produced from animal blood and fetal bovine serum is the most widely used because it contains an especially high amount of growth factors due to its origin – the blood of fetuses is a by-product of slaughtered cattle.

Advantages of PAN-Biotech

- Own raw material resources in different countries: Australia, South Africa, South America (Brazil) and the United States of America (USA)
- Certificate of Suitability (COS) no. R1-CEP 2002-167-Rev 00
- Licensed according to the EU-decree no. 1774/2002 with vet. no. DE 09 275 0001 14
- Every single batch is fully documented from the country of origin to the end product
- Every process, from collection of raw serum to production, is specified in standard operating instructions (SOP) which will be provided upon request
- We offer special types of serum: charcoal absorbed, delipidized, dialyzed, gamma irradiated, heat inactivated and gamma globulin reduced
- Highest production and safety standards for serum manufacturing
- Best references from industry and research
- Extensive analyses and tests are presented in Certificate of Analysis (CoA)

We exclusively use serum from guaranteed BSE-free collection areas. In addition, we can also confirm that South Africa, as a country of origin, is free from scrapie. No serum batches from Great Britain as the country of origin are processed by PAN-Biotech. We warrant the submission of a complete documentation, consisting of a certificate of origin and a veterinary certificate, shipping documents and a certificate of analysis. Furthermore, every single procedure during an individual production process is documented and then summarized in a production protocol.

Certificate of Suitability

Declaration

The manufacturing process and quality control testing are performed in accordance with the submitted records and with a suitable quality assurance system in compliance with ISO 9001 quality standards. This quality assurance system verifies traceability and batch consistency. PAN-Biotech conducts internal and external audits for its quality system on an annual basis. In addition, PAN-Biotech audits its raw material serum suppliers on a cyclical basis and reviews the facilities, manufacturing processes and documentation for the collection, handling, storage and transport of raw serum. PAN-Biotech is willing to be inspected, in accordance with the relevant legislation, on request of a relevant authority before and/or after being granted a certificate of suitability.

Production site:

PAN-Biotech GmbH Am Gewerbepark 13 94501 Aidenbach / GERMANY

Serum Hotline: +49(0)8543/6016-55









Certification of Substances Division

Certificate of suitability No. R1-CEP 2002-167-Rev 00

- Name of the substance: **FOETAL BOVINE SERUM**
- Name of holder:
 - PAN BIOTECH GMBH
- Am Gewerbepark 13
 - Germany-94501 Aidenbach
- Site(s) of production:
- PAN BIOTECH GMBH
- Am Gewerbepark 13
- Germany-94501 Aidenbach 10
- 11 THIS CERTIFICATE SUPERSEDES THE PREVIOUS CERTIFICATE 12 R0-CEP 2002-167-REV 01
- After examination of the information provided on the origin of raw material(s) and type of
- tissue(s) used and on the manufacturing process for this substance on the site(s) of 14
- production mentioned above, we certify that the substance FOETAL BOVINE SERUM 15
- meets the criteria described in the current version of the monograph Products with risk
- of transmitting agents of animal spongiform encephalopathies no. 1483 of the European 17 Pharmacopoeia, current edition including supplements. 18
- country(ies) of origin of source materials:
- Australia and Brazil
- nature of animal tissues used in manufacture: 20
- Foetal bovine blood
- The submitted dossier must be updated after any significant change that may after the 21
- quality, safety or efficacy of the substance, or that may alter the risk of transmitting 22
- 23 animal spongiform encephalopathy agents.
- 24 Manufacture of the substance shall take place in accordance with a suitable quality
- assurance system such as ISO 9001, and in accordance with the dossier submitted. 25
- Failure to comply with these provisions will render this certificate void. 26
- The certificate is valid provided there has been no deterioration in the TSE status of the 27
- country(ies) of origin of the source material.

Address: 7, allée Kastner, CS 30026 - F - 67081 Strasbourg (France) Telephone: 33 (0) 3 88 41 30 30 - Fax: 33 (0) 3 88 41 27 71 - e-mail: cep@edqm.eu Internet : http://www.edqm.eu



Bovine Serum

Bovine serum is the blood fraction remaining after the natural coagulation of blood, followed by centrifugation to remove any remaining red blood cells. The production of bovine serum at PAN-Biotech is tightly controlled, from the collection of serum at the slaughterhouse and throughout the whole production cycle which is performed without exception in our own production facilities in Aidenbach, Germany. All serum lots are virus and mycoplasma tested.

Fetal Bovine Serum, Australia origin	100 ml 500 ml	P30-1301 P30-1302
Fetal Bovine Serum, South Africa origin	100 ml 500 ml	P30-1505 P30-1506
Fetal Bovine Serum, South America origin	100 ml 500 ml	P30-3305 P30-3306
Fetal Bovine Serum, US admissible	100 ml 500 ml	P30-1701 P30-1702
Fetal Bovine Serum, US origin	100 ml 500 ml	P30-1401 P30-1402
Fetal Bovine Serum Premium, South Africa origin	100 ml 500 ml	P30-1501 P30-1502
Fetal Bovine Serum Premium, South America origin	100 ml 500 ml	P30-3301 P30-3302
Sera Pro, Fetal bovine serum, low Endotoxin, EU approved	100 ml 500 ml	P30-5100 P30-5500 NEW
FBS EU Professional, Filtrated bovine serum, EU approved	100 ml 500 ml	P30-8100 P30-8500
Bovine Serum, variable origins	100 ml 500 ml	P30-0601 P30-0602
Calf serum - newborn, variable origins	100 ml 500 ml	P30-0401 P30-0402

FBS Good Product Family

The FBS Good product family contains specially processed serum products. Serum of selected batches is filtrated and separated into individual components by a sophisticated chromatographic method. The growth promoting components contained in the serum are then combined and restored in a defined process. Compared to conventional fetal bovine serum the FBS Good product family has been shown to support and promote cell growth of many different cell types equally well or even better.

Advantages

- Innovative new products
- Minor batch to batch variation
- Once tested always same quality
- No batch testing required
- No lot reservation required



FBS Good Product Family

FBS Good, Filtrated bovine serum, Australia origin	100 ml 500 ml	P40-39100 P40-39500	NEW
FBS Good, Filtrated bovine serum, EU approved	100 ml 500 ml	P40-37100 P40-37500	NEW
FBS Good, Filtrated bovine serum, US origin	100 ml 500 ml	P40-38100 P40-38500	NEW
FBS Good Forte, Filtrated bovine serum with Additive Fortifier, Australia origin	100 ml 500 ml	P40-49100 P40-49500	NEW
FBS Good Forte, Filtrated bovine serum with Additive Fortifier, EU approved	100 ml 500 ml	P40-47100 P40-47500	NEW
FBS Good Forte, Filtrated bovine serum with Additive Fortifier, US origin	100 ml 500 ml	P40-48100 P40-48500	NEW
FBS Good Biotech, Filtrated bovine serum, Australia origin, tested acc. EMEA 1793 and Ph. Eur. 2262*	100 ml 500 ml	P40-59100 P40-59500	NEW
FBS Good Biotech, Filtrated bovine serum, US origin, tested acc. EMEA 1793 and Ph. Eur. 2262*	100 ml 500 ml	P40-58100 P40-58500	NEW

^{*}tested upon request and produced after receipt of order

FBS Good

By developing FBS Good we wanted to create a naturally defined serum with a sustained growth promoting property and a higher safety. FBS Good only contains serum of highest quality from defined countries as specified. It is not blended or enhanced by addition of growth factors or proteins.

FBS Good Forte

By developing FBS Good Forte we wanted to create a naturally defined serum with an increased growth promoting property and a higher safety. Therefore, additional growth fortifying compounds have been added to increase cell proliferation. FBS Good Forte only contains serum of highest quality from defined countries as specified. In addition, growth promoting and stabilizing compounds (e.g. proteins, salts, sugars, vitamins) have been added to further enhance the stability of the serum as well as the proliferation of many different cell types.

FBS Good Forte advantages

- Reproducible enhanced growth properties
- Very low endotoxin level
- Suitable for many different cell types
- Continued high quality
- No more batch testing required

FBS Good advantages

- Reproducible growth properties
- Very low endotoxin level
- Suitable for a great variety of cells
- Continuous quality control

FBS Good Biotech

By developing FBS Good Biotech we wanted to create a naturally defined serum with a sustained growth promoting property and a higher safety. It is tested according to EMA (also known as EMEA) and Ph. Eur. guidelines for use in the bio-pharmaceutical industry. FBS Good Biotech only contains serum of highest quality from defined countries as specified. It is not blended or enhanced by addition of growth factors or proteins.

FBS Good Biotech advantages

- Tested according to EMEA CPMP/BWP/1793/02
- Tested according to Ph. Eur. 01/2008:2262
- Very low endotoxin level
- Suitable for bio-pharmaceutical processing
- Expanded quality control



Special Fetal Bovine Serum

Pansera ES, Fetal bovine serum, Australia origin, special designed for embryonic stem cells	100 ml 500 ml	P30-2605 P30-2606	NEW
Pansera ES, Fetal bovine serum, EU approved, special desgined for embryonic stem cells	100 ml 500 ml	P30-2601 P30-2602	
Pansera ES, Fetal bovine serum, US origin, special designed for embryonic stem cells	100 ml 500 ml	P30-2608 P30-2609	NEW
Fetal Bovine Serum Biotech, Australia origin, tested acc. EMEA 1793 and Ph. Eur. 2262*	100 ml 500 ml	P40-1301 P40-1302	NEW
Fetal Bovine Serum Biotech, US origin, tested acc. EMEA 1793 and Ph. Euro. 2262*	100 ml 500 ml	P40-1401 P40-1402	NEW

^{*}tested upon request and produced after receipt of order

Pansera ES

Our specially developed, proprietary processing methodology for serum enables us to offer a special fetal bovine serum for embryonic stem cells (ES).

Advantages

- Reproducible constant growth properties
- Improved cloning efficiency
- More undifferentiated clones
- Permanent strict quality control
- No need for further testing of different batches

Fetal Bovine Serum Biotech

The bio-pharmaceutical industry is facing a constantly growing demand for high quality, extensively tested fetal bovine serum originating from FDA-approved regions. PAN-Biotech is meeting this demand and offers a new product, Fetal Bovine Serum Biotech, which is tested according to EMA (also known as EMEA) and Ph. Eur. guidelines.

Fetal Bovine Serum Biotech originates exclusively from Australia or the USA, both of which are approved regions by FDA's CFR. The entire production process is followed by close inspections and quality controls – from the collection of raw material to the final production and sterile filtration procedure, all steps are documented and traceable.

Especially a possible contamination with bovine viruses has to be excluded. Therefore, a multitude of tests is performed to meet highest safety requirements.

This product is tested according to EMEA CPMP/BWP/1793/02 and Ph. Eur. 01/2008:2262 upon request. In addition, EMA/410/01 rev. 3 and EMA/CHMP/BWP/457920/2012 rev. 1 are also included as guidelines for testing procedures. Besides extensive testing for viral contamination, supplementary sterility testing is performed before, during and after filling of the product.

Application

Fetal Bovine Serum Biotech is particularly suited for the production of virus, vaccine, monoclonal antibodies, recombinant protein and growth factors, as well as the manufacture of other bio-pharmaceutical products.





Certificate of Analysis

Fetal Bovine Serum

Origin: Australia

I	Product	Description	Catalogue-No.	Size
	FBS	Fetal bovine serum, Australia origin	P30-1301 P30-1302	100 ml 500 ml

Lot No.: P140118

Date of production: January 09, 2014

Storage, stability, shipping:

Storage:

Stability: 6 years from date of production

Shipping: on dry ice

Parameter	Result	Units
Appearance	amber liquid	n.a.
pH value	7.46	n.a.
Osmolality	296	mOsm/kg
Hemoglobin	13.0	mg/100 ml
Endotoxin	0.113	ng/ml
Total protein Albumin alpha-Globulin beta-Globulin lgG	38.79 29.57 5.03 4.19 254	mg/ml mg/ml mg/ml mg/ml µg/ml
Glucose	109.1	mg/100 ml
Cholesterol	51.3	mg/100 ml
Triglycerides	45.0	mg/100 ml

- Contraction		Specification	Result
Sterility	Incubation at 32°C Incubation at 20°C	sterile sterile	sterile sterile
	Mycoplasma	not detected	not detected
Virus testin	ng		
Bovine herp	diarrhoea virus (BVDV) es virus (BHV-1) iza virus type 3 (PI-3)	negative negative negative	negative negative negative
Antibody te	esting		
Bovine herp	diarrhoea virus (BVDV) es virus (BHV-1) iza virus type 3 (PI-3)	serological titer serological titer serological titer	< 1:2 < 1:2 < 1:2

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Performance (cell culture tested)

Cell growth (SP2/0-Ag14)	Seed	day 2	day 5	day 7	[cells per ml]
Lot no. P140118	1.00x10e3	6.28x10e3	8.90x10e5	1.13x10e6	
Control serum	1.00x10e3	5.34x10e3	8.40x10e5	1.11x10e6	
Cell growth (L929)		5 to 200 to 200 to 200	200000000000000000000000000000000000000		[cells per ml]
Lot no. P140118	1.00x10e4	1.50x10e4	8.25x10e5	1.09x10e6	
Control serum	1.00x10e4	1.75x10e4	8.35x10e5	1.10x10e6	
No. of	color	ies/clones	absolute %	relative %	
Plating efficiency (L929)	257500	100000000000000000000000000000000000000	\$ 0000 Bayes 30 U.S.	100000000000000000000000000000000000000	
Lot no. P140118	286		58	105	
Control serum	257		55	100	
Cloning efficiency (SP2/0-Ag	14)			-05	
Lot. no. P140118	63		66	102	
Control serum	62		65	100	

TABLE 1:

Test	Method
pH value	Measured with pH-electrode
Osmolulity	Analyzed by freezing point depression
Hemoglobin	Determined spectrophotometrically at three different wave lengths
Endotoxin	Kinetic limulus amoebocyte lysate test (LAL)
Total protein	Colorimetric test (Biuret reaction)
Albumin, Globulins	Serum protein electrophoresis (SPEP)
lgG	Redal immune diffusion
Glucose	Colorimetric test (Trinder reaction)
Cholesterol	Colorimetric test (CHOD PAP)
Triglycerides	Colorimetric test (Trinder reaction)
Sterility	The absence of bacterial or fungal contamination is verified by dual incubation with Caso-Bouillon or Thioglycolat- Bouillon according to Ph. Eur. at 32 °C and 20 °C
Mycoplasma	Three different detection systems are used: DNA-binding fluorescence dye (DAPI), microscopic analysis of microbial outures and test kits which detect mycoplasma specific enzymes
Virus testing	The following viruses and the presence of their arribodies are tested by cytopathic effect: Bovine viral diarrhoea virus (BVDV), bovine herpes virus (BHV 1) and parainfluenza virus (PH3)
Cell growth	Growth test of murine myeloma cells (SP2/0-Ag14) and murine fibroblasts (L929)
Plating efficiency	Murine (broblasts (1,929) are plated into a Petri dish, stained with Giernsa and after 14 days of incubation the fixed cell colonies are counted (+ absolute plating efficiency). The results are normalized to a previously tested reference serum (+ relative plating efficiency).
Cloning efficiency	Murine myeloma cells (SP2/0-Ag14) are plated on microtiter plates (one cell per well). After 7 days of incubation the developed cell proposes are counted (absolute classical efficiency). The results are normalized to a previously tested

Suitability

FOR RESEARCH USE ONLY!

These products are intended for research or manufacturing use only. Not for use in animal or human clinical or diagnostic application.

reference serum (= relative cloning efficiency).

Raw material is collected in regularly inspected facilities and processed by PAN Biotech in compliance with current Ph. Eur. guidelines for Bovine Sera, Processing of raw material into finished serum product is performed by employees of PAN Biotech.

Since raw serum is not pre-aged before filtration, turbidity or floculent debris in form of precipitate may develop upon thawing or storage of the product. This occurrence does not adversely affect the performance of the serum.

Results shown in this compilation have been obtained by carefully performing standard test methods (see table 1). Since results for any specific test may vary depending on methodology, technical equipment, or test substances used, it is suggested that results for particularly important parameters be repeated by the end user of this product.

PAN Biotech has been assigned a Certificate of Suitability (Ref. No. R1-CEP 2002-167-Rev 00; renewed Nov/11/2008) by the European Directorate for the Quality of Medicines (EDQM) for production of bovine serum.

* n.a. = not available

Date: Feb/12/2014

Iris Podolski Quality Assurance

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2



100 ml	P30-2301
500 ml	P30-2302
100 ml	P30-3401
500 ml	P30-3402
100 ml	P30-2101
500 ml	P30-2102
100 ml	P30-2008
500 ml	P30-2009
100 ml	P30-2081
500 ml	P30-2085
100 ml	P30-1905
500 ml	P30-1906
100 ml	P30-1908
500 ml	P30-1909
100 ml	P30-1981
500 ml	P30-1985
100 ml	P30-2801
500 ml	P30-2802
100 ml	P30-3601
500 ml	P30-3602
	500 ml 100 ml 500 ml

Activated charcoal treated serum

Fetal bovine serum is heated in a water bath with dextran and activated charcoal. The activated charcoal, together with the substances bound to it, is then removed by centrifugation and filtration.

Application

- Work involving reduced hormone content (steroids)
- Work involving reduced growth factors (prevention of cell differentiation)
- Receptor studies (e. g. estrogens)
- Minimizes lot-to-lot variations in serum

Delipidized serum

Lipids are removed from serum by affinity chromatography.

Application

- Lipid metabolism studies
- Arteriosclerosis research

Dialyzed serum

Serum is dialyzed with a 10,000 Dalton exclusion membrane against physiological saline solution (alternative DPBS) until the glucose content is below 10 mg/100 ml.

Application

- Radioactive labeling studies
- Hormone-free applications
- Tests intolerant for small molecules such as nucleotides (hypoxanthine, thymidine), amino acids (serine, alanine etc.), sugars or metabolites

Gamma irradiated serum

Serum is exposed to irradiation > 25 kGy

Application

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- Biopharmaceutical production
- Virus production
- Vaccine production
- Manufacturing of diagnostic products



Heat inactivated serum

Serum is heated for 30 min to 56 °C in a water bath under repeated gentle mixing.

Application

- Measurements of lactate dehydrogenase in the culture supernatant as a marker for cell damage (serum LDH is inactivated by heat)
- Minimizes lot-to-lot variations in serum (all thermolabile components are removed)
- Studies on vitamins and growth factors
- Enhance viral safety, since heat-labile viruses are inactivated
- Tests that do not tolerate presence of complement (complement destruction)

Ultra low IgG serum

The average IgG level in fetal bovine serum is in the range of 70 to 330 μ g/ml. The IgG content in our Ultra low IgG serum is reduced by affinity chromatography (protein-G affinity column) to max. 5 μ g/ml. The biological activity of serum is not affected.

Application

- Antibody production
- Monoclonal antibodies
- Radioactive labeling

Tetracycline-free serum

Fetal bovine serum is tested for absence of tetracycline using the TET-off system (luciferase).

Application

- TET-on / TET-off regulated gene expression
- Transfections
- Expression studies

Other Animal Serum

Chicken serum	100 ml 500 ml	P30-0301 P30-0302
Donkey serum	100 ml 500 ml	P30-0101 P30-0102
Goat serum	100 ml 500 ml	P30-1001 P30-1002
Hamster serum	10 ml	P30-0210
Horse serum	100 ml 500 ml	P30-0701 P30-0702
Lamb serum	100 ml 500 ml	P30-0801 P30-0802
Mouse serum	10 ml 100 ml	P30-0200 P30-0201
Pig serum	100 ml 500 ml	P30-0901 P30-0902
Rabbit serum	100 ml 500 ml	P30-1101 P30-1102
Rat serum	10 ml 100 ml	P30-01901 P30-01901E
Sheep serum	100 ml 500 ml	P30-4101 P30-4102

All serum tested for virus and mycoplasma. Other serum upon request.



Human Serum

Human serum is manufactured from human plasma by addition of calcium chloride. This results in clotting of the plasma. After removing the clot, the human serum is washed and concentrated by ultra-filtration and finally filtered through a combination of depth- and membrane-filters.

Off-the-clot serum (True human serum)

Off-the-clot serum is prepared from human whole blood collected without anti-coagulant, allowed to clot at room temperature and then centrifuged to remove the clot. We provide single donor units as well as pooled off-the-clot serum. Off-the-clot serum is filtered through depth and membrane filters before filling.

Human serum	100 ml 500 ml	P30-2401 P30-2402
Human AB serum	100 ml 500 ml	P30-2501 P30-2502
Human AB serum (male)	100 ml 500 ml	P30-2901 P30-2902
Human serum off-the-clot	100 ml 500 ml	P30-2701 P30-2702

Serum Services

PAN-Biotech offers a variety of services and test procedures for your serum. We deliver these services fast and cost efficient, using the latest up-to-date techniques.

Profit from our expertise! If you need further special testing or particular services please contact PAN-Biotech. In most cases we can find a solution.

Special processing of serum lots

Treatment	Method
Charcoal absorbed	Serum is heated in a water bath with dextran and activated charco- al. The activated charcoal, together with the substances bound to it, is then removed by centrifugation and filtration.
Delipidized	Lipids are removed from serum by affinity chromatography
Dialyzed	Serum is dialyzed with a 10,000 Dalton exclusion membrane against physiological saline solution
Heat inactivated	Serum is heated for 30 min to 56 °C in a water bath under repeated gentle mixing
IgG reduced	The IgG content in serum is reduced by affinity chromatography to max. 5 μ g/ml. The biological activity of serum is not affected.
Sterile filtered	Serum passes a series of filters with decreasing pore sizes. The last filtration step is done with a 0.2 μm pore size sterile filter.

Virus testing according EMEA guidelines

The following virus tests are performed according to EMEA guideline CPMP/BWP/1793/02:

- Bluetongue and related orbi viruses
- Bovine adenovirus
- Bovine parvovirus

- Bovine respiratory syncytial virus (BRSV)
- Bovine viral diarrhoea virus (BVDV)
- Rabies virus (rabies)
- Reo virus
- Bovine polyoma virus (BPyV)



Serum testing

Test	Method
Albumin, Globulins	Serum protein electrophoresis
Bacterial count	Detection of total number of viable aerobic germs will be either done by membrane filtration or plate-flush-method or as surface method. The microorganisms are detected as colony forming units per ml (CFU/ml) on Caso agar plates.
Cell growth	Growth test of murine myeloma cells (SP2/0-Ag14) and murine fibroblasts (L929)
Cholesterol	Colorimetric test (CHOD-PAP)
Cloning efficiency	Murine myeloma cells (SP2/0-Ag14) are plated on microtiter plates (one cell per well). After 7 days of incubation the developed cell colonies are counted (= absolute cloning efficiency). The results are normalized to a previously tested reference serum (= relative cloning efficiency).
Endotoxin	Kinetic limulus amoebocyte lysate test (LAL)
Glucose	Colorimetric test (Trinder reaction)
Hemoglobin	Determined spectrophotometrically at three different wave lengths
IgG	Radial immune diffusion
Mycoplasma	Three different detection systems are used: DNA-binding fluore- scence dye (DAPI), microscopic analysis of microbial cultures and test kits which detect mycoplasma specific enzymes
Osmolality	Analyzed by freezing point depression
pH value	Measured with pH-electrode
Plating efficiency	Murine fibroblasts (L929) are plated into a Petri dish. After 14 days of incubation the fixed cell colonies are stained with Giemsa and counted (= absolute plating efficiency). The results are normalized to a previously tested reference serum (= relative plating efficiency).
Sterility	The absence of bacterial or fungal contamination is verified by dual incubation with Caso-Bouillon or Thioglycolat-Bouillon according to Ph. Eur. at 32 °C and 20 °C
Tetracycline	Tested by a TET-off system of a CHO-luc cell line. Absence of tetracyline induces expression of luciferase, which is quantified using the luciferase test system from Promega.
Total protein	Colorimetric test (Biuret reaction)
Triglycerides	Colorimetric test (Trinder reaction)
Virus testing	The following viruses and the presence of their antibodies are tested by cytopathic effect: Bovine viral diarrhoea virus (BVDV), bovine herpes virus (BHV 1) and parainfluenza virus (PI-3)



Serum-free Systems

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Introduction

Basics

Cell culture, the cultivation of cells isolated from live tissue in vitro (in the test tube), is an acknowledged and valuable tool in biomedical research for the acquisition of reproducible data. In addition, highly effective substances are produced in large-scale for medicine or biopharmaceutical and academic research by means of cell cultures to an ever increasing degree (e.g. insulin, growth factors, monoclonal antibodies, or clotting factors).

The cell culture with serum

Cell cultures in vitro need nutrient solutions, so-called media, which provide an as close as possible simulation of the in vivo situation (in live organism). For this purpose these cell culture media – mixtures of nutrients, salts, trace elements, buffers, growth factors, protective binding and transport proteins and many additional components – have to be supplemented by a most natural, highly complex additive mixture. For many years, animal-derived and also human sera were the means of choice for production related reasons as well as for a lack of alternatives in more defined cell culture nutrients.

Function of serum in cell cultures

- Hormone factors stimulate cell growth, proliferation and differentiation
- Attachment factors favour or enable the attachment of cells to the culture dish (biomatrix)
- Transport and binding proteins support the supply with hormones, minerals and lipids
- Serum proteins bind toxic substances and stabilize sensitive growth factors

However, the use of serum in cell cultures, usually fetal bovine serum (FBS), is problematic for several reasons.

Disadvantages of serum in cell culture

- The composition of serum is variable and depends on the age of the fetus, on the origin and feeding of the animals, and on the time of year at slaughter
- Serum batches have to be tested extensively for their suitability before use
- Test results are often unsatisfactory and often not comparable because of the undefined and inconsistent nature of serum
- Risk of a contamination with bacteria, fungi, mycoplasma and virus from serum
- Risk of contamination with TSE agents (transmissible spongiform encephalopathy)
- Possibility of impurities in the end products due to residual serum proteins or pyrogens
- Time-consuming purification of the end products from culture media containing serum
- Uncertain availability and increasing cost of serum.

The serum-free cell culture

Because of the numerous disadvantages of a serumcontaining cell culture, for many years considerable research and development efforts have been undertaken to finally establish cell cultures under serum-free conditions.

Advantages of a serum-free cell culture

- Lower risk of contamination with bacteria, fungi, mycoplasma or virus
- Defined and reproducible formulations result in more convincing and comparable research data
- Time-consuming batch tests are dispensable
- Elimination of a source for possible infectious agents (prions)
- Ease of purification of end products
- Fulfilment of legal requirements for the manufacturing of medical products
- Reduction of impurities in end products by culture residues





Definitions

Serum-free media

Serum-free media can be used without any addition of serum or FBS and cell cultures usually perform similar to or even better as with serum. In some compositions, defined and purified components or sub-fractions of serum proteins, growth factors, or hormones (e.g. albumin, transferrin or insulin) are contained, or protein hydrolysates or gland extracts (BPE, BBE) are used.

Protein-free media

Protein-free media support the cell growth without addition of any protein. They contain higher concentrations of amino acids or herbal hydrolysates. Chemically defined formulation do not contain peptones or hydrolysates.

Chemically defined media

Chemically defined media are completely free of any animal or human components. In addition, these formulations do not contain peptone or hydrolysate. All components have a known chemically defined structure and composition. This results in a very constant and stable formulation with positive effects on quality, reproducibility and reduction of inter-batch variability.

PAN-Biotech serum-free media products

Serum substitute

Many users strive to keep their basal medium, because the cells are acquainted to these media over a long time or extensive efforts have been made to find a suitable basal medium. With this in mind, PAN-Biotech has developed easy to use serum substitutes which can fully replace FBS in the medium. Since different cell types (e.g. adherent or suspension cells) require different nutritional and attachment factors, we have developed two different serum substitutes for these kinds of cells. Panexin NTA is designed for adherent cells and Panexin NTS is designed for suspension cells.

These serum substitutes can be used in many cases without an adaptation of the cells and no or little weaning. In this case, our Panexins give an instant advantage over conventional serum containing cultures, eliminating many of the above described limitations of cell cultures with FBS.

Stem and progenitor cell media

Research and development in the field of stem cell biology has been tremendously advanced in the last decade. Today, some cell types are being used in clinical studies or applications and several more are close to being employed in cellular therapy. One important aspect for any application of stem and progenitor cells in patients is the isolation and expansion of these cells under defined conditions. For this purpose, the presence of FBS in such cell cultures is undesirable.

PAN-Biotech is offering a full range of serum-free media for stem and progenitor cells for the most important fields of research and development. Some of these stem cell media are free of animal-derived components, enabling the culture of cells in conditions close to clinical application.

Quality assurance

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Each batch of serum-free media is produced only with pretested premium raw materials to ensure the highest quality standards. Water is the main and determining basal ingredient for any cell culture medium. The condition of our pyrogen-free water is of extra purity with a conductance value of 0.055 µS/cm. It is regularly tested, since a minimal variation in water quality will have detrimental effects on the cells in a serum-free culture. Each batch of Panexin or Panserin will not be released unless the quality control process is finished and all the required specifications have been met.



Product Numbers

Serum Substitutes

Panexin NTA ⁽¹⁾	100 ml 500 ml	P04-95700 P04-95750
Panexin NTS ⁽¹⁾	100 ml 500 ml	P04-95800 P04-95850
Panexin BMM ⁽¹⁾	100 ml	P04-951SA2

Serum-free Media

Panserin 401 ⁽¹⁾	100 ml	P04-710401M
	500 ml	P04-710401
Panserin 411 ⁽¹⁾	100 ml	P04-710411M
	500 ml	P04-710411
Pasnerin 411S ⁽¹⁾	100 ml	P04-7411S0
	500ml	P04-7411S1
	1 L	P04-71411S
Panserin 412 ⁽¹⁾	100 ml 500ml	P04-710412M
2		P04-710412
Panserin 413 with 1 supplement ⁽³⁾	500 ml	P04-710413
Panserin 416 with 1 supplement ⁽³⁾	500 ml	P04-710416
Panserin H4000 ⁽¹⁾	100 ml	P04-714000M
	500 ml	P04-714000
Panserin H8000 ⁽¹⁾	100 ml	P04-718000M
	500 ml	P04-718000
Panserin C6000 ⁽¹⁾	100 ml	P04-716000M
	500 ml	P04-716000
Panserin 293A ⁽¹⁾	100 ml	P04-710608M
	500 ml	P04-710608
Panserin 293S ⁽¹⁾	100 ml	P04-710609M
	500 ml	P04-710609
Panserin T3 ⁽³⁾	100 ml	P04-710110
	500 ml	P04-710100
Panserin ProVero ⁽³⁾	100 ml	P04-710613M
D 1 =04(1)	500 ml	P04-710613
Panserin 701 ⁽¹⁾	100 ml 500 ml	P04-710701M P04-710701
D (2)		
Panserin 801 with 6 supplements ⁽³⁾	500 ml	P04-710801
Panserin PX10 ⁽¹⁾	500 ml	P04-710PX10
Panserin PX40 ⁽¹⁾	500 ml	P04-710PX40
Spodopan ⁽¹⁾	100 ml	P04-850100
	500 ml	P04-850500
Panserin S2 ⁽¹⁾	100 ml	P04-710210
	500 ml	P04-710200
Endopan 300 SL ready-to-use (3)	500 ml	P04-00650
Endopan 300 SL kit (3)	500 ml	P04-0065K



Serum Substitutes

Panexin NTA

Panexin NTA is a defined serum substitute for the cultivation of adherent cells under serum-free conditions. Panexin NTA is developed with an unique technology and contains a special 3-dimensional substance release system (3D-SRS) for an optimal support of cells with nutrients and growth stimulants.

The ready to use, sterile solution is added to the culture medium in a final concentration of 10%. It supports the adherent growth of many cell types in an optimum manner

Composition

Panexin NTA contains purified proteins, lipids, salts, amino acids, trace elements, attachment factors and hormones in an optimized formulation and a new 3-dimensional substance release system (3D-SRS). Panexin NTA contains no growth factors, undefined hydrolysates or peptones.

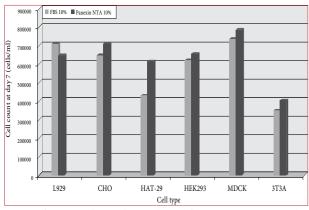


Fig.1: Efficiency and Growth Stimulation of Panexin NTA compared to FBS (each 10 % in DMEM/F12)

Suitability

Panexin NTA is suitable for the cultivation of a variety of adherent cells under serum-free culture conditions.

Special advantages

It has been shown for many cell lines that Panexin NTA can fully replace FBS. Due to selected and pretested raw materials Panexin NTA batches are very homogeneous. Therefore the complex batch testing known from FBS can be omitted with the use of Panexin NTA. In addition, there is no need to change the previously used basal medium.

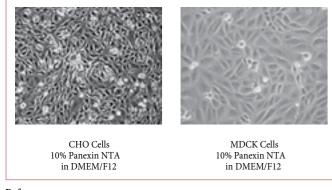
Panexin NTA is completely defined and contains no undefined peptones or hydrolysates. Therefore, the interpretation of results from studies on effects of individually added growth factors is easier and more reliable in serum-free conditions. For cell lines which require specific growth factors these should be added in a concentration as previously used. As a basal medium you may use classical standard media such as RPMI 1640, DMEM (high or low glucose), DMEM/F12, IMDM and so on. Make sure that L-glutamine is present in sufficient quantity (possibly supplement glutamine).

Depending on the cell type, some differences in morphology or proliferation rate may be observed with various standard media. Many applications were performed with DMEM or DMEM/F12 for adherent cells. With these combinations very good growth stimulation was achieved in a range of 5% to 15% Panexin NTA.

For more demanding cells an adaptation to Panexin NTA may be necessary.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panexin NTA and can also be found at www.pan-biotech.com.



References

- a) Hashimoto J et al. (2006) Regulation of Proliferation and Chondrogenic Differentation of Human Mesenchymal Stem Cells by Laminin-5 (Laminin-332). Stem Cells 24:2346
- b) Traeger T et al. (2008) Detrimental Role of CC Chemokine Receptor 4 in Murine Polymicrobial Sepsis. Infection and Immunity 11:5285

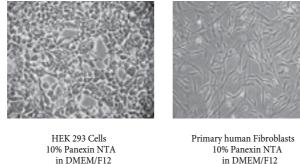


Fig.2: Different Cell Lines in DMEM/F12 with 10 % Panexin NTA

Panexin NTA ⁽¹⁾	100 ml	P04-95700
Panexin NTA	500 ml	P04-95750

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Serum Substitutes

Panexin NTS

Panexin NTS is a defined serum substitute for the cultivation of suspension cells under serum-free conditions. Panexin NTS is developed with a unique technology and contains a special 3-dimensional substance release system (3D-SRS) for an optimal support of cells with nutrients and growth stimulants. The ready-to-use, sterile solution is added to the cell culture medium in a final concentration of 10%. It supports the growth of many cell types in an optimum manner.

Composition

Panexin NTS contains purified proteins, lipids, salts, amino acids, trace elements, and hormones in an optimized formulation and a new 3-dimensional substance release system (3D-SRS). Panexin NTS contains no growth factors, undefined hydrolysates or peptones.

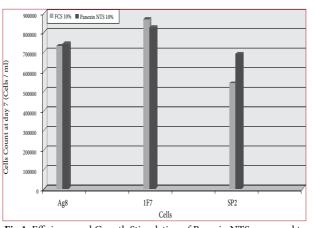


Fig.1: Efficiency and Growth Stimulation of Panexin NTS compared to FBS (each 10 % in RPMI)

Suitabilit

Panexin NTS is suitable for the cultivation of a variety of non-adherent suspension cells under serum-free conditions

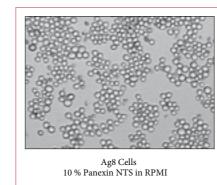
Special advantages

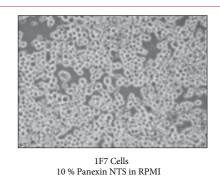
Panexin NTS can be used for many cell lines to replace FBS. Due to selected and pretested raw materials Panexin NTS batches are very homogeneous. Therefore the complex batch testing known from FBS can be omitted with the use of Panexin NTS. In addition, there is no need to change the previously used basal medium. Panexin NTS is completely chemically defined and contains no growth factors, undefined peptones or hydrolysates. Therefore, the interpretation of results from studies on effects of individually added growth factors is easier and more reliable in serum-free conditions. For cell lines which require specific growth factors, these should be added in a concentration as previously used.

For more demanding cells an adaptation to Panexin NTS may be necessary.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panexin NTS. In addition, instructions for use can also be found at www.pan-biotech.com.





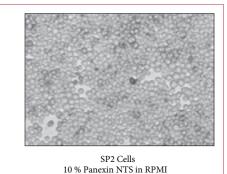


Fig.2: Different Cell Lines in RPMI with 10 % Panexin NTS

Panexin NTS ⁽¹⁾	100 ml	P04-95800
Panexin N 15°	500 ml	P04-95850

Reference

a) Breitbach K et al. (2009) Caspase-1 Mediates Resistance in Murine Melioidosis. Infection and Immunity 4:1589

b) Into T et al. (2008) Regulation of MyD88-Dependent Signaling Events by S Nitrosylation Retards Toll-Like Receptor Signal Transduction and Initiation of Acute-Phase Immune Responses. Molecular and Cellular Biology 4:1338

) usually on stock, (2) minimum order 10 l, (3) available upon reque



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Serum Substitutes

Panexin BMM

Panexin BMM is a defined serum substitute for the cultivation of macrophages from mouse bone marrow (murine bone marrow derived macrophages, BMM) under serum-free conditions. The ready-to-use sterile solution in a final concentration of 5 % is added to the basal medium RPMI 1640, supplemented with 50 µM Mercaptoethanol and 2 ng/ml GM-CSF mur. rec.

Composition

Panexin BMM contains purified proteins, lipids, salts, amino acids, trace elements, attachment factors and hormones in an optimized formulation. It contains no growth factors, undefined hydrolysates or lysates (e. g. Peptones).

Suitability

Panexin BMM has been developed for the generation of murine macrophages from bone marrow under serum-free conditions. This achieves standardized conditions and reproducible results.

Panexin BMM⁽¹⁾ P04-951SA2 100 ml

Panexin BMM allows the generation of murine macrophages from bone marrow under standardized serum-free conditions. The results will be more comparable, as undefined components - like in serum-containing cultures - are eliminated. In Panexin BMM matured macrophages will show excellent attachment capabilities.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panexin BMM and can also be found at www.pan-biotech.com.

Kristin Eske, Katrin Breitbach, Jens Köhler, Patimaporn Wongprompitak and Ivo Steinmetz (2008). Generation of murine bone marrow derived macrophages in a standardised serum-free cell culture system, Journal of Immunological Methods.



Serum-free Media

Panserin 401

Panserin 401 is a complete ready-to-use medium for the serum-free cultivation of a multitude of adherent and non adherent cells.

Composition

Based on Iscove's MEM, trace elements, albumin, cholesterol, soya lipids and vitamins were added to the medium. It does not contain any growth or attachment factors.

Panserin 401 is a multi-purpose medium suitable for a variety of cells. As the medium contains no growth factors there is a possibility to investigate the effects of specific growth factors added to the cell culture. Panserin 401 does not contain any attachment factors. With some cell types a pre-treatment of the cell culture vessels with gelatine, collagen, poly-D-lysine or fibronectin may support or enable a culture under serum-free conditions. Please note that a coating may be especially important with low seeding densities. With every adaption to serum-free media, changes of the cells should be taken into consideration. These changes may concern morphology, karyotype, surface markers and so on. Thus cells in serum-free medium may not be identical with those from cultures containing serum in which they originated (selection).

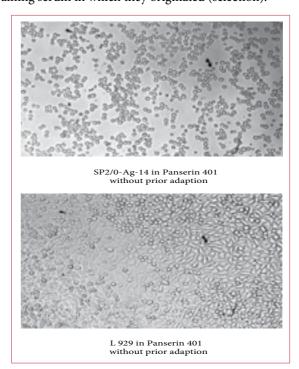


Fig 1.: SP2/0-Ag-14 and L929 in Panserin 401

Panserin 401 ⁽¹⁾	100 ml	P04-710401M
1 4110 01111 101	500 ml	P04-710401



Among others the following cells have been cultivated successfully:

- Hybridoma
- Lymphocytes
- Macrophages
- Fibroblasts
- Melanocytes
- Carcinoma cells
- HEK-cells
- HeLa-cells
- CHO-cells

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 401. In addition, instructions for use can also be found at www.pan-biotech.com.

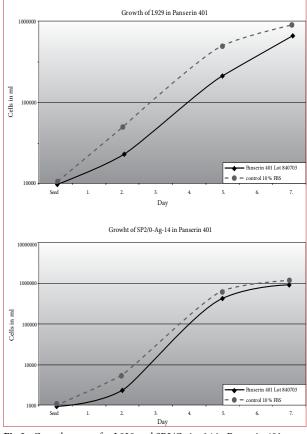


Fig 2.: Growth curves for L929 and SP2/O-Ag-14 in Panserin 401

www.pan-biotech.com

- a) Pilar S et al. (2002) Contribution of CD3y to TCR regulation and signaling in human mature T lymphocytes. International Immunology
- b) Toptan T et al. (2010) Rhadinovirus vector-derived human telomerase reverse transcriptase expression in primary T cells. Gene Therapy 17:653
- c) Martin F et al. (2005) Lentiviral vectors transcriptionally targeted to hematopoietic cells by WASP gene proximal promotor sequences. Gene Therapy 12:715
- d) Montzka K et al. (2010) Expansion of human bone marrow derived mesenchymal stromal cells: serum-reduced medium is better than conventional medium. Cytotherapy 5:587

Panserin 411

Panserin 411 is a complete, ready-to-use medium for the serum-free cultivation of a multitude of adherent and non adherent cells which are Insulin-dependent (e.g. CHO-cells).

Composition

Based on Iscove's MEM, trace elements, albumin, cholesterol, soya lipids, vitamins and insulin were added to the medium. It does not contain any growth or attachment factors.

Suitability

Panserin 411 is a multi-purpose medium suitable for a variety of cells. In Panserin 411 adherent as well as non adherent cells can be cultivated. As the medium contains no growth factors there is a possibility to investigate the effects of specific growth factors added to the cell culture. Panserin 411 does not contain any attachment factors. With some cell types a pre-treatment of the cell culture vessels with gelatine, collagen, poly-D-lysine or fibronectin may support or enable a culture under serum-free conditions. Please note that a coating may be especially important with low seeding densities.

With every adaption to serum-free media, changes of the cells should be taken into consideration. These changes may concern morphology, karyotype, surface markers and so on. Thus cells in serum-free medium may not be identical with those from cultures containing serum in which they originated (selection).

Instructions for use

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Detailed instructions will be provided with the accompanying datasheet for Panserin 411 and can also be found at www.pan-biotech.com.

Panserin 411 ⁽¹⁾	100 ml 500 ml	P04-710411M P04-710411
Panserin 411S ⁽¹⁾	100 ml 500 ml 1 L	P04-7411S0 P04-7411S1 P04-71411S

Panserin 411S

Panserin 411S is a complete, ready-to-use medium for the serum-free cultivation of myeloid and lymphoid cells for cytological examination.

Composition

Based on RPMI 1640 medium, additional trace elements, albumin, cholesterol, soy lipids, vitamins and hormones are added.

Suitability

Panserin 411S is a serum-free complete medium for the cultivation of myeloid and lymphoid cells from peripheral blood or bone marrow. It is therefore suitable for a rapid expansion of blood cells in order to investigate leukemic diseases (ALL, AML, CLL, CML, MPN, MDS). The state of the art diagnostic techniques of leukemic diseases are based on the interaction of cytomorphology including cytochemistry with immunophenotyping, chromosome banding analysis, FISH and molecular genetics. In Panserin 411S the number and quality of metaphases are significantly higher and independent of individual batches as compared to serum-containing media.

Suitability

Cells $(1x10^7)$ are seeded in 5 ml Panserin 411S. Depending on the assay or quality of raw material, an un-stimulated culture and another 1-3 cultures with appropriate growth factors are prepared. The culture time is 24 to 72 hours at 37° C in an incubator with 5% CO₂.

The processing of the metaphases is done with hypotonic KCl solution and Carnoy's fixative.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 411S and can also be found at www.pan-biotech.com.

(1) usually on stock, (2) minimum order 10 i, (3) available upon request



Serum-free Media

Panserin 412

Panserin 412 is a complete, ready-to-use medium for the serum-free cultivation of a multitude of adherent cells.

Composition

Based on Iscove's MEM, trace elements, albumin, cholesterol, soya lipids, vitamins and insulin were added to the medium. It does not contain any growth or attachment factors.

Suitability

Panserin 412 is a multi-purpose medium suitable for a variety of adherent cells. Panserin 412 contains special attachment factors for the successful cultivation of cells that hardly attach. With every adaption to serum-free media, changes of the cells should be taken into consideration. These changes may concern the morphology, the karyotype, the surface marker etc. Thus cells in serum-free medium don't always have to be identical with those from the culture containing serum in which they originate (selection).

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 412 and can also be found at www.pan-biotech.com.

Panserin 413

Panserin 413 is a medium for the serum-free cultivation of lymphocytes from whole blood.

Composition

Based on RPMI 1640/DMEM-F12, trace elements, albumin, cholesterol, soya-lipids and vitamins were added to the medium. A growth factor mixture is also supplied which has to be added to the medium immediately before use.

Suitability

Panserin 413 has been developed for the cultivation of lymphocytes from whole blood. Normally blood cells die rather quickly in culture, only lymphocytes can be cultivated over multiple divisions in culture. To achieve a division of non-proliferating cells, the cells must be stimulated with certain mitogens. These mitogens are mostly herbal lectins (phytohemagglutinin, PHA).

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 413 and can also be found at www.pan-biotech.com.

Panserin 416

Panserin 416 is a serum-free medium (basal medium) which is, after supplementation with growth factors, suitable for the production of dendritic cells.

Composition

Based on RPMI 1640/DMEM/F-12, trace elements, albumin, cholesterol, soya-lipids and vitamins were added to the medium. A growth factor mixture is also supplied which has to be added to the medium just before use.

Suitability

Dendritic cells are highly specialized antigen-presenting cells and can initiate and regulate antigen-specific immune responses. This ability can be used in order to generate immune responses against certain proteins of tumour cells and thus the immune system itself could be able to fight against tumours. Dendritic cells have been isolated from a great variety of non-lymphatic and lymphatic tissues of human beings, mice and other species.

For the generation of tumour vaccines, dendritic cells can be produced from the peripheral blood of tumour patients. In clinical studies the principal effectiveness of a vaccination with dendritic cells has been shown.

Production and serum-free cultivation of dendritic cells from mononuclear cells of peripheral blood (PBMC).

Serum-free cultivation of dentritic cells in Panserin 416 After the last washing step the mononuclear cells are transferred with a cell density of 1 x 10^7 cells/ml into Panserin 416. In order to remove non-adherent cells, culture dishes are put into the incubator for 2 hours. Then the supernatant is carefully taken off and replaced by new Panserin 416. GM-CSF (800 U/ml) and interleukin-4 (500 U/ml) are added as growth factors. The culture dishes are incubated for another 6 days in the incubator and every day half of the medium is replaced by new medium which is supplemented with GM-CSF and IL-4.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 416 and can also be found at www.pan-biotech.com.

Panserin 412 ⁽¹⁾	100 ml 500 ml	P04-710412M P04-710412
Panserin 413 ⁽³⁾	500 ml	P04-710413
Panserin 416 ⁽³⁾	500 ml	P04-710416



) usually on stock, (2) minimum order 10 l, (3) available upon request

Panserin H4000

Panserin H4000 is a protein-free ready-to-use medium for an optimized growth of myeloma and hybridomacell lines in suspension culture for the production of monoclonal antibodies. Panserin H4000 is suitable for spinner cultures, roller bottles and tissue culture bioreactors.

Composition

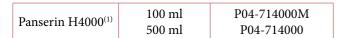
Panserin H4000 consists of a balanced mixture of salts, amino acids, vitamins, trace elements, hormones and is enriched with selected herbal hydrolysates for an optimized growth of myeloma and hybridoma cell lines. As Panserin H4000 is free of animal or human components it is predestined for the use in sensitive production areas (e.g. production of diagnostic or therapeutic tools) where safety requirements prohibit the use of human or animal components.

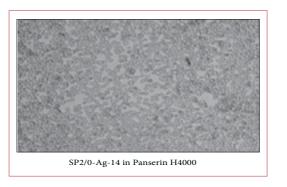
Suitability

Cultivation of myeloma and hybridoma cell lines for the production of monoclonal antibodies.

Special advantages

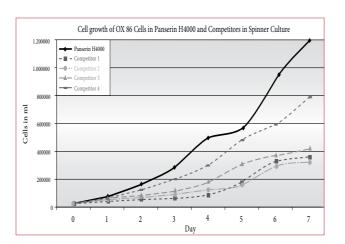
The formulation of the protein-free Panserin H4000 with a low concentration of plant hydrolysates enables a high cell yield in combination with excellent production rates of monoclonal antibodies. The ready to use protein-free medium allows easy handling and therefore reduces contamination risks and ensures for an easy and economic purification of the final products in downstream processes.





Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin H4000. In addition, instructions for use can also be found at www.pan-biotech.com.



Serum-free Media

Panserin H8000

Panserin H8000 is a protein-free, ready-to-use medium for an optimized growth of cholesterol-dependent myeloma and hybridoma cell lines in suspension culture for the production of monoclonal antibodies. Panserin H8000 is suitable for spinner cultures, roller bottles and bioreactors.

Composition

Panserin H8000 consists of a balanced mixture of salts, amino acids, vitamins, trace elements, hormones, bioavailable cholesterol and is enriched with selected herbal hydrolysates for an optimized growth of cholesterol dependent myeloma and hybridoma cell lines.

Suitability

Cultivation of cholesterol-dependent myeloma and hybridoma cell lines for the production of monoclonal antibodies.

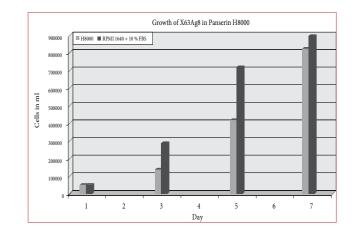
Special advantages

The formulation of the protein-free Panserin H8000 with a low concentration of plant hydrolysates enables a high cell yield in combination with excellent production rates of monoclonal antibodies. As Panserin H8000 is free of animal or human components it is predestined for the use in sensitive production areas (e.g. production of diagnostic or therapeutic tools) where safety requirements prohibit the use of human or animal components. The ready-to-use protein-free medium allows easy handling and therefore reduces contamination risks and ensures an easy and economic purification of final products in the downstream processing.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin H8000. In addition, instructions for use can also be found at www.pan-biotech.com.

Most hybridoma cell lines can be directly transferred from a serum-containing culture into a protein-free suspension culture. It should be noted here that the seeding density should be at least $1-3 \times 10^5$ cells.



Panserin H8000 ⁽¹⁾	100 ml	P04-718000M
Panserin fi8000	500 ml	P04-718000

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Panserin C6000

Panserin C6000 is a protein-free ready to use medium for an optimized growth of CHO-cells (Chinese Hamster Ovary) and their recombinant derivates in suspension culture. These cells are often used for the production of recombinant proteins for diagnostic or therapeutic purposes. Panserin C6000 is suitable for spinner cultures, roller bottles and tissue culture flasks and bioreactors.

Composition

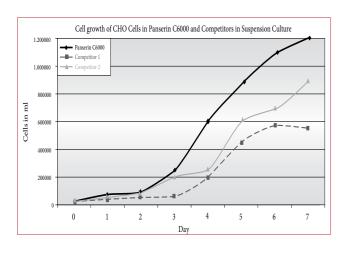
Panserin C6000 consists of a balanced mixture of salts, amino acids, vitamins, trace elements, hormones and is enriched with select herbal hydrolysates for an optimized growth of CHO-cells in suspension culture. As Panserin C6000 is free of animal or human components it is predestined for the use in sensitive production areas (e.g. production of diagnostic or therapeutic tools) where safety requirements prohibit the use human or animal components.

Suitability

Protein-free cultivation of CHO-cells and their recombinant derivates in suspension culture for the production of recombinant proteins for diagnostics or therapeutic purposes.

Special advantages

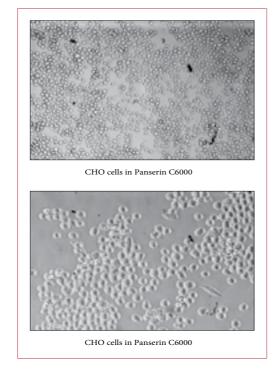
The formulation of the protein-free Panserin C6000 with a low concentration of plant hydrolysates enables a high cell yield in combination with excellent production rates of recombinant proteins. The ready to use complete protein-free medium allows easy handling and therefore reduces contamination risks and ensures for an easy and economic purification of the final products in downstream processes. Due to the optimized composition of Panserin C6000 the cells expand and grow in single-cell suspension with a very low tendency to form aggregates.



Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin C6000. In addition, instructions for use can also be found at www.pan-biotech.com.

Most CHO-cells can be directly transferred from a serum containing adherent culture into the protein-free suspension culture. In most cases the stable suspension culture is developing within about 2 weeks.



Panserin C6000 ⁽¹⁾	100 ml	P04-716000M
Panserin Coudo	500 ml	P04-716000

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Serum-free Media

Panserin 293A

Panserin 293A is a complete ready to use medium for the serum-free cultivation of HEK293 cells (Human Embryonic Kidney) in adherent culture.

Composition

Based on DMEM additional trace elements, albumin, cholesterol, soy lipids, vitamins and hormones have been added to the medium.

Suitability

Panserin 293A is a particularly enriched medium optimized for the growth of HEK293 cells in adherent culture. HEK293 is frequently used for the expression of recombinant proteins and the proliferation of adenoviruses. Panserin 293A promotes a rapid attachment of the cells and guarantees high cell growth rates.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 293A. In addition, instructions for use can also be found at www.pan-biotech.com.

Panserin 293A ⁽¹⁾	100 ml	P04-710608M
Panserin 295A	500 ml	P04-710608
Panserin 293S ⁽¹⁾	100 ml	P04-710609M
Panserin 2935	500 ml	P04-710609

Panserin 293S

Panserin 293S is a complete ready to use medium for the serum-free cultivation of HEK293 cells (Human Embryonic Kidney) in suspension culture.

Composition

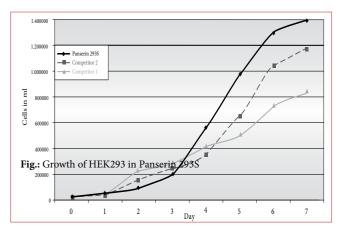
Based on DMEM/F12 medium additional trace elements, cholesterol and herbal hydrolysates have been added. Panserin 293S does not contain any proteins or components of animal or human origin.

Suitability

Panserin 293S is a particularly enriched medium optimized for the growth of HEK293 cells in suspension culture and quickly provides high cell densities. Due to its protein-free formulation the purification of final products (recombinant proteins, viruses) from the cell culture is more convenient and economic. Cell clustering - often seen in serum-free suspension cultures – will be reduced significantly in Panserin 293S.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 293S. In addition, instructions for use can also be found at www.pan-biotech.com.



1) usually on stock, (2) minimum order 10 l, (3) available upon request



Panserin T3

Panserin T3 is a ready-to-use serum-free complete medium for the cultivation of 3T3 cells in suspension culture.

Composition

Panserin T3 is a defined serum-free complete medium. Based on Iscove's MEM, this medium was supplemented with cholesterol, soy lipids, albumin, vitamins and trace elements. It contains no growth and attachment factors.

Suitability

Panserin T3 was developed for the serum-free cultivation of mouse fibroblasts (3T3A) in suspension.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin T3. In addition, instructions for use can also be found at www.pan-biotech.com.

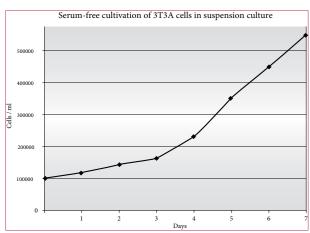
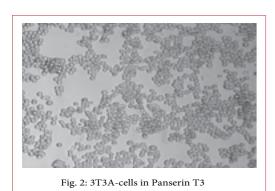


Fig.1: Growth of 3T3A cells in Panserin T3



Panserin T3⁽³⁾

100 ml
500 ml

P04-710110
P04-710100

Panserin ProVero⁽³⁾

100 ml
500 ml
P04-710613M
P04-710613

Panserin ProVero

Panserin ProVero is a complete serum-free medium ready to use for the cultivation of Vero cells (kidney epithelial cells from African green monkey) in an adherent culture.

Composition

Panserin ProVero is based on DMEM/F12. It contains trace elements, albumin, cholesterol, soy lipids, vitamins, hormones and attachment factors.

Suitability

Cultivation of Vero cells in adherent culture (e.g. roller bottles)

Special advantages

Highly enriched medium for the fast growth and culture of adherent Vero cells.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin ProVero. In addition, instructions for use can also be found at www.pan-biotech.com.

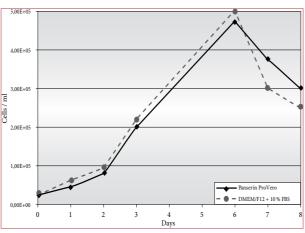


Fig.1: Growth of Vero cells in Panserin ProVero



(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Serum-free Media

Panserin 701

Panserin 701 is a complete ready-to-use serum-free medium for the cultivation of lymphocytes from whole blood

Composition

Based on Iscove's MEM the medium is enriched with additional trace elements, albumin, cholesterol, lipids and vitamins. It contains the mitogen phythemagglutinin (PHA) for a growth stimulation of lymphocytes.

Suitability

Panserin 701 has been developed for the serum-free cultivation of lymphocytes from whole blood. The herbal lectin (PHA) in Panserin 701 stimulates cell division.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 701. In addition, instructions for use can also be found at www.pan-biotech.com.

Panserin 701 ⁽¹⁾	100 ml 500 ml	P04-710701M P04-710701
Panserin 801 ⁽³⁾	500 ml	P04-710801

Panserin 801

Panserin 801 is a serum-free, ready-to-use medium for the cultivation of human keratinocytes.

Composition

MCDB-153 is used as basal medium to which the supplied supplements have to be added just before use.

These supplements are:

- Epidermal Growth Factor (EGF)
- Insulin
- Hydrocortisone
- Ethanolamine
- Phosphoethanolamine
- Pituitary Extract (BPE)

Suitability

Panserin 801 has been developed for the serum-free cultivation of human keratinocytes. Panserin 801 selectively supports the growth of human keratinocytes and concurrently prevents the overgrowth with fibroblasts.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin 801. In addition, instructions for use can also be found at www.pan-biotech.com.

usually on stock, (2) minimum order 10 l, (3) available upon request



Panserin PX10

Panserin PX10 is a ready-to-use serum-free complete medium for the cultivation of myeloma- and hybridoma cells for the production of monoclonal antibodies.

Composition

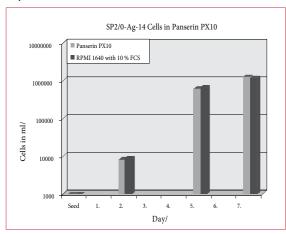
Based on RPMI 1640/DMEM/F-12, trace elements, albumin, cholesterol, soy lipids, vitamins and hormones were added to the medium. The medium does not contain any growth factors.

Suitability

Cultivation of myeloma- and hybridoma cells for the production of monoclonal antibodies.

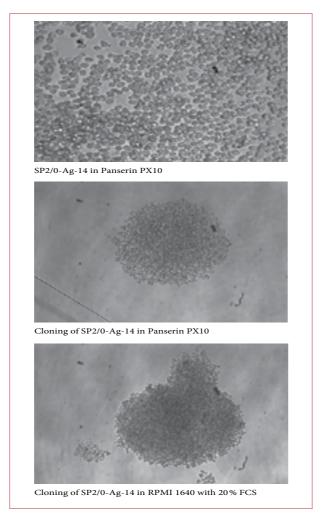
Special advantages

Panserin PX10 is a ready-to-use, serum-free medium for the production of monoclonal antibodies. It contains no undefined peptones or hydrolysates. Due to its optimized composition Panserin PX10 shows significant growth stimulation even at low seeding densities. In addition to an excellent cell growth Panserin PX10 shows very good cloning properties. Conventional serum-free systems often require long and laborious adaptation steps and seeding densities of up to 10⁵ cells/ml. In contrast, most clones can be directly transferred into Panserin PX10 culture. With Panserin PX10 clones can be obtained easily.

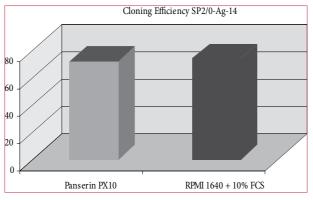


Typical growth curve of SP2/O-Ag-14 in Panserin PX10

Panserin PX10 ⁽¹⁾	500 ml	P04-710PX10
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Sp2/0-Ag-14 cells were transferred from serum containing culture (RPMI 1640 with 10 % FCS) directly into Panserin PX10. Seeding density 1.000 cells/ml. In comparison Sp2/0-Ag-14 in RPMI 1640 with 10 % FCS.



Instructions for use

www.pan-biotech.com

Detailed instructions will be provided with the accompanying datasheet for Panserin PX10. In addition, instructions for use can also be found at www.pan-biotech.com.

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Serum-free Media

Panserin PX40

Panserin PX40 is a ready-to-use complete medium for the serum-free cultivation of a variety of cells.

Composition

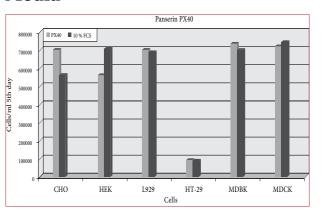
Based on RPMI 1640/DMEM/F-12, trace elements, albumin, lipoproteins, vitamins, hormones and attachment factors were added to the medium. The medium does not contain any growth factors.

Suitability

Cultivation of a variety of adherent cells under serum-free conditions (e. g. HEK, L929, CHO, MDCK, MDBK, 3T3A).

Special advantages

Panserin PX40 is a ready-to-use serum-free medium for the cultivation of a variety of adherent cells. The addition of attachment factors allows the cultivation of even highly demanding cells after a short adaptation phase.



Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin PX40. In addition, instructions for use can also be found at www.pan-biotech.com.

Panserin PX40 ⁽¹⁾	500 ml	P04-710PX40

) usually on stock, (2) minimum order 10 l, (3) available upon request



Spodopan

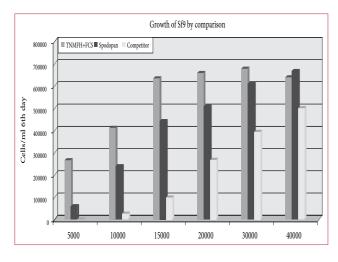
Spodopan is a protein-free medium for an optimized growth of insect cells such as Sf9 and Sf21 (Spodoptera frugiperda) in suspension culture. Insect cells are often used for the industrial production of recombinant proteins.

Composition

Spodopan contains amino acids, vitamins, salts, trace elements, lipids and growth promoting factors in a formulation optimized for insect cells. It contains no protein or any orther components of human or animal origin.

Suitability

Spodopan is suitable for the cultivation of insect cells and the production of recombinant proteins. (Baculovirus expression vector system, BEVS)

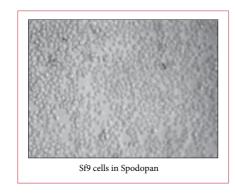


Special advantages

Spodopan with its protein-free formulation is free of human and animal components. This allows the production of recombinant proteins for medical and therapeutic purposes. The protein-free formulation also facilitates an easier and more economic purification of final products from the cell culture. Spodopan guarantees a high cell density with increased production of recombinant proteins (Baculovirus expression vector system).

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Spodopan. In addition, instructions for use can also be found at www.pan-biotech.com.



Cmadaman(1)	100 ml	P04-850100
Spodopan ⁽¹⁾	500 ml	P04-850500

Serum-free Media

Panserin S2

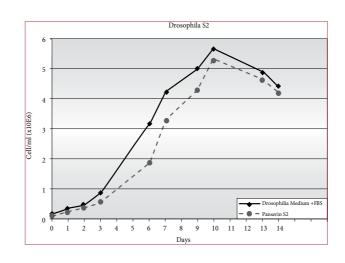
Panserin S2 is a protein-free medium for an optimized growth of insect Drosophila S2 cells in suspension culture. Insect cells are widely used for the industrial production of recombinant proteins.

Composition

Panserin S2 contains amino acids, vitamins, salts, trace elements, lipids and growth promoting factors in a formulation optimized for the growth of insect cells. It contains no protein or any further components of human or animal origin.

Suitability

Panserin S2 is suitable for the cultivation of Drosophila S2 cells and the production of recombinant protein. (e.g. Baculovirus expression vector system, BEVS)



Panserin S2 ⁽¹⁾	100 ml	P04-710210
Palisellii 52\	500 ml	P04-710200

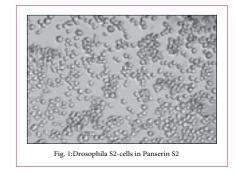
Special advantages

Panserin S2 with its protein-free formulation is free of human and animal components. This allows the production of recombinant proteins for medical and therapeutic purposes. The protein-free formulation also facilitates convenient and economic purification of final products from the cell culture. Panserin S2 guarantees a high cell density and viability resulting in an increased production and easy and economic purification of recombinant protein.

(Baculovirus expression vector system)

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Panserin S2. In addition, instructions for use can also be found at www.pan-biotech.com.









Endopan 300 SL

Endopan 300 SL is the first complete medium specially developed for the serum-free in vitro culture of human endothelial cells containing all components necessary for optimal growth.

Endothelial cells line blood and lymphatic vessels and the internal cavities of the heart. They display a strongly flattened, polygonal form and mostly rest on a basal membrane. They adhere to each other by desmosomes and tight-junctions. With a total cell number of about one trillion (10¹²), the endothelium is one of the biggest organs of the body and plays a key role in many physiological and patho-physiological processes (e.g. cell-based immune response, wound healing, inflammation, allergy, cardiovascular diseases, tumour growth). A huge number of soluble factors circulating in the blood or released by neighbouring cells control proliferation or apoptosis of endothelial cells and the invasion and migration of leucocytes to the endothelium, thereby regulating the maintenance, degeneration, or regeneration of blood vessels.

Composition and application

Endopan 300 SL ready-to-use is a complete medium specially developed for serum-free in vitro culture of human endothelial cells and it contains all components necessary for optimal growth. It is designed for use in an incubator at 37° C with a 5% CO2 atmosphere. Endopan 300 SL kit is provided with a serum substitute (Panexin SL-S) and supplements in separate sterile packing.

Endopan 300 SL has been designed for serum-free culture of endothelial cells directly after isolation. This exclusive medium is optimized for the maintenance and expansion of endothelial cells under serum-free culture conditions. HUVEC cultured in Endopan 300 SL exhibit a typical endothelial morphology and express endothelial specific markers such as CD31 or von Willebrand Factor and bind UEA-1 lectin. Additionally, HUVEC in Endopan 300 SL have been shown to maintain endothelial cell signal transduction pathways. When using complete Endopan 300 SL the growth rate of HUVEC is similar to that obtained for cells cultured in endothelial growth media containing bovine serum and supplements.

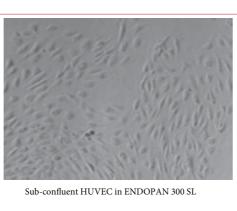
Although not extensively tested, it has been shown that Endopan 300 SL can also be used with endothelial cells of bovine, pig, rat, and rabbit origin.

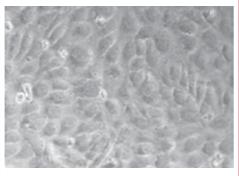
Endothelial cell biology has been greatly advanced by studying cultured vascular endothelial cells in vitro. Traditionally, complete endothelial growth media contain animal serum. The advance of so-called low-serum media for endothelial cells has improved the quality of experimental data acquired in recent years. However, endothelial cells may synthesize substances which can not be detected due to their low quantity or masking effects from serum.

In the past, cellular signalling pathways in endothelial cells have not been decipherable experimentally because even low concentrations of serum present in traditional media induce an undefined and undesired stimulation of cell surface receptors or intracellular signalling which only may become evident under serum-free conditions. As endothelial cells move into the field of interest for vascular tissue engineering with potential therapeutic application, the presence of whole animal serum is undesirable for such applications.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for Endopan 300 SL. In addition, instructions for use can also be found at www.pan-biotech.com.





Confluent HUVEC in ENDOPAN 300 SL

Endopan 300 SL ready-to-use(3)	500 ml	P04-00650
Endopan 300 SL kit ⁽³⁾	500 ml	P04-0065K

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Introduction

You only want highest quality media for cell cultivation?

At PAN-Biotech, perfect raw materials combined with state-of-the-art technologies guarantee a first-class quality of our media.

Water is the most important component of liquid media, which is why water purity is of outstanding importance for the quality of media. The water we use generally has a very low endotoxin level of < 0.005 EU/ml, and therefore is of highest purity.

Our media are placed in quarantine until quality control procedures are finished. This guarantees an excellent quality of the final product.

Advantages of our cell culture media

- Raw materials used are tested according to the highest possible quality standards
- Standard filling in sterile, high-class PET bottles
- Batches of 10 litre up to 4000 litre
- Custom service of product optimisation and further development for specific applications and purposes
- CE-label according to medical product law available upon request

Other sizes and custom formulation

Almost all media available from PAN-Biotech can be filled in special containers as per customer requirement. Besides standard bottles in 100, 500, and 1000 ml, medium can be filled in cans (up to 10 L), bags (up to 500 L), or other containers with fittings according to customer specifications for special applications such as continous feed process or production purposes.

Delivery time

Standard media:

In principle within 3 working days in Germany; otherwise we will inform you.

Special media and custom products:

Within Germany in 4 to 6 weeks after receipt of order.

Shelf life

Powder media	2 years
Liquid media without Glutamine	2 years
Liquid media with stable Glutamine	2 years
Liquid media with L-Glutamine	1 year

Liquid media with L-Glutamine can be used also after the expiry date, but have to be supplemented with new L-Glutamine in this case. Shelf life starts on date of production!

Storage

Powder media 2 – 8° C

Liquid media 2 – 8° C protected from light

Benefit from the experience and know-how of PAN-Biotech. Our state-of-the-art production facilities, with a production line specifically installed for these requirements, allow us to produce the formulations especially developed for your needs in constant high quality also for longer periods of time, and to make batch sizes adapted to your need. Our team of scientists will be pleased to advise you regarding your proprietary formulation.

For further information regarding the dependency of ph-values in media on the ${\rm CO_2}$ concentration in the incubator please refer to our website at www.pan-biotech.com.





Alpha MEM

Media

Description

Alpha MEM is a different formulation of MEM Eagle and contains a higher concentration of amino acids. It also has a higher concentration of lipoic acid, vitamins and pyruvate. Primarily it was developed for the cultivation of hamster kidney cells, but today it is used for a broad range of mammalian cells. Among others the alpha MEM promotes the growth and progeny of bone marrow cells in suspension culture and monolayer. A further possibility is the use as a separation medium or for the out-breeding of amniotic cells.

Liquid Media

Alpha MEM Eagle⁽¹⁾
without L-Glutamine
without Ribonucleosides
without Deoxyribonucleosides
with 2.2 g/L NaHCO₃ 500 ml P04-21050

Alpha MEM Eagle⁽¹⁾
with L-Glutamine
with Ribonucleosides
with Deoxyribonucleosides
with 2.2 g/L NaHCO₃ 500 ml P04-21500

Alpha MEM Eagle⁽¹⁾
with stab. Glutamin
with Ribonucleosides
with Deoxyribonucleosides
with 2.2 g/L NaHCO₃ 500 ml P04-21250

Special Media

Alpha MEM Eagle⁽²⁾
without L-Glutamine
with Ribonucleosides
with Deoxyribonucleosides
with 2.2 g/L NaHCO₂ 500 ml P04-21150

Alpha MEM Eagle⁽²⁾
with L-Glutamine
without Ribonucleosides
without Deoxyribonucleosides
with 2.2 g/L NaHCO₂ 500 ml P04-21060

Alpha MEM Eagle ⁽²⁾
with stab. Glutamine
without Ribonucleosids
without Deoxyribonucleosids
with 2.2 g/L NaHCO, 500 ml P04-21350

Alpha MEM Eagle ⁽²⁾
with L-Glutamine
without Glucose
with Ribonucleosides
with Deoxyribonucleosides
with 2.2 g/L NaHCO₂ 500 ml P04-21502

Alpha MEM Eagle (2)
without Glutamine
without Phenol red
without Ribonucleosides
without Deoxyribonucleosides
with 2.2 g/L NaHCO₃ 500 ml P04-21051

Alpha MEM

Media

Powder Media

Alpha MEM Eagle(1) 10 L P03-2410 without L-Glutamine 50 L P03-2450 with Ribonucleosides with Deoxyribonucleosides without NaHCO, Alpha MEM Eagle(1) 10 L P03-2510 with L-Glutamine 50 L P03-2550 with Ribonucleosides with Deoxyribonucleosides without NaHCO3 Alpha MEM Eagle(1) 10 L P03-2310 with L-Glutamine 50 L P03-2350 without Ribonucleosides without Deoxyribonucleosides with 2.2 g/L NaHCO₃

Alpha MEM Eagle⁽¹⁾

with L-Glutamine

with 25 mM Hepes

with Ribonucleosides

with Deoxyribonucleosides

without NaHCO₃

10 L P03-2610

50 L P03-2650

Composition

	Components	mg/L
Inorganic	Calcium chloride x 2H,O	264.92
Salts	Magnesium sulfate dried	139.52
	Potassium chloride	400.00
	Sodium chloride	6,800.00
	Sodium dihydrogen	140.00
	phosphate x H ₂ O	
Other	D(+)-Glucose anhydrous	1,000.00
Compo-	Hepes	5,958.00
nents	Lipoic acid	0.20
	Phenol red	10.00
	Sodium pyruvate	110.00
Amino	L-Alanine	25.00
Acids	L-Arginine x HCl	126.64
	L-Asparagine x H ₂ O	50.00
	L-Aspartic acid	30.00
	L-Cysteine x HCl x H ₂ O	100.00
	L-Cystine ²	24.00
	L-Glutamine	292.00
	L-Glutamic acid	75.00
	Glycine	50.00
	L-Histidine x HCl x H ₂ O	42.00
	L-Isoleucine	52.40
	L-Leucine	52.40
	L-Lysine x HCl	72.47
	L-Methionine	15.00
	L-Phenylalanine	32.00
	L-Proline	40.00
	L-Serine	25.00
	L-Threonine	48.00
	L-Tryptophan	10.00
	L-Tyrosine	36.20
	L-Valine	46.00
Vitamins	L-Ascorbic acid	50.00
, 100111110	D(+)-Biotin	0.10
	D-Calcium pantothenate	1.00
	Choline chloride	1.00
	Folic acid	1.00
	myo-Inositol	2.00
	Nicotinamide	1.00
	Pyridoxal x HCl	1.00
	Riboflavin	0.10
	Thiamine x HCl	1.00
	Vitamine B12	1.33
Ribonu-	Adenosine	10.00
cleosides	Cytidine	10.00
Sicosiacs	Guanosine	10.00
	Uridine	10.00
Deoxy-	2'-Deoxyadenosine x H ₂ O	10.00
ribonu-	2'-Deoxycytidine x HCl	11.00
cleosides	2'-Deoxyguanosine	10.00
21COOIUCO	2'-Deoxyguanosine	10.00
	2 DOONY HIS HILLING	10.00

1) usually on stock, (2) minimum order 10 l, (3) available upon reques



(1) usually on stock, (2) minimum order 10 l, (3) available upon requ



BME with Hank's Salts

Description

In the fifties of the last century it was found that mammalian cells do not only need the 10 essential amino acids, but also cystine, tyrosine and glutamine. In addition to these three amino acids BME includes

Composition

Components	mg/L
Calcium chloride x 2H ₂ O	185.44
Magnesium sulfate dried	139.52
Potassium chloride	400.00
Potassium dihydrogen	60.00
phosphate anhydrous	
Sodium chloride	8,000.00
di-Sodium hydrogen	47.88
phosphate	
D(+)-Glucose anhydrous	1,000.00
Hepes	5,958.00
Phenol red	10.00
L-Arginine x HCl	21.00
L-Cystine	12.00
L-Glutamine	292.00
L-Histidine Base	8.00
L-Isoleucine	26.00
L-Leucine	26.00
L-Lysine x HCl	36.47
L-Methionine	7.50
L-Phenylalanine	16.50
L-Threonine	24.00
L-Tryptophan	4.00
L-Tyrosine	18.00
L-Valine	23.50
D(+)-Biotin	1.00
D-Calcium pantothenate	1.00
Choline chloride	1.00
Folic acid	1.00
myo-Inositol	2.00
Nicotinamide	1.00
Pyridoxal x HCl	1.00
Riboflavin	0.10
Thiamine x HCl	1.00
	Calcium chloride x 2H ₂ O Magnesium sulfate dried Potassium chloride Potassium dihydrogen phosphate anhydrous Sodium chloride di-Sodium hydrogen phosphate D(+)-Glucose anhydrous Hepes Phenol red L-Arginine x HCl L-Cystine L-Glutamine L-Histidine Base L-Isoleucine L-Leucine L-Lysine x HCl L-Methionine L-Phenylalanine L-Threonine L-Tryrosine L-Tyrosine L-Valine D(+)-Biotin D-Calcium pantothenate Choline chloride Folic acid myo-Inositol Nicotinamide Pyridoxal x HCl Riboflavin

When 5,958.00 mg/L HEPES is included there is only 7,500.00 mg/L sodium chloride.

Liquid Media

BME with HBSS⁽¹⁾ without L-Glutamine with 0.35 g/L NaHCO₃ 500 ml P04-26050

BME with Earle's Salts

also eight B-vitamins. Originally BME was used for the cultivation of murine L-cells and HeLa cells. With its many variations it is used in many fields of science today. Along with the cultivation of normal mammalian cells BME is very suitable for transformed cells.

Composition

0.1: 11:1 211.0	
Calcium chloride x 2H,O	264.92
Magnesium sulfate dried	139.52
Potassium chloride	400.00
Sodium chloride	6,800.00
Sodium dihydrogen	140.00
phosphate x H ₂ O	
D(+)-Glucose anhydrous	1,000.00
Hepes	5,958.00
Phenol red	10.00
L-Arginine x HCl	21.00
L-Cystine	12.00
L-Glutamine	292.00
L-Histidine Base	8.00
L-Isoleucine	26.00
L-Leucine	26.00
L-Lysine x HCl	36.47
L-Methionine	7.50
L-Phenylalanine	16.50
L-Threonine	24.00
	4.00
•	18.00
L-Valine	23.50
D(+)-Biotin	1.00
D-Calcium pantothenate	1.00
Choline chloride	1.00
Folic acid	1.00
myo-Inositol	2.00
Nicotinamide	1.00
Pyridoxal x HCl	1.00
Riboflavin	0.10
Thiamine x HCl mg/L HEPES is included there is only 6.	1.00
	Potassium chloride Sodium chloride Sodium dihydrogen phosphate x H ₂ O D(+)-Glucose anhydrous Hepes Phenol red L-Arginine x HCl L-Cystine L-Glutamine L-Histidine Base L-Isoleucine L-Leucine L-Lysine x HCl L-Methionine L-Phenylalanine L-Threonine L-Tryptophan L-Tyrosine L-Valine D(+)-Biotin D-Calcium pantothenate Choline chloride Folic acid myo-Inositol Nicotinamide Pyridoxal x HCl Riboflavin

sodium chloride.

Liquid Media

BME with EBSS⁽¹⁾ without L-Glutamine with 2.2 g/L NaHCO₃

500 ml P04-25050

Special Media

BME with EBSS(2) with L-Glutamine with 2.2 g/L NaHCO₃

500 ml P04-25500



CMRL-1066 Medium

Description

The CMRL is a nucleosid- and vitamin-rich medium. In the past it was developed to clone monkey-kidney cells and as long time culture medium for L-cells. It is suitable for many types of human and monkey cells and also for other mammalian cells, especially by using horse and calf serum.

Liquid Media

CMRL - 1066(1) without L-Glutamine without Phenol red with 2.2 g/L NaHCO₃ 500 ml P04-84600

Special Media

CMRL - 1066⁽²⁾ with L-Glutamine without Phenol red with 2.2 g/L NaHCO,

500 ml P04-84500

Commonition

Composition			
Components mg/l			
Inorganic	Calcium chloride x 2H,O	264.92	
Salts	Potassium chloride	400.00	
	Magnesium sulfate dried	139.55	
	Sodium acetate anhydrous	50.00	
	Sodium chloride	6,799.00	
	di-Sodium hydrogen	139.75	
	phosphate x H ₂ O		
Other	Cholesterol	0.20	
Compo-	D(+)-Glucose anhydrous	1,000.00	
nents	Glutathione (red.)	10.00	
	Sodium glucuronate x H ₂ O	4.00	
	Tween 80	5.00	
Coenzy-	Cocarboxylase x HCl	1.00	
me	Coenzyme A Trilithiumsalt x 2H ₂ O	2.60	
	NAD	7.00	
	NADP sodium salt	1.00	
	UTP	1.0	
Amino	L-Alanine	25.0	
Acids	L-Arginine x HCl	70.0	
	L-Aspartic acid	30.0	
	L-Cysteine x HCl x H ₂ O	260.0	
	L-Cystine	20.0	
	L-Glutamine	100.0	
	L-Glutamic acid	75.0	
	Glycine	50.0	
	L-Histidine x HCl x H ₂ O	20.0	
	L-Hydroxyproline	10.0	
	L-Isoleucine	20.0	
	L-Leucine	60.0	
	L-Lysine x HCl	70.0	
	L-Methionine	15.0	
	L-Phenylalanine	25.0	
	L-Proline	40.0	
	L-Serine	25.0	
	L-Threonine	30.0	
	L-Tryptophan	10.0	
	L-Tyrosine	40.0	
	L-Valine	25.0	
Vitamins	L-Ascorbic acid	50.0	
	D(+)-Biotin	0.10	
	D-Calcium pantothenate	1.0	
	Choline chloride	1.0	
	Folic acid	1.0	
	myo-Inositol	2.0	
	Nicotinamide	1.0	
	Pyridoxal x HCl	1.0	
	Riboflavin	0.10	
	Thiamine x HCl	1.0	
_	Vitamine B12	1.3	
Deoxy-	2'-Deoxyadenosine x H ₂ O	10.0	
ribonu-	2'-Deoxycytidine x HCl	11.0	
cleosides	2'-Deoxyguanosine	10.0	
	2'-Deoxythymidine	10.0	

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Dulbecco's Modified Eagle Medium

Description

Intrinsically developed for the cultivation of murine embryonic cells, DMEM is tailor-made for the cultivation of a broad range of cells, especially if the medium is supplemented with FBS. DMEM is an Eagle medium modification with a four-fold content of amino acids and vitamins. DMEM with 1.0 g/L Glucose is the standard medium, whereas DMEM with 4.5 g/L Glucose is for cells which have a high energy demand.

Liquid Media without Glucose

DMEM without Glucose(1) without L-Glutamine without Sodium pyruvate with 3.7 g/L NaHCO, 500 ml P04-01548S1

DMEM without Glucose⁽¹⁾ without L-Glutamine with Sodium pyruvate

with 3.7 g/L NaHCO₃ 500 ml P04-01549

DMEM without Glucose⁽¹⁾ without L-Glutamine without Sodium pyruvate without Phenol red with 3.7 g/L NaHCO₃ 500 ml P04-01548

Special Media without Glucose

DMEM without Glucose⁽²⁾ with L-Glutamine with Sodium pyruvate

with 3.7 g/L NaHCO, 500 ml P04-01551

Powder Media without Glucose

DMEM without Glucose(1) without L-Glutamine without Sodium pyruvate

without Phenol red P03-0010 10 L without NaHCO, 50 L P03-0050

Liquid Media with 1.0 g/L Glucose

DMEM with 1.0 g/L Glucose(1) without L-Glutamine with Sodium pyruvate with 3.7 g/L NaHCO₃ 500 ml P04-01500

DMEM with 1.0 g/L Glucose⁽¹⁾

with L-Glutamine with Sodium pyruvate

with 3.7 g/L NaHCO₃ 500 ml P04-01550

DMEM with 1.0 g/L Glucose⁽¹⁾ with stab. Glutamine with Sodium pyruvate

with 3.7 g/L NaHCO₃ 500 ml P04-02500

DMEM with 1.0 g/L Glucose(1) with L-Glutamine with Sodium pyruvate without Phenol red

with 3.7 g/L NaHCO, 500 ml P04-01515

DMEM with 1.0 g/L Glucose⁽¹⁾ with L-Glutamine with Sodium pyruvate with 25 mM Hepes

with 3.7 g/L NaHCO, 500 ml P04-05551

Powder Media with 1.0 g/L Glucose

DMEM with 1.0 g/L Glucose(1) with L-Glutamine

with Sodium pyruvate P03-0510 10 L with 3.7 g/L NaHCO, 50 L P03-0550

DMEM with 1.0 g/L Glucose⁽¹⁾ with L-Glutamine with Sodium pyruvate

without Phenol red P03-01510 10 L with 3.7 g/L NaHCO, 50 L P03-01550

Dulbecco's Modified Eagle Medium

Media

Special Media with 1.0 g/L Glucose

DMEM with 1.0 g/L Glucose⁽²⁾ without L-Glutamine with Sodium pyruvate without Calcium

with 3.7 g/L NaHCO. 500 ml P04-01501

SILAC-DMEM⁽²⁾ with 1.0 g/L Glucose with stab. Glutamine with Sodium pyruvate without L-Arginin without L-Lysin

with 3.7 g/L NaHCO, 500 ml P04-02501

DMEM with 1.0 g/L Glucose⁽²⁾ without L-Glutamine without Sodium pyruvate without Phenol red

with 3.7 g/L NaHCO₃ 500 ml P04-03556

DMEM with 1.0 g/L Glucose⁽²⁾ without L-Glutamine with Sodium pyruvate without Phenol red 500 ml P04-01159

with 3.7 g/L NaHCO,

DMEM with 1.0 g/L Glucose(2) with stab. Glutamine with Sodium pyruvate without Phenol red with 3.7 g/L NaHCO₃ 500 ml P04-02500S1

DMEM with 1.0 g/L Glucose⁽²⁾ with L-Glutamine with Sodium pyruvate without Phenol red with 3.7 g/L NaHCO, 500 ml P04-01516

DMEM with 1.0 g/L Glucose(2) with L-Glutamine without Sodium pyruvate

with 3.7 g/L NaHCO. 500 ml P04-01555

Liquid Media with 4.5 g/L Glucose

DMEM with 4.5 g/L Glucose⁽¹⁾ without L-Glutamine without Sodium pyruvate

with 3.7 g/L NaHCO₃ 500 ml P04-03500

DMEM with 4.5 g/L Glucose(1) without L-Glutamine with Sodium pyruvate with 3.7 g/L NaHCO, 500 ml P04-03600

DMEM with 4.5 g/L Glucose⁽¹⁾ with L-Glutamine without Sodium pyruvate with 3.7 g/L NaHCO, 500 ml P04-03550

DMEM with 4.5 g/L Glucose⁽¹⁾ with L-Glutamine with Sodium pyruvate with 3.7 g/L NaHCO,

500 ml P04-03590

DMEM with 4.5 g/L Glucose(1) with L-Glutamine

with Sodium pyruvate with 1.5 g/L NaHCO3 500 ml P04-03596

DMEM with 4.5 g/L Glucose⁽¹⁾ with stab. Glutamine without Sodium pyruvate with 3.7 g/L NaHCO, 500 ml P04-04500

DMEM with 4.5 g/L Glucose⁽¹⁾ with stab. Glutamine with Sodium pyruvate with 3.7 g/L NaHCO, 500 ml P04-04510

DMEM with 4.5 g/L Glucose(1) without L-Glutamine without Sodium pyruvate without Phenol red with 3.7 g/L NaHCO, 500 ml P04-01161

DMEM with 4.5 g/L Glucose⁽¹⁾ without Glutamine with Sodium pyruvate without Phenol red with 3.7 g/L NaHCO, 500 ml P04-01158

DMEM with 4.5 g/L Glucose⁽¹⁾ with L-Glutamine with Sodium pyruvate without Phenol red 500 ml P04-03591 with 3.7 g/L NaHCO₃

(1) usually on stock, (2) minimum order 10 l, (3) available upon request (1) usually on stock, (2) minimum order 10 l, (3) available upon request





www.pan-biotech.com

Dulbecco's Modified Eagle Medium

DMEM with 4.5 g/L Glucose⁽¹⁾ with L-Glutamine with 25 mM Hepes without Sodium pyruvate with 3.7 g/L NaHCO₃ 500 ml P04-05540

DMEM with 4.5 g/L Glucose⁽¹⁾ with L-Glutamine with 25 mM Hepes with Sodium pyruvate

500 ml P04-05550 with 3.7 g/L NaHCO₃

Special Media with 4.5 g/L Glucose

DMEM with 4.5 g/L Glucose⁽²⁾ with stab. L-Glutamine with 25 mM Hepes without Sodium pyruvate

with 3.7 g/L NaHCO, 500 ml P04-04550

DMEM with 4.5 g/L Glucose⁽²⁾ with L-Glutamin without Sodium pyruvate with 25 mM Hepes without Phenol red

with 3.7 g/L NaHCO₃ 500 ml P04-05545

DMEM with 4.5 g/L Glucose⁽²⁾ with stab. Glutamine with Sodium pyruvate without Phenol red

with 3.7 g/L NaHCO₃ 500 ml P04-03588

DMEM with 4.5 g/L Glucose⁽²⁾ without L-Glutamine with 25 mM Hepes with Sodium pyruvate with 3.7 g/L NaHCO, 500 ml P04-01597 DMEM $(10 \text{ x})^{(3)}$ with 4.5 g/L Glucose without L-Glutamine without Sodium pyruvate with NEAA

without NaHCO3 500 ml P04-03510

DMEM with 4.5 g/L Glucose⁽²⁾ without L-Glutamine with Sodium pyruvate without L-Arginine with 3.7 g/L NaHCO, 500 ml P04-03598

DMEM with 4.5 g/L Glucose⁽²⁾ with L-Glutamine with 25 mM Hepes without Sodium pyruvate with 2.2 g/L NaHCO₃ 500 ml P04-04057

DMEM with 4.5 g/L Glucose⁽²⁾ with L-Glutamine without Sodium pyruvate without Sodium chloride without NaHCO3 500 ml P04-03560

DMEM with 5.5 g/L Glucose⁽²⁾ with L-Glutmamin without Sodium pyruvate with 3.7 g/L NaHCO₃ 500 ml P04-03551

DMEM with 4.5 g/L Glucose⁽²⁾ with stab. Glutamine with Sodium pyruvate with 25 mM Hepes without Phenol red with 0.5 g/L NaHCO3 500 ml P04-01163

DMEM with 4.5 g/L Glucose⁽²⁾ with L-Glutamine without Sodium pyruvate without Isoleucine with 3.7 g/L NaHCO₃ 500 ml P04-03503

Dulbecco's Modified Eagle Medium

Media

Powder Media with 4.5 g/L Glucose

DMEM with 4.5 g/L Glucose⁽¹⁾ without L-Glutamine without Sodium pyruvate 10 L P03-6510 without NaHCO, 50 L P03-6550 DMEM with 4.5 g/L Glucose⁽¹⁾ with L-Glutamine without Sodium pyruvate 10 L P03-0710 without NaHCO, 50 L P03-0750 DMEM with 4.5 g/L Glucose⁽¹⁾ with L-Glutamine with Sodium pyruvate 10 L P03-0810 without NaHCO3 P03-0850 50 L DMEM with 4.5 g/L Glucose(1) with L-Glutamine without Sodium pyruvate with 25 mM Hepes 10 L P03-0910 without NaHCO, 50 L P03-0950 DMEM with 4.5 g/L Glucose⁽¹⁾ with L-Glutamine with Sodium pyruvate with 25 mM Hepes 10 L P03-1010

50 L

P03-1050

without NaHCO,

Composition

		/т
	Components	mg/L
Inorganic	Calcium chloride anhydrous	200.00
Salts	Iron(III) nitrate x 9H ₂ O	0.10
	Magnesium sulfate anhydrous	97.66
	Potassium chloride	400.00
	Sodium chloride	6,400.00
	Sodium dihydrogen phosphate	108.69
	anhydrous	
Other	D(+)-Glucose anhydrous	4,500.00
Compo-	Hepes	5,958.00
nents	Phenol red	15.00
	Sodium pyruvate	110.00
Amino	L-Arginine x HCl	84.00
Acids	L-Cystine x 2HCl	62.58
	L-Glutamine	584.00
	Glycine	30.00
	L-Histidine x HCl x H,O	42.00
	L-Isoleucine 2	104.80
	L-Leucine	104.80
	L-Lysine x HCl	146.20
	L-Methionine	30.00
	L-Phenylalanine	66.00
	L-Serine	42.00
	L-Threonine	95.20
	L-Tryptophan	16.00
	L-Tyrosine x 2Na	103.79
	L-Valine	93.60
Vitamins	D-Calcium pantothenate	4.00
	Choline chloride	4.00
	Folic acid	4.00
	myo-Inositol	7.00
	Nicotinamide	4.00
	Pyridoxine x HCl	4.00
	Riboflavin	0.40
	Thiamine x HCl	4.00
When 5,957.50	mg/L HEPES is included there is only 5,4	00.00 mg/L

sodium chloride.

(1) usually on stock, (2) minimum order 10 l, (3) available upon request





Powder Media

DMEM/F12

Description

This medium supports the growth of almost all cell lines. For example, it is used for pancreas cells, Sertoli cells or to culture cells, which are used for human protein production. It combines the advantages of both media DMEM (high concentration of amino acids and vitamins) and Ham's F12 (higher concentration of zinc sulphate, putrescine and linoleic acid).

Composition

	Components	mg/L
Inorganic	Calcium chloride x 2H ₂ O	154.45
Salts	Iron(III)-nitrate x 9H,O	0.05
	Iron(II)-sulfate x 7H,O	0.42
	Potassium chloride	311.83
	Copper(II)-sulfate x 5H ₂ O	0.001
	Magnesium chloride	28.57
	Magnesium sulfate	48.85
	Sodium chloride	6,999.50
	Sodium dihydrogen phosphate	54.35
	di-Sodium hydrogen phosphate	70.98
	Zinc sulfate x 7H,O	0.43
Other	D(+)-Glucose anhydrous	3,151.0
Compo-	Hypoxanthine	2.04
nents	Linoleic acid	0.04
	DL-68-Lipoic acid	0.103
	Sodium pyruvate	110.00
	Phenol red	8.10
	Putrescin x 2HCl	0.081
	Thymidine	0.36
Amino	L-Alanine	4.45
Acids	L-Arginine x HCl	147.35
	L-Asparagine x H ₂ O	7.50
	L-Aspartic acid	6.65
	L-Cystine x 2HCl	31.29
	L-Cysteine x HCl x H ₂ O	17.56
	L-Glutamine	365.00
	L-Glutamic acid	7.35
	Glycine	18.75
	L-Histidine x HCl x H ₂ O	31.48
	L-Isoleucine	54.37
	L-Leucine	58.96
	L-Lysine x HCl	91.37
	L-Methionine	17.24
	L-Phenylalanine	35.48
	L-Proline	17.27
	L-Serine	26.25
	L-Threonine	53.55
	L-Tryptophan	9.02
	L-Tyrosine x 2Na x 2H ₂ O	55.81
	L-Valine	53.00
Vitamins	D-(+)-Biotine	0.004
	D-Calcium pantothenate	2.12
	Cholin chloride	8.98
	Folic acid	2.66
	myo-Inositol	12.51
	Nicotinamide	2.02
	Pyridoxine x HCl	2.03
	Riboflavin	0.22
	Thiamine x HCl	2.17
	Vitamine B12	0.68

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Media

DMEM/F12 (1:1) ⁽¹⁾ without L-Glutamine with 1.2 g/L NaHCO ₃	500 ml P04-41450	DMEM/F12 (1:1) ⁽¹⁾ without L-Glutamine without NaHCO ₃	10 L 50 L	P03-6010 P03-6050
DMEM/F12 (1:1) ⁽¹⁾ with L-Glutamine with 1.2 g/L NaHCO ₃	500 ml P04-41500	DMEM/F12 (1:1) ⁽¹⁾ with L-Glutamine without NaHCO ₃	10 L 50 L	P03-1110 P03-1150
DMEM/F12 (1:1) ⁽¹⁾ with stab. Glutamine with 1.2 g/L NaHCO ₃	500 ml P04-41150	DMEM/F12 (1:1) ⁽¹⁾ with L-Glutamine with 15 mM Hepes without NaHCO ₃	10 L 50 L	P03-6110 P03-6150
DMEM/F12 (1:1) ⁽¹⁾ without L-Glutamine with 15 mM Hepes with 1.2 g/L NaHCO ₃	500 ml P04-41550	DMEM/F12 (1:1) ⁽¹⁾ with L-Glutamine with 25 mM Hepes	10 L	P03-1210
DMEM/F12 (1:1) ⁽¹⁾ with L-Glutamine with 15 mM Hepes		without NaHCO ₃	50 L	P03-1250

www.pan-biotech.com

(1) usually on stock, (2) minimum order 10 l, (3) available upon request

Liquid Media

with 1.2 g/L NaHCO₃

DMEM/F12 (1:1)⁽²⁾ without L-Glutamine without Glucose with 1.2 g/L NaHCO3

DMEM/F12 (1:1)⁽²⁾ with L-Glutamine with 25 mM Hepes with 1.2 g/L NaHCO₃

DMEM/F12 (1:1)⁽²⁾ with L-Glutamine without Phenol red with 1.2 g/L NaHCO₃

DMEM/F12 (1:1)⁽²⁾ with stab. Glutamine with 15 mM Hepes without Calcium chloride with 1.2 g/L NaHCO₃

Special Media

500 ml P04-41250

500 ml P04-41151

500 ml P04-41252

500 ml P04-41650

500 ml P04-41251



Glasgow MEM (BHK 21)

Description

The GMEM was developed as a modification of BME to culture primary baby hamster kidney cells. This version has twice the concentration of vitamins and amino acids.

Liquid Media

Diquia Media		
Glasgow-MEM (BHK 21) ⁽¹⁾ without L-Glutamine without Tryptose phosphate with 2.75 g/L NaHCO ₃	500 ml	P04-97500
Glasgow-MEM (BHK 21) ⁽¹⁾ with L-Glutamine with Tryptose phosphate with 2.75 g/L NaHCO ₃	500 ml	P04-96500
Glasgow-MEM (BHK 21) ⁽²⁾ without L-Glutamine with Tryptose phosphate with 2.75 g/L NaHCO ₃ Powder Media	500 ml	P04-98500
Glasgow-MEM (BHK 21) ⁽¹⁾ without L-Glutamine without Tryptose phosphate without NaHCO ₃	10 L 50 L	P03-3110 P03-3150

Glasgow-MEM (BHK 21) (1)

Glasgow-MEM (BHK 21)⁽¹⁾

without Tryptose phosphate

with L-Glutamine with Tryptose phosphate

without NaHCO3

with L-Glutamine

without NaHCO3

10 L

50 L

10 L

50 L

P03-6910

P03-6950

P03-6810

P03-6850

Composition

		,	• . 1
	Components	w/o	with
		Tryptose	Trytose
		Phosph-	Phosph-
		ate mg/L	ate mg/L
Inor-	Calcium chloride x 2H ₂ O	264.92	238.43
ganic	Iron(III) nitrate x 9H ₂ O	0.10	0.09
Salts	Magnesium sulfate dried	139.57	125.64
	Potassium chloride	400.00	360.00
	Sodium chloride	6,400.00	6,260.00
	di-Sodium hydrogen	0.00	250.00
	phosphate		
	Sodium dihydrogen	124.00	111.60
	phosphate x H ₂ O		
Other	D(+)-Glucose anhydrous	4,500.00	4,250.00
Com-	Phenol red	15.00	13.50
po-	Pepton from Casein	0.00	1,000.00
nents	Pepton from meat	0.00	500.00
	Yeast extract	0.00	500.00
Ami-	L-Arginine x HCl	42.00	37.80
no	L-Cystine	24.00	21.60
Acids	L-Glutamine	292.00	262.80
	L-Histidine x HCl x H ₂ O	21.00	18.90
	L-Isoleucine 2	52.40	47.16
	L-Leucine	52.40	47.16
	L-Lysine x HCl	73.10	65.79
	L-Methionine	15.00	13.50
	L-Phenylalanine	33.00	29.70
	L-Threonine	47.60	42.84
	L-Tryptophan	8.00	7.20
	L-Tyrosine	36.20	32.52
	L-Valine	46.80	42.12
Vita-	D-Calcium pantothenate	2.00	1.80
mins	Choline chloride	2.00	1.80
	Folic acid	2.00	1.80
	myo-Inositol	3.60	3.24
	Nicotinamide	2.00	1.80
	Pyridoxal x HCl	2.00	1.80
	Riboflavin	0.20	0.18

Thiamine x HCl

Grace's Insect Medium

www.pan-biotech.com

Description

The Grace's Insect Medium was originally developed to culture insect cells including SF9 and SF21 cells. Moreover it supports a broad range of lepidopteran cells.

Special Media

Grace's Insect Medium(2) without L-Glutamine with 0.35 g/L NaHCO₃

500 ml P04-81500

Grace's Insect Medium(2) with L-Glutamine

with 0.35 g/L NaHCO₃ 500 ml P04-82500

Powder Media

Grace's Insect Medium(1) without L-Glutamine P03-9010 10 L without NaHCO3 50 L P03-9050 Grace's Insect Medium(1)

with L-Glutamine 10 L P03-9110 without NaHCO3 50 L P03-9150

Composition

	Components	mg/L
Inorganic	Calcium chloride x 2H ₂ O	1,324.6
Salts	Potassium chloride	2,240.00
	Magnesium chloride x 6H ₂ O	2,278.86
	Magnesium sulfate dried	1,939.81
	di-Sodium hydrogen	876.92
	phosphate	
Other	DL-Malic acid	670.00
Compo-	Succinic acid	60.00
nents	Fructose	400.00
	Fumaric acid	55.00
	D(+)-Glucose anhydrous	700.00
	α-Ketoglutaric acid sodium	425.66
	salt	
	D-Sucrose	26,680.00
Amino	ß-Alanine	200.00
Acids	L-Alanine	200.00
	L-Arginine x HCl	700.00
	L-Asparagine x H ₂ O	350.00
	L-Aspartic acid	350.00
	L-Cystine	19.18
	L-Glutamine	600.00
	L-Glutamic acid	600.00
	Glycine	650.00
	L-Histidine Base	2,500.00
	L-Isoleucine	50.00
	L-Leucine	75.00
	L-Lysine x HCl	625.00
	L-Methionine	50.00
	L-Phenylalanine	150.00
	L-Proline	350.00
	L-Serine	550.00
	L-Threonine	175.00
	L-Tryptophan	100.00
	L-Tyrosine	50.00
	L-Valine	100.00
Vitamins	p-Aminobenzoic acid	0.02
	D(+)-Biotin	0.01
	D-Ca-Pantothenate	0.02
	Choline chloride	0.20
	Folic acid	0.02
	myo-Inositol	0.02
	Niacin	0.02
	Pyridoxine x HCl	0.02
	Riboflavin	0.02
	Thiamine x HCl	0.02

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



2.00

1.80



Ham's F10 Medium

Description

Ham's F10 is an alternative to Ham's F12 and it was used primarily to culture CHO-cells. Today, Ham's F10 can be used with or without FBS for many different cell cultures. It is used for example for primary cells of rat and chicken, but also for human diploid cells.

Liquid Media

Ham's F10 Medium ⁽¹⁾ with L-Glutamine with 1.2 g/L NaHCO ₃	500 ml	P04-12500
Special Media		
Ham's F10 Medium ⁽²⁾ without L-Glutamine with 1.2 g/L NaHCO ₃	500 ml	P04-12050
Ham's F10 Medium ⁽²⁾ with stab. Glutamine with 1.2 g/L NaHCO ₃	500 ml	P04-13500
Ham's F10 Medium ⁽²⁾ with L-Glutamine with 25 mM Hepes with 1.2 g/L NaHCO ₃	500 ml	P04-13050
Ham's F10 Medium ⁽²⁾ without L-Glutamine without Phenol red with 1.2 g/L NaHCO ₃	500 ml	P04-12049
Powder Media		
Ham's F-10 Medium ⁽¹⁾ without L-Glutamine without NaHCO ₃	10 L 50 L	P03-5010 P03-5050
Ham's F-10 Medium ⁽¹⁾ with L-Glutamine without NaHCO ₃	10 L 50 L	P03-3910 P03-3950
Ham's F-10 Medium ⁽¹⁾ with L-Glutamine with 25 mM Hepes without NaHCO ₃	10 L 50 L	P03-4010 P03-4050

Composition

	Components	ma/I
	Components	mg/L
Inorganic	Calcium chloride x 2H ₂ O	44.09
Salts	Copper(II) sulfate x 5H ₂ O	0.003
	Iron(II) sulfate x 7H ₂ O	0.834
	Magnesium sulfate dried	106.06
	Potassium chloride	285.00
	Potassium dihydrogen	83.00
	phosphate	7400.00
	Sodium chloride	7400.00
	di-Sodium hydrogen	153.70
	phosphate	0.020
	Zinc sulfate x 7H ₂ O	0.029
Other	D(+)-Glucose anhydrous	1100.00
Compo-	Hypoxanthine	4.08
nents	DL-α-Lipoic acid	0.21
	Hepes	5958.00
	Phenol red	1.20
	Sodium pyruvate	110.00
	2'- Deoxythymidine	0.73
Amino	L-Alanine	8.91
Acids	L-Arginine x HCl	211.00
	L-Asparagine x H ₂ O	15.00
	L-Aspartic acid	13.30
	L-Cysteine x HCl x H ₂ O	35.12
	L-Glutamine	146.20
	L-Glutamic acid	14,70
	Glycine	7.51
	L-Histidine x HCl x H ₂ O	21.00
	L-Isoleucine	2.60
	L-Leucine	13.10
	L-Lysine x HCl	29.30
	L-Methionine	4.48
	L-Phenylalanine	4.96
	L-Proline L-Serine	11.50
		10.50 3.57
	L-Threonine	0.60
	L-Tryptophan L-Tyrosine	1.81
	L-Valine	3.50
X7:4		
Vitamins	D(+)-Biotin	0.024
	D-Calcium pantothenate	0.715
	Choline chloride	0.698
	Folic acid	1.32
	myo-Inositol Nicotinamide	0.541 0.615
		0.615
	Pyridoxine x HCl Riboflavin	0.21
	Thiamine x HCl	1.01
	Vitamin B12	1.01
When 5,958.00	mg/L HEPES is included there is only 6,	

sodium chloride.

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Ham's F12 Medium

Description

In the past Ham's F12 was the first choice for a serum-free cultivation of CHO-cells and is now substituted through better serum-free systems like our Panserin C6000, which is protein-free in addition. However, Ham's F12 is an appropriate medium for mammalian cells when it is supplemented with FBS. It contains a high concentration of vitamins, amino acids and trace elements. The content of zinc sulphate is increased and it contains putrescine and linoleic acid.

Liquid Media

Ham's F12 Medium(1) without L-Glutamine with 1.176 g/L NaHCO, 500 ml P04-14550

Ham's F12 Medium(1) with L-Glutamine 500 ml P04-14500 with 1.176 g/L NaHCO₃

Ham's F12 Medium(1) with stab. Glutamine with 1.176 g/L NaHCO₂ 500 ml P04-15500

Special Media

Ham's F12 Medium(2) with L-Glutamine without Phenol red with 25 mM Hepes with 1.176 g/L NaHCO₃ 500 ml P04-14501

Ham's F12 Medium(2) without L-Glutamine without Phenol red with 1.176 g/L NaHCO₃ 500 ml P04-14559 Ham's F12K Medium(2) with L-Glutamine

500 ml P04-15600

Powder Media

with 2.5 g/L NaHCO₃

Ham's F12 Medium(1) with L-Glutamine 10 L P03-4110 without NaHCO3 50 L P03-4150

Composition

	Components	mg/L
Inorganic	Calcium chloride anhydrous	33.30
Salts	Copper(II) sulfate x 5H,O	0.003
	Iron(II) sulfate x 7H2O	0.834
	Magnesium chloride x 6H ₂ O	122.00
	Potassium chloride	223.65
	Sodium chloride	7599.9
	di-Sodium hydrogen	142.04
	phosphate anhydrous	
	Zinc sulfate x 7H ₂ O	0.86
Other	D(+)-Glucose anhydrous	1801.60
Compo-	Hepes	5958.00
nents	Hypoxanthine	4.08
	Linoleic acid	0.084
	DL-Lipoic acid	0.21
	Phenol red	1.20
	Putrescine x 2HCl	0.16
	Sodium pyruvate	110.00
	Thymidine	0.73
Amino	L-Alanine	8.91
Acids	L-Arginine x HCl	210.70
	L-Asparagine x H ₂ O	15.01
	L-Aspartic acid	13.31
	L-Cysteine x HCl x H ₂ O	35.12
	L-Glutamine	146.20
	L-Glutamic acid	14.71
	Glycine	7.51
	L-Histidine x HCl x H ₂ O	20.96
	L-Isoleucine	3.94
	L-Leucine	13.12
	L-Lysine x HCl	36.54
	L-Methionine	4.48
	L-Phenylalanine	4.96
	L-Proline	34.53
	L-Serine	10.51
	L-Threonine	11.91
	L-Tryptophan	2.04
	L-Tyrosine	5.44
	L-Valine	11.71
Vitamins	D(+)-Biotin	0.007
	D-Calcium pantothenate	0.24
	Choline chloride	13.96
	Folic acid	1.32
	myo-Inositol	18.00
	Nicotinamide	0.037
	Pyridoxine x HCl	0.037
	Riboflavin	0.062
	Thiamine x HCl	0.038
	Vitamin B12 mg/L HEPES is included there is only 7	1.36

sodium chloride.

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



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Iscove's Modified Dulbecco's Medium

Description

The IMDM is a modified DMEM, with a higher content of vitamins, selenium and amino acids. As it is supplemented with albumin, transferrin and soy lipids it can be excellently applied for culturing lymphocytes, marrow cells or hybridoma cells. Note: for hybridomas there is a better and highly efficient protein-free medium available: our Panserin H4000.

Liquid Media

IMDM⁽¹⁾ without L-Glutamine with 3.024 g/L NaHCO₃ 500 ml P04-20250

IMDM⁽¹⁾
with L-Glutamine

with 3.024 NaHCO₃ 500 ml P04-20350

IMDM⁽¹⁾
without L-Glutamine
with 25 mM Hepes
with 3 024 g/L NaHC0

with 3.024 g/L NaHCO₃ 500 ml P04-20050

IMDM⁽¹⁾
with L-Glutamine
with 25 mM Hepes

with 3.024 g/L NaHCO₃ 500 ml P04-20150

Composition

	Components	mg/L
Inorganic	Calcium chloride anhydrous	165.00
Salts	Potassium chloride	330.00
	Potassium nitrate	0.076
	Magnesium sulfate anhydrous	97.66
	Sodium chloride	5,005.00
	Sodium dihydrogen phosphate	125.00
	x H ₂ O	
	Sodium selenite x 5H ₂ O	0.01
Other	D(+)-Glucose anhydrous	4500.00
Compo-	Hepes	5958.00
nents	Sodium pyruvate	110.00
	Phenol red	15.00
Amino	L-Alanine	25.00
Acids	L-Arginine x HCl	84.00
	L-Asparagine x H ₂ O	28.40
	L-Aspartic acid	30.00
	L-Cystine x 2HCl	91.24
	L-Glutamine	584.00
	L-Glutamic acid	75.00
	Glycine	30.00
	L-Histidine x HCl x H ₂ O	42.00
	L-Isoleucine	105.00
	L-Leucine	105.00
	L-Lysine x HCl	146.00
	L-Methionine	30.00
	L-Phenylalanine	66.00
	L-Proline	40.00
	L-Serine	42.00
	L-Threonine	95.00
	L-Tryptophan	16.00
	L-Tyrosine x 2Na x 2H ₂ O	104.2
	L-Valine	94.00
Vitamins	D(+)-Biotin	0.0130
	D-Calcium pantothenate	4.00
	Choline chloride	4.00
	Folic acid	4.00
	myo-Inositol	7.20
	Nicotinamide	4.00
	Pyridoxine x HCl	4.00
	Riboflavin	0.40
	Thiamine x HCl	4.00
	Vitamin B12	0.013
When 5,958.00	mg/L HEPES is included there is only 4,50	05.00 mg/L

When 5,958.00 mg/L HEPES is included there is only 4,505.00 mg/L sodium chloride.

Iscove's Modified Dulbecco's Medium

Powder Media

IMDM ⁽²⁾ without L-Glutamine with 1.0 g/L Glucose with 3.024 g/L NaHCO ₂	500 ml	P04-20259	IMDM ⁽¹⁾ without L-Glutamine without NaHCO ₃	10 L 50 L	P03-5210 P03-5250
William 6.021 8/21 will 6.03	000 1111	10120209	$IMDM^{(1)}$		
$IMDM^{(2)}$			with L-Glutamine	10 L	P03-1310
with stab. Glutamine			without NaHCO ₃	50 L	P03-1350
with 25 mM Hepes					
without Phenol red			$IMDM^{(1)}$		
315 mOsm			with L-Glutamine		
with 3.024 g/L NaHCO ₃	500 ml	P04-20451S1	with 25 mM Hepes	10 L	P03-1410
3			without NaHCO.	50 L	P03-1450

IMDM⁽²⁾
with stab. G

 $IMDM^{(2)}$

320 mOsm

 $IMDM^{(2)}$

 $IMDM^{(2)}$

with L-Glutamine

with 25 mM Hepes

with L-Glutamine

with 1.5 g/L NaHCO₃

with stab. Glutamine

with 25 mM Hepes

Special Media

with stab. Glutamine with 25 mM Hepes without Phenol red

with 3.024 g/L NaHCO₃ 500 ml P04-20451

with 3.024 g/L NaHCO₂ 500 ml P04-20450

with 3.024 g/L NaHCO₃ 500 ml P04-20150S2

500 ml P04-20351

(1) usually on stock, (2) minimum order 10 l, (3) available upon request





 $\left(1\right)$ usually on stock, (2) minimum order 10 l, (3) available upon request

IPL-41 Insect Medium

Composition

Composition			
	Components	mg/L	
Inorganic Salts	Ammonium heptamolybdate x 4H,O	0.04	
	Calcium chloride x 2H ₂ O	662.31	
	Cobalt(II) chloride x 6H ₂ O	0.05	
	Copper chloride x 2H ₂ O	0.20	
	Iron(II) sulfate x 7H ₂ O	0.55	
	Magnesium sulfate dried	1311.40	
	Manganese chloride x 4H ₂ O	0.02	
	Potassium chloride	1200.00	
	Sodium chloride	2850,00	
	Sodium dihydrogen phosphate x H ₂ O	1160.00	
	Zinc chloride	0.04	
Other	Fumaric acid	4.40	
Compo-	D(+)-Glucose anhydrous	2500.00	
nents	α-Ketoglutaric acid sodium salt	34.05	
	DL-Malic acid	53.60 1052.58	
	D-Maltose x H ₂ O Succinic acid	4.80	
	Sucrose	1650.00	
Amino	β-Alanine	300.00	
Acids	L-Arginine x HCl	800.00	
ricids	L-Aspartic acid	1300.00	
	L-Asparagine x H ₂ O	1477.14	
	L-Cystine	100.00	
	L-Glutamine	1000.00	
	L-Glutamic acid	1500.00	
	Glycine	200.00	
	L-Histidine Base	200.00	
	L-Hydroxyproline	800.00	
	L-Isoleucine	750.00	
	L-Leucine	250.00	
	L-Lysine x HCl	700.00	
	L-Methionine	1000.00	
	L-Phenylalanine	1000.00	
	L-Proline	500.00	
	L-Serine	200.00	
	L-Threonine	200.00 100.00	
	L-Tryptophan L-Tyrosine	250.02	
	L-Valine	500.00	
Vitamins	p-Aminobenzoic acid	0.32	
	D(+)-Biotin	0.16	
	D-Calcium pantothenate	0.008	
	Choline chloride	20.00	
	Folic acid	0.08	
	myo-Inositol	0.40	
	Nicotinic acid	0.16	
	Nicotinamide	0.16	
	Pyridoxine x HCl	0.40	
	Riboflavin	0.08	
	Thiamine x HCl	0.08	
	Vitamine B12	0.24	

Description

IPL-41 is primarly used for the growth and maintenance of lepidopteran cells and for the propagation of viruses in these cells lines. The medium is also used for long time culture of baculo-virus infected spodotera cells.

Special Media

IPL-41 Insect Medium⁽²⁾
with L-Glutamine
with 0.35 g/L NaHCO₃ 500 ml P04-85600

Powder Media

IPL-41 Insect Medium⁽¹⁾
without L-Glutamine 10 L P03-9210
without NaHCO₃ 50 L P03-9250

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Joklik-MEM

Description

Joklik's MEM is a modification of MEM for suspension cultures. Due to the absence of calcium chloride in this formulation the attachment of cells is reduced.

Composition

	Components	mg/L
Inorganic	Magnesium chloride x 6H,O	200.00
Salts	Potassium chloride	400.00
	Sodium chloride	6,500.00
	Sodium dihydrogen phosphate	1,327.00
	x H ₂ O	
Other	D(+)-Glucose anhydrous	2,000.00
Compo-	Phenol red	10.00
nents		
Amino	L-Arginine x HCl	126.00
Acids	L-Cystine	24.00
	L-Glutamine	294.00
	L-Histidine Base	31.00
	L-Isoleucine	52.00
	L-Leucine	52.00
	L-Lysine x H ₂ O	65.00
	L-Methionine	15.00
	L-Phenylalanine	32.00
	L-Threonine	48.00
	L-Tryptophan	10.00
	L-Tyrosine	32.60
	L-Valine	46.00
Vitamins	D-Calcium pantothenate	1.00
	Choline chloride	1.00
	Folic acid	1.00
	myo-Inositol	2.00
	Nicotinamide	1.00
	Pyridoxal x HCl	1.00
	Riboflavin	0.10
	Thiamine x HCl	1.00

Liquid Media

Joklik – MEM⁽¹⁾ Hepes Medium with L-Glutamine

with 3.6 g/L Hepes 500 ml P04-21300

Special Media

Joklik - MEM⁽²⁾
modified for spinner culture
with EBSS (modified)
without L-Glutamine
without Antibiotica
without Calciumchloride
with 2.0 g/L NaHCO₃ 500 ml P04-21200

Powder Media

Joklik - MEM⁽¹⁾
modified for spinner culture
with EBSS (modified)
without L-Glutamine
without Antibiotica
without Calciumchloride 10 L P03-02010P
without NaHCO₃ 50 L P03-02050P

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Leibovitz's L-15 Medium

Description

L-15 contains no sodium hydrogen carbonate and no bicarbonate, because it is buffered already by a high concentration of amino acids. The L-15 medium supports the growth of established cells like Hep-2, but also human nerve cells and primary tissue explants. With 10 % tryptose phosphate broth it is also ideally suited for the cultivation of insect cell lines.

Composition

	Components	mg/L
Inorganic	Calcium chloride x 2H ₂ O	185.44
Salts	Magnesium chloride x 6H ₂ O	200.00
	Magnesium sulfate dried	139.53
	Potassium chloride	400.00
	Potassium dihydrogen	60.00
	phosphate	
	Sodium chloride	8,000.00
	di-Sodium hydrogen phosphate	190.00
Other	D(+)-Galactose anhydrous	900.00
Compo-	Hepes	5,958.00
nents	Phenol red	10.00
	Sodium pyruvate	550.00
Amino	L-Alanine	225.00
Acids	L-Arginine Base	500.00
	L-Asparagine x H ₂ O	250.00
	L-Cysteine	120.00
	L-Glutamine	300.00
	Glycine	200.00
	L-Histidine Base	250.00
	L-Isoleucine	250.00
	L-Leucine	125.00
	L-Lysine x HCl	93.75
	L-Methionine	75.00
	L-Phenylalanine	125.00
	L-Serine	200.00
	L-Threonine	300.00
	L-Tryptophan	20.00
	L-Tyrosine	300.00
	L-Valine	100.00
Vitamins	D-Calcium pantothenate	1.00
	Choline chloride	1.00
	Folic acid	1.00
	myo-Inositol	2.00
	Nicotinamide	1.00
	Pyridoxine x HCl	1.00
	Riboflavin-5'-phosphate	0.1075
	sodium salt x 2H ₂ O	1.00
	Thiamine monophosphate chloride x2 H ₂ O	1.00

When 5,958.00 mg/L HEPES is included there is only 7,500.00 mg/L sodium chloride.

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Liquid Media

Leibovitz's L-15 Medium⁽¹⁾ without L-Glutamine

without NaHCO₃ 500 ml P04-27055

Leibovitz's L-15 Medium⁽¹⁾ with L-Glutamine

without NaHCO, 500 ml P04-27500

Special Media

Leibovitz's L-15 Medium⁽²⁾ with stab. Glutamine

without NaHCO₃ 500 ml P04-27050

Leibovitz's L-15 Medium⁽²⁾ without L-Glutamine without Phenol red

without NaHCO, 500 ml P04-27054

Powder Media

Leibovitz's L-15 Medium⁽¹⁾

with L-Glutamine 10 L P03-1510 without NaHCO₃ 50 L P03-1550

Leibovitz's L-15 Medium⁽¹⁾ with L-Glutamine

with 25 mM Hepes 10 L P03-1610 without NaHCO₃ 50 L P03-1650

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Mc Coy's 5A Medium

Description

Mc Coy's 5A Medium is a complete medium with all amino acids and vitamins. It is used for growing primary cultures. This group contains marrow cells, gingival cells, adrenal cells, spleen cells, lung cells, rat embryos and other cell types.

Liquid Media

Mc Coy's 5A Medium (modified)⁽¹⁾

with L-Glutamine

with 2.2 g/L NaHCO₃ 500 ml P04-05500

Mc Coy's 5A Medium (modified)⁽¹⁾

with stab. Glutamine

with 2.2 g/L NaHCO₃ 500 ml P04-06500

Special Media

Mc Coy's 5A Medium (modified)⁽²⁾

with L-Glutamine

with 25 mM Hepes

with 2.2 g/L NaHCO $_3$ 500 ml P04-05050

Mc Coy's 5A Medium⁽²⁾ without L-Glutamine

without Phenol red

with 2.2 g/L NaHCO₃ 500 ml P04-05610

Powder Media

McCoy's 5A Medium (modified)(1)

with L-Glutamine 10 L P03-1710 without NaHCO, 50 L P03-1750

McCoy's 5A Medium (modified)(1)

with L-Glutamine

with 25 mM Hepes 10 L P03-1810 without NaHCO₃ 50 L P03-1850

Composition

	_	
	Components	mg/L
Inorganic	Calcium chloride x 2H ₂ O	132.46
Salts	Magnesium sulfate dried	139.53
	Potassium chloride	400.00
	Sodium chloride	6,460.00
	Sodium dihydrogen phosphate	580.00
	x H ₂ O	
Other	<u></u>	2 000 00
	D(+)-Glucose anhydrous	3,000.00
Compo-	Glutathione (red.)	0.50
nents	Hepes	5,958.00
	Bacto - Peptone	600.00
	Phenol red	10.00
Amino	L-Alanine	13.36
Acids	L-Arginine x HCl	42.10
	L-Asparagine x H ₂ O	45.00
	L-Aspartic acid	19.97
	L-Cysteine	24.24
	L-Glutamine	219.20
	L-Glutamic acid	22.10
	Glycine	7.50
	L-Histidine x HCl x H ₂ O	20.76
	L-Hydroxyproline ²	19.70
	L-Isoleucine	39.36
	L-Leucine	39.36
	L-Lysine x HCl	36.50
	L-Methionine	14.90
	L-Phenylalanine	16.50
	L-Proline	17.30
	L-Serine	26.30
	L-Threonine	17.90
	L-Tryptophan	3.10
	L-Tyrosine	18.10
	L-Valine	17.60
Vitamins		
vitamins	p-Aminobenzoic acid	1.00
	Ascorbic acid	0.50
	D(+)-Biotin	0.20
	D-Calcium pantothenate	0.20
	Choline chloride	5.00
	Folic acid	10.00
	myo-Inositol	36.00
	Nicotinamide	0.50
	Nicotinic acid	0.50
	Pyridoxal x HCl	0.50
	Pyridoxine x HCl	0.50
	Riboflavin	0.20
	Thiamine x HCl	0.20
	Vitamin B12	2.00

When 5,958.00 mg/L HEPES is included there is only 5,960.00 mg/L sodium chloride.

1) usually on stock, (2) minimum order 10 l, (3) available upon request



MCDB 131 Medium

www.pan-biotech.com

Composition

Composition				
	Components	mg/L		
Inorganic	Ammonium Metavandate	0.0006		
Salts	Calcium Chloride x 2H ₂ O	235.05		
	Copper(II) Sulfate x 5H ₂ O	0.0012		
	Iron (III) sulfate x 7H ₂ O	0.283		
	Magnesiumsulfate dried	1565.20		
	Manganese Sulfate x H ₂ O	0.0002		
	Ammonium Molybdate x 4H ₂ O	0.0037		
	Nickel Chloride x 6H ₂ O	0.0007		
	Potassium Chloride	298.00		
	Sodium Chloride	6,430.00		
	Sodium Metasilicate x 5H,O	2.09		
	Di-Sodium hydrogen phosphate	71.00		
	Sodium Selenite anhydrous	0.0039		
	Zinc Sulfate x 7H ₂ O	0.0003		
Other	Adenine	0.135		
Compo-	D-Glucose	1,000.00		
nents	DL-α-Lipoic acid	0.0021		
	Phenol Red	10.00		
	Putrescine x 2HCl	0.002		
	Sodium pyruvate	110.00		
	2'Deoxythymidine	0.024		
Amino	L-Alanine	2.70		
Acids	L-Arginine x HCl	63.20		
110103	L-Asparagine x H,O	15.00		
	L-Aspartic Acid	13.30		
	L-Cysteine x HCl x H ₂ O	35.00		
	L-Glutamic Acid	4.00		
	L-Glutamine	1,461.00		
	Glycine	2.30		
	L-Histidine x HCl x H,O	42.00		
	L-Isoleucine	66.00		
	L-Leucine	131.00		
	L-Lysine x HCl	182.00		
	L-Methionine	15.00		
	L-Phenylalanine	33.00		
	L-Proline	11.50		
	L-Serine	32.00		
	L-Threonine	12.00		
	L-Tryptophan	4.10		
	L-Tyrosine	18.10		
	L-Valine	117.10		
Vitamins	D-Biotin	0.0073		
	Choline Chloride	13.96		
	Folic Acid	0.60		
	myo-Inositol	7.20		
	Niacinamide	6.10		
	D-Calcium-pantothenate	12.00		
	Pyridoxine x HCl	2.10		
	Riboflavin	0.0038		
	Thiamine x HCl	3.40		
	Vitamin B12	0.0136		
	1			

Description

MCDB 131 is a medium for the cultivation of human micro-vascular endothelial cells under reduced serum content. For this purpose it has be supplemented with dialyzed serum, EGF and hydrocortisone.

Liquid Media

MCDB 131⁽¹⁾ without L-Glutamine with 1.176 g/L NaHCO₃ 500 ml P04-80057

Special Media

MCDB 131⁽²⁾
with L-Glutamine
with 1.176 g/L NaHCO₃ 500 ml P04-80053

MCDB 131⁽²⁾
without Glutamine
with 25 mM Hepes
with 1.176 g/L NaHCO₃ 500 ml P04-80054

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Medium 199 with Earle's Salts

Description

The M199 was originally developed to assay the nutrient demand of embryonic chicken fibroblasts. But it works very well with cells from many different animal species. For example, it is used for vaccine production in virology. For long term cultures serum should be added.

Composition

	Composition	
	Components	mg/L
Inorganic	Calcium chloride x 2H ₂ O	264.92
Salts	Iron(III) nitrate x 9H ₂ O	0.72
	Magnesium sulfate dried	139.52
	Potassium chloride	400.00
	Sodium acetate x 3H ₂ O	82.95
	Sodium chloride	6,800.00
	Sodium dihydrogen phosphate	140.00
Other	Adenine sulfate	10.00
Compo-	AMP	0.20
nents	ATP	1.00
	Cholesterol	0.20
	2'-Deoxyribose	0.50
	D(+)-Glucose anhydrous	1,000.00
	Glutathione (red,)	0.05
	Guanine x HCl	0.30
	Hepes	5,958.00
	Hypoxanthine	0.30
	Phenol red	10.00
	D-Ribose	0.50
	Thymine	0.30
	Tween 80	4.90
	Uracil	0.30
	Xanthine	0.30
Amino	L-Alanine	25.00
Acids	L-Arginine x HCl	70.00
	L-Aspartic acid	30.00
	L-Cysteine x HCl x H ₂ O	0.10
	L-Cystine	20.00
	L-Glutamine	100.00
	L-Glutamic acid	67.00
	Glycine	50.00
	L-Histidine x HCl x H ₂ O	21.88
	L-Hydroxyproline	10.00
	L-Isoleucine	20.00
	L-Leucine	60.00
	L-Lysine x HCl	70.00
	L-Methionine	15.00
	L-Phenylalanine	25.00
	L-Proline	40.00
	L-Serine	25.00
	L-Threonine	30.00
	L-Tryptophan	10.00
	L-Tyrosine	40.00
	L-Valine	25.00

(I)	usually on	stock, (2)	minimum	order	101, (3)	available upon reque	st

Vitamins	p-Aminobenzoic acid	0.05
	Ascorbic acid	0.05
	D(+)-Biotin	0.01
	Calciferol	0.10
	D-Calcium pantothenate	0.01
	Choline chloride	0.50
	Folic acid	0.01
	myo-Inositol	0.05
	Menadione	0.01
	Nicotinic acid	0.025
	Nicotinamide	0.025
	Pyridoxal x HCl	0.025
	Pyridoxol x HCl	0.025
	Riboflavin	0.01
	DL-α-Tocopherol phosphate-	0.01
	Na2	
	Thiamine x HCl	0.01
	Vitamin A acetate	0.14

When 5,958.00 mg/L HEPES is included there is only 6,300.00 mg/L sodium chloride.

Liquid Media

M199 with EBSS⁽¹⁾
without L-Glutamine
with 2.2 g/L NaHCO₃
500 ml P04-07500

Special Media

Powder Media

 $\begin{array}{lll} \text{M199 with EBSS}^{(1)} & & \\ \text{with L-Glutamine} & & 10 \text{ L} & \text{P03-1910} \\ \text{without NaHCO}_3 & & 50 \text{ L} & \text{P03-1950} \\ \end{array}$



Medium 199 with Hank's Salts

Special Media

M199 with HBSS(2) without L-Glutamine with 0.35 g/L NaHCO₃

500 ml P04-07753

M199 with HBSS(2) with L-Glutamine with 25 mM Hepes

with 0.35 g/L NaHCO, 500 ml P04-07450

M199 with HBSS (10X)(2) without L-Glutamine

without NaHCO, 500 ml P04-07600

Powder Media

M199 with HBSS(1)

with L-Glutamine P03-2110 10 L without NaHCO, 50 L P03-2150

Composition

	Components	mg/L
Inorganic	Calcium chloride x2H ₂ O	185.45
Salts	Iron(III) nitrate x 9H,O	0.72
	Magnesium sulfate dried	139.68
	Potassium chloride	400.00
	Potassium dihydrogen	60.00
	phosphate	
	Sodium acetate x 3H ₂ O	83.00
	Sodium chloride	8,000.00
	di-Sodium hydrogen phosphate	47.68
Other	Adenine sulfate	10.00
Compo-	AMP	0.20
nents	ATP	1.00
	Cholesterol	0.20
	2'-Deoxyribose	0.50
	D(+)-Glucose anhydrous	1,000.00
	Glutathione (red.)	0.05
	Guanine x HCl	0.30
	Hepes	5,958.00
	Hypoxanthine	0.30
	Phenol red	10.00
	D-Ribose	0.50
	Thymine	0.30
	Tween 80	4.90
	Uracil	0.30
	Xanthine	0.30

Amino	L-Alanine	25.00
Acids	L-Arginine x HCl	70.00
Acius	L-Aspartic acid	30.00
	L-Aspartic acid L-Cysteine x HCl x H,O	0.10
	L-Cystine	20.00
	L-Glutamine	100.00
	L-Glutamic acid	67.00
	Glycine	50.00
	L-Histidine x HCl x H ₂ O	21.88
	L-Hydroxyproline L-Isoleucine	10.00
	L-Isoleucine L-Leucine	20.00
		60.00
	L-Lysine x HCl	70.00
	L-Methionine	15.00
	L-Phenylalanine	25.00
	L-Proline	40.00
	L-Serine	25.00
	L-Threonine	30.00
	L-Tryptophan	10.00
	L-Tyrosine	40.00
	L-Valine	25.00
Vitamins	p-Aminobenzoic acid	0.05
	Ascorbic acid	0.05
	D(+)-Biotin	0.01
	Calciferol	0.10
	D-Calcium pantothenate	0.01
	Choline chloride	0.50
	Folic acid	0.01
	myo-Inositol	0.05
	Menadione	0.01
	Nicotinic acid	0.025
	Nicotinamide	0.025
	Pyridoxal x HCl	0.025
	Pyridoxol x HCl	0.025
	Riboflavin	0.01
	DL-α-Tocopherol phosphate-	0.01
	Na,	
	Thiamine x HCl	0.01
	Illiallille X I I CI	0.01



MEM with Earle's Salts

Media

Description

MEM is an advancement of the BME and the base medium of many further modifications. Because BME did not fulfil all requirements for some mammalian and HeLa cells, a better variation had to be developed. Today, MEM is one of the most used synthetic media and shows its versatility by supplementing with amino acids including Hank's or Earle's salts. Even the addition of only small amounts of FBS results in a positive effect on cell growth.

Composition

	Components	mg/L
Inorganic	Calcium chloride x2H,O	264.92
Salts	Magnesium sulfate dried	139.53
	Potassium chloride	400.00
	Sodium chloride	6800.00
	Sodium dihydrogen phosphate x H ₂ O	140.00
Other	D(+)-Glucose anhydrous	1000.00
Compo-	Hepes	5958.00
nents	Phenol red	10.00
Amino	L-Alanine	8.90
Acids	L-Arginine x HCl	126.00
	L-Asparagine x H ₂ O	13.20
	L-Aspartic acid	13.30
	L-Cystine	24.00
	L-Glutamine	292.00
	L-Glutamic acid	14.70
	Glycine	7.50
	L-Histidine x HCl x H,O	42.00
	L-Isoleucine 2	52.00
	L-Leucine	52.00
	L-Lysine x HCl	72.50
	L-Methionine	15.00
	L-Phenylalanine	32.00
	L-Proline	11.50
	L-Serine	10.50
	L-Threonine	48.00
	L-Tryptophan	10.00
	L-Tyrosine	36.00
	L-Valine	46.00
Vitamins	D-Calcium pantothenate	1.00
	Choline chloride	1.00
	Folic acid	1.00
	myo-Inositol	2.00
	Nicotinamide	1.00
	Pyridoxal x HCl	1.00
	Riboflavin	0.10
	Thiamine x HCl	1.00

When 5,958.00 mg/L HEPES is included there is only 6,300.00 mg/L sodium chloride.



Liquid Media without Glutamine

MEM Eagle with EBSS⁽¹⁾ without L-Glutamine

with 2.2 g/L NaHCO3 500 ml P04-08050

MEM Eagle with EBSS⁽¹⁾ without L-Glutamine with 25 mM Hepes

with 2.2 g/L NaHCO₃ 500 ml P04-08150

Special Media without Glutamine

MEM Eagle with EBSS⁽²⁾ without L-Glutamine without Phenol red

with 2.2 g/L NaHCO3 500 ml P04-00507

MEM Eagle with EBSS⁽²⁾ without L-Glutamine with NEAA

500 ml P04-08509 with 2.2 g/L NaHCO,

MEM Eagle with EBSS⁽²⁾ without L-Glutamine

without NaHCO, 500 ml P04-09050

Powder Media without Glutamine

MEM Eagle with EBSS(1)

without L-Glutamine 10 L P03-7410 without NaHCO3 50 L P03-7450

(1) usually on stock, (2) minimum order 10 l, (3) available upon request

MEM with Earle's Salts

Liquid Media with L-Glutamine

MEM Eagle with EBSS⁽¹⁾ with L-Glutamine with 2.2 g/L NaHCO3 500 ml P04-08500

MEM Eagle with EBSS⁽¹⁾ with L-Glutamine

with 1.5 g/L NaHCO, 500 ml P04-00509

Special Media with L-Glutamine

MEM Eagle with EBSS⁽²⁾ with L-Glutamine without Phenol red with 1.5 g/L NaHCO3

500 ml P04-00508 MEM Eagle with EBSS⁽²⁾

with L-Glutamine with 20 mM Hepes with 2.2 g/L NaHCO3

500 ml P04-08549

MEM Eagle with EBSS⁽²⁾ with L-Glutamine with NEAA

with 2.2 g/L NaHCO₃ 500 ml P04-08510

MEM Eagle with EBSS⁽²⁾ with 2 mM Glutamine with 1 mM Pyruvate

with NEAA

with 1.5 g/L NaHCO₃ 500 ml P04-08056

Powder Media with L-Glutamine

MEM Eagle with EBSS⁽¹⁾ with L-Glutamine with NEAA 10 L P03-2910 without NaHCO3 50L P03-2950 MEM Eagle with EBSS⁽¹⁾ with L-Glutamine with NEAA with 25 mM Hepes 10 L P03-3010 50 L P03-3050 with NaHCO₃ MEM Eagle with EBSS⁽¹⁾ with L-Glutamine 10 L P03-2710 without NaHCO3 50 L P03-2750 MEM Eagle with EBSS⁽¹⁾ with L-Glutamine with 25 mM Hepes 10 L P03-2810 without NaHCO, 50 L P03-2850

Liquid Media with stab. Glutamine

MEM Eagle with EBSS⁽¹⁾ with stab. Glutamine

with 2.2 g/L NaHCO, 500 ml P04-09500

MEM Eagle with EBSS⁽¹⁾ with stab. Glutamine with 25 mM Hepes

with 2.2 g/L NaHCO₃ 500 ml P04-08250

MEM with Hank's Salts

Media

Composition

	Components	mg/L
Inorganic	Calcium chloride x 2H,O	185.44
Salts	Potassium chloride	400.00
	Potassium dihydrogen	60.00
	phosphate anhydrous	
	Magnesium sulfate dried	139.53
	Sodium chloride	8,000.00
	di-Sodium hydrogen phosphate	47.88
Other	D(+)-Glucose anhydrous	1,000.00
Compo-	Hepes	5,958.00
nents	Phenol red	10.00
Amino	L-Alanine	8.90
Acids	L-Arginine x HCl	126.00
	L-Asparagine x H ₂ O	13.20
	L-Aspartic acid	13.30
	L-Cystine	24.00
	L-Glutamine	292.00
	L-Glutamic acid	14.70
	Glycine	7.50
	L-Histidine x HCl x H ₂ O	42.00
	L-Isoleucine	52.00
	L-Leucine	52.00
	L-Lysine x HCl	72.50
	L-Methionine	15.00
	L-Phenylalanine	32.00
	L-Proline	11.50
	L-Serine	10.50
	L-Threonine	48.00
	L-Tryptophan	10.00
	L-Tyrosine	36.00
	L-Valine	46.00
Vitamins	D-Calcium pantothenate	1.00
	Cholin chloride	1.00
	Folic acid	1.00
	myo-Inositol	2.00
	Nicotinamide	1.00
	Pyridoxal x HCl	1.00
	Riboflavin	0.10
TATL	Thiamine x HCl	1.00

When 5,958.00 mg/L HEPES is included there is only 7,500.00 mg/L

Special Media

MEM Eagle with HBSS ⁽²⁾ without L-Glutamine with 0.35 g/L NaHCO ₃	500 ml	P04-10050
MEM Eagle with HBSS ⁽²⁾ with L-Glutamine with 0.35 g/L NaHCO ₃	500 ml	P04-10500
MEM Eagle with HBSS ⁽²⁾ with L-Glutamine with 0.60 g/L NaHCO ₃	500 ml	P04-10599
Powder Media		
MEM Eagle with HBSS ⁽¹⁾ with L-Glutamine	10 L	P03-3310
without NaHCO ₃	50 L	P03-3350

sodium chloride.



(1) usually on stock, (2) minimum order 10 l, (3) available upon request





RPMI 1640

Description

The medium was developed for culture of normal and neoplastic leukocytes, but also marrow cells and hybridoma cells. Meanwhile there are better, serum-free media for hybridoma cells such our Panserin H4000. Just by supplementing RPMI 1640 with varying amounts of FBS, a very good medium for many different cell lines can be obtained.

Composition

	Components	mg/L
Inorganic	Calcium nitrate x 4H ₂ O	100.00
Salts	Potassium chloride	400.00
	Magnesium sulfate anhydrous	48.83
	Sodiumchloride	6000.00
	di-Sodium hydrogen phosphate	800.49
Other	D(+)-Glucose anhydrous	2000.00
Compo-	Glutathion (red.)	1.00
nents	Hepes	5958.00
	Phenol red	5.00
Amino	L-Arginine x HCl	241.86
Acids	L-Asparagine x H ₂ O	50.00
	L-Aspartic acid	20.00
	L-Cystine x 2HCl	65.19
	L-Glutamine	300.00
	L-Glutamic acid	20.00
	Glycine	10.00
	L-Histidine x HCl x H,O	20.27
	L-Hydroxyproline ²	20.00
	L-Isoleucine	50.00
	L-Leucine	50.00
	L-Lysine x HCl	40.00
	L-Methionine	15.00
	L-Phenylalanine	15.00
	L-Proline	20.00
	L-Serine	30.00
	L-Threonine	20.00
	L-Tryptophan	5.00
	L-Tyrosine x 2Na	28.83
	L-Valine	20.00
Vitamins	p-Aminobenzoic acid	1.00
	D-(+)-Biotin	0.20
	D-Calcium pantothenate	0.25
	Choline chloride	3.00
	Folic acid	1.00
	myo-Inositol	35.00
	Nicotinamide	1.00
	Pyridoxine x HCl	1.00
	, Riboflavin	0.20
	Thiamine x HCl	1.00

When 5,958.00 mg/L HEPES is included there is only 5,000.00 mg/L sodium chloride.

Liquid Media without Glutamine

RPMI 1640⁽¹⁾ without L-Glutamine with 2.0 g/L NaHCO3 500 ml P04-17500

RPMI 1640⁽¹⁾ without L-Glutamine without Phenol red

with 2.0 g/L NaHCO₃ 500 ml P04-16516

RPMI 1640⁽¹⁾ without L-Glutamine with 25 mM Hepes

with 2.0 g/L NaHCO3 500 ml P04-18000



RPMI 1640

Media

Special Media without C	Glutamine	!
RPMI 1640 ⁽²⁾ without L-Glutamine without Calcium with 2.0 g/L NaHCO ₃	500 ml	P04-16151
RPMI 1640 ⁽²⁾ without L-Glutamine without L-Tryptophan with 2.0 g/L NaHCO ₃	500 ml	P04-17599
RPMI 1640 ⁽²⁾ without L-Glutamine without Glucose with 2.0 g/L NaHCO ₃	500 ml	P04-17550
RPMI 1640 ⁽²⁾ without L-Glutamine with 15 mM Hepes without Phosphate with 2.0 g/L NaHCO ₃	500 ml	P04-21049
RPMI 1640 (10X) ⁽²⁾ without L-Glutamine without NaHCO ₃	500 ml	P04-17510
RPMI 1640 ⁽²⁾ without L-Glutamine with 25 mM Hepes without NaHCO ₃	500 ml	P04-17850
RPMI 1640 ⁽²⁾		

500 ml P04-22500

500 ml P04-19500

without L-Glutamine with 25 mM Hepes with 2.2 g/L NaHCO₃

without L-Glutamine with 20 mM Hepes with 0.85 g/L NaHCO₃

RPMI 1640⁽²⁾

Powder Media without Glutamine

RPMI 1640 ⁽¹⁾		
without L-Glutamine	10 L	P03-7210
without $NaHCO_3$	50 L	P03-7250
RPMI 1640 ⁽¹⁾		
without L-Glutamine		
without Phenol red	10 L	P03-7710
without $NaHCO_3$	50 L	P03-7750
RPMI 1640 ⁽¹⁾		
without L-Glutamine		
with 25 mM Hepes	10 L	P03-4410
without NaHCO ₃	50 L	P03-4450



RPMI 1640

Liquid Media with L-Glutamine

RPMI $1640^{(1)}$ with L-Glutamine with $2.0~\rm{g/L~NaHCO_3}$ 500 ml P04-16500

RPMI 1640⁽¹⁾ with L-Glutamine without Phenol red

with 2.0 g/L NaHCO₃ 500 ml P04-16515

RPMI 1640⁽¹⁾

with 2 mM L-Glutamine with 1 mM Sodium pyruvate with 4.5 g/L Glucose with 10 mM Hepes

with 1.5 g/L NaHCO₃ 500 ml P04-18047

RPMI 1640⁽¹⁾ with L-Glutamine with 25 mM Hepes with 2.2 g/L NaHCO

with 2.2 g/L NaHCO₃ 500 ml P04-22100

Special Media with L-Glutamine

RPMI 1640⁽²⁾ with L-Glutamine without Glucose

with 2.0 g/L NaHCO₃ 500 ml P04-17545

RPMI 1640⁽²⁾ with L-Glutamine without L-Arginine

with 2.0 g/L NaHCO₃ 500 ml P04-16598

RPMI 1640⁽²⁾ with L-Glutamine without L-Tryptophan

with 2.0 g/L NaHCO₃ 500 ml P04-17598

RPMI 1640⁽²⁾ with L-Glutamine with 20 mM Hepes with 0.85 g/L NaHCO₃

Powder Media with L-Glutamine

RPMI 1640⁽¹⁾ with L-Glutamine 10 L P03-4310 without NaHCO3 50 L P03-4350 RPMI 1640⁽¹⁾ with L-Glutamine with 25 mM Hepes 10 L P03-7310 without NaHCO3 50 L P03-7350 RPMI 1640⁽¹⁾ with L-Glutamine without Phenol red 10 L P03-7610 50 L P03-7650 without NaHCO,

Liquid Media with stab. Glutamine

RPMI 1640⁽¹⁾ with stab. Glutamine with 2.0 g/L NaHCO₃ 500 ml P04-18500

with 2.0 g/L NaHCO₃ 500 ml P04-18500

RPMI 1640⁽¹⁾
with stab. Glutamine
with 25 mM Hepes
with 2.2 g/L NaHCO₃ 500 ml P04-18050

Special Media with stab. Glutamine

RPMI 1640⁽²⁾ with stab. L-Glutamine without Phenol red

with 2.0 g/L NaHCO₃ 500 ml P04-16520

with stab. Glutamine without Glucose

RPMI 1640⁽²⁾

with 2.0 g/L NaHCO₃ 500 ml P04-17546

RPMI 1640⁽²⁾ with stab. Glutamine without Phenol red without Glucose

with 2.0 g/L NaHCO₃ 500 ml P04-16530

Schneider's Drosophila Medium

Description

Originally developed for the culture of Drosophila cells, this medium is also suitable for the culture of other dipteran cell lines.

Liquid Media

Schneider's Drosophila Medium⁽¹⁾
without L-Glutamine
with 0.35 g/L NaHCO₂ 500 ml P04-90500

Schneider's Drosophila Medium⁽¹⁾ with L-Glutamine with 0.35 g/L NaHCO₃ 500 ml P04-91500

Powder Media

Schneider's Drosophila Medium⁽¹⁾
without L-Glutamine 10 L P03-9310
without NaHCO₂ 50 L P03-9350

Composition

	Components	mg/L
Inorganic	Potassium chloride	1600.00
Salts	Potassium dihydrogen	450.00
	phosphate	
	Magnesium sulfate dried	2585.71
	Sodium chloride	2100.00
	di-Sodium hydrogen phosphate	700.00
Other	DL-Malic acid	600.00
Compo-	Succinic acid	60.00
nents	Fumaric acid	60.00
	D(+)-Glucose anhydrous	2000.00
	Yeast extract	2000.00
	α-Ketoglutaric acid sodium salt	402.66
	D(+)-Trehalose x 2H ₂ O	2210.00
Amino	β-Alanine	500.00
Acids	L-Arginine Base	600.00
	L-Asparatic acid	400.00
	L-Cysteine free base	60.00
	L-Cystine	16.60
	L-Glutamine	1800.00
	L-Glutamic acid	800.00
	Glycine	250.00
	L-Histidine Base	400.00
	L-Isoleucine	150.00
	L-Leucine	150.00
	L-Lysine x HCl	1650.00
	L-Methionine	150.00
	L-Proline	1700.00
	L-Serine	250.00
	L-Threonine	350.00
	L-Tryptophan	100.00
	L-Tyrosine	500.00
	L-Valine	300.00

(1) usually on stock, (2) minimum order 10 l, (3) available upon request





500 ml P04-19550

TC 100 Insect Medium

www.pan-biotech.com

Description

The TC 100 Insect Medium is an absolutely serum-free formula (Oxford formulation) for the growth of insect cells, especially for SF9 cells and the breeding of viruses. If you would like to work with a modern protein-free insect medium, our Spodopan is the ideal choice.

Liquid Media

TC 100 Insect Medium(1) with L-Glutamine with 0.35 g/L NaHCO₂ 500 ml P04-92500

Special Media

TC 100 Insect Medium(2) without L-Glutamine

with 0.35 g/L NaHCO₃ 500 ml P04-93500

Powder Media

TC 100 Insect Medium⁽¹⁾

with L-Glutamine P03-9610 10 L without NaHCO3 50 L P03-9650

Composition

	Components	mg/L
Inorganic	Calcium chloride x 2H ₂ O	1298.13
Salts	Potassium chloride	2900.00
	Magnesium chloride x 6H ₂ O	2282.59
	Magnesium sulfate dried	1781.00
	Sodium dihydrogen phosphate	970.00
	x H ₂ O	
Other	D(+)-Glucose anhydrous	1000.00
Compo-	Bacto - Tryptose	2600.00
nents		
Amino	L-Alanine	225.00
Acids	L-Arginine Base	550.00
	L-Aspartic acid	350.00
	L-Asparagine x H ₂ O	391.97
	L-Cystine	20.00
	L-Glutamine	600.00
	L-Glutamic acid	600.00
	Glycine	650.00
	L-Histidine x HCl x H ₂ O	3400.00
	L-Isoleucine	50.00
	L-Leucine	75.00
	L-Lysine x HCl	630.00
	L-Methionine	50.00
	L-Phenylalanine	150.00
	L-Proline	350.00
	L-Serine	550.00
	L-Threonine	180.00
	L-Tryptophan	100.00
	L-Tyrosine	55.00
	L-Valine	100.00
Vitamins	p-Amino benzoic acid	0.02
	D-(+)-Biotin	0.01
	D-Calcium pantothenate	0.11
	Folic acid	0.02
	myo-Inositol	0.02
	Nicotinic acid	0.02
	Pyridoxine x HCl	0.02
	Riboflavin	0.02
	Thiamine x HCl	0.02
	Vitamin B12	0.01

TNM-FH Medium

www.pan-biotech.com

Description

The TNM-FH is a variation of the Grace medium. This modification has proved as a good culture medium for many lepidopteran cells.

Liquid Media

TNM-FH Medium(1) with L-Glutamine with Lactalbumine-Hydrolysate with Yeast extract with 0.35 g/L NaHCO₃ 500 ml P04-80500

Special Media

TNM-FH Medium(2) with L-Glutamine with Lactalbumin-Hydrolysate with Yeast extract with 10 % Fetal Bovine Serum with 0.35 g/L NaHCO₂ 500 ml P04-83500

Powder Media

TNM-FH Insect Medium(1) without L-Glutamine with Lactalbumine-Hydrolysate with Yeast extract P03-9710 10 L without NaHCO3 50 L P03-9750

Composition

	Components	mg/L
Inorganic	Calcium chloride x 2H,O	1,324.62
Salts	Potassium chloride	2,240.00
	Magnesium chloride x 6H ₂ O	2,278.86
	Magnesium sulfate dried	1,939.80
	di-Sodium hydrogen	876.92
	phosphate	
Other	DL-Malic acid	670.00
Compo-	Succinic acid	60.00
nents	D-Fructose	400.00
	Fumaric acid	55.00
	D(+)-Glucose anhydrous	700.00
	Yeast extract	3,333.33
	α-Ketoglutaric acid	425.66
	sodium salt	
	Lactalbumin Hydrolysate	3,333.33
	Sucrose	26,680.00
Amino	β-Alanine	200.00
Acids	L-Alanine	225.00
	L-Arginine x HCl	700.00
	L-Asparagine x H ₂ O	350.00
	L-Aspartic acid	350.00
	L-Cystine	19.18
	L-Glutamine	600.00
	L-Glutamic acid	600.00
	Glycine	650.00
	L-Histidine Base	2,500.00
	L-Isoleucine	50.00
	L-Leucine	75.00
	L-Lysine x HCl	625.00
	L-Methionine	50.00
	L-Phenylalanine	150.00
	L-Proline	350.00
	L-Serine	550.00
	L-Threonine	175.00
	L-Tryptophan	100.00
	L-Tyrosine	50.00
	L-Valine	100.00
Vitamins		0.02
vitaiiiiis	p-Aminobenzoic acid D-(+)-Biotin	0.02
	D-(+)-Blothi D-Ca-Pantothenate	
	Cholin chloride	0.02
	Folic acid	0.20
		0.02
	myo-Inositol	0.02
	Nicotinic acid	0.02
	Pyridoxol x HCl	0.02
	Riboflavin	0.02
	Thiamine x HCl	0.02

(1) usually on stock, (2) minimum order 10 l, (3) available upon request







Waymouth's MB 752/1 Medium

Description

Waymouth's MB 752/1 Medium was developed for studies concerning nutrition and metabolism. It also can be used for growing strain L sub-lines, NCTC clone 929.

Special Media

Waymouth's MB 752/1 Medium $^{(2)}$ with L-Glutamine with 2.24 g/L NaHCO $_3$ 500 ml P04-28500

Powder Media

Waymouth's MB 752/1 Medium(1)

with L-Glutamine 10 L P03-4510 without NaHCO₃ 50 L P03-4550

Composition

	Components	mg/L
Inorganic	Calcium chloride x 2H ₂ O	120.02
Salts	Magnesium chloride x 6H,O	240.00
	Magnesium sulfate dried	130.96
	Potassium chloride	150.00
	Potassium dihydrogen	80.00
	phosphate	
	Sodium chloride	6,000.00
	di-Sodium hydrogen	300.00
	phosphate anhydrous	
Other	D(+)-Glucose anhydrous	5,000.00
Compo-	Glutathione (red.)	15.00
nents	Hepes	4,766.40
	Hypoxanthine	25.00
	Phenol red	10.00
Amino	L-Arginine x HCl	75.00
Acids	L-Aspartic acid	60.00
	L-Cysteine x HCl x H,O	100.26
	L-Cystine	15.00
	L-Glutamine	350.00
	L-Glutamic acid	150.00
	Glycine	50.00
	L-Histidine x HCl x H ₂ O	164.10
	L-Isoleucine	25.00
	L-Leucine	50.00
	L-Lysine x HCl	240.00
	L-Methionine	50.00
	L-Phenylalanine	50.00
	L-Proline	50.00
	L-Threonine	75.00
	L-Tryptophan	40.00
	L-Tyrosine	40.00
	L-Valine	65.00
Vitamins	L-Ascorbic acid	17.50
	D(+)-Biotin	0.02
	D-Calcium pantothenate	1.00
	Choline chloride	250.00
	Folic acid	0.40
	myo-Inositol	1.00
		1.00
	,	1.00
		1.00
	Thiamine x HCl	10.00
		0.20
When 4 766 40	Nicotinamide Pyridoxine x HCl Riboflavin	1.00 1.00 1.00 10.00 0.20

When 4,766.40 mg/L HEPES is included there is only 5,500.00 mg/L sodium chloride.

 $\left(1\right)$ usually on stock, (2) minimum order 10 l, (3) available upon request



William's Medium E

Description

The William's Medium E is used for long-term cultivation of adult rat liver epithelial cells.

Liquid Media

 $\label{eq:williams} Williams Medium \ E^{(1)} \\ without \ L\text{-}Glutamine$

with 2.24 g/L NaHCO₃ 500 ml P04-29050

William's Medium E⁽¹⁾ with L-Glutamine

with 2.24 g/L NaHCO₃ 500 ml P04-29500

William's Medium $E^{(1)}$ with stab. Glutamine

with 2.24 g/L NaHCO₃ 500 ml P04-29150

William's Medium E⁽¹⁾
without L-Glutamine
without Phenol red

with 2.24 g/L NaHCO₃ 500 ml P04-29510

Special Media

William's Medium E⁽²⁾
without L-Glutamine
without Glucose
with 2.24 g/L NaHCO₃ 500 ml P04-29050S1

Powder Media

William's Medium E⁽¹⁾
with L-Glutamine
with 25 mM Hepes 10 L P03-4810
without NaHCO, 50 L P03-4850

Composition

	Components	mg/I
Inorganic	Calcium chloride x 2H ₂ O	264.92
Salts	Iron(III)-nitrat x 9H ₂ O	0.0001
	Potassium chloride	400.00
	Copper(II)-sulfate x 5H ₂ O	0.0001
	Magnesium sulfate dried	139.57
	Manganese chloride x 4H ₂ O	0.0001
	Sodium chloride	6,800.00
	Sodium dihydrogen phospha-	140.00
	te x H ₂ O	
	Zinc sulfate x 7H ₂ O	0.0002
Other	D(+)-Glucose anhydrous	2,000.00
Compo-	Hepes	5,958.00
nents	Glutathion (red.)	0.05
	Methyllinoleat	0.03
	Sodium pyruvate	25.00
	Phenol red	10.00
Amino	L-Alanine	90.00
Acids	L-Arginine free base	50.00
110100	L-Asparagine x H ₂ O	20.00
	L-Aspartic acid	30.00
	L-Cysteine	40.00
	L-Cystine	20.00
	L-Glutamine	292.00
	L-Glutamine L-Glutamic acid	50.00
		50.00
	Glycine L-Histidine Base	15.00
	L-Isoleucine	
		50.00
	L-Leucine	75.00
	L-Lysine x HCl	87.50
	L-Methionine	15.00
	L-Phenylalanine	25.00
	L-Proline	30.00
	L-Serine	10.00
	L-Threonine	40.00
	L-Tryptophan	10.00
	L-Tyrosine	35.00
	L-Valine	50.00
Vitamins	L-Valine L-Ascorbic acid	
Vitamins	L-Ascorbic acid D(+)-Biotin	2.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol	2.00 0.50
Vitamins	L-Ascorbic acid D(+)-Biotin	2.00 0.50 0.10
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride	2.00 0.50 0.10 1.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate	2.00 0.50 0.10 1.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride	2.00 0.50 0.10 1.00 1.50
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride Folic acid	2.00 0.50 0.10 1.00 1.50 1.00 2.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride Folic acid myo-Inositol	2.00 0.50 0.10 1.00 1.50 1.00 2.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride Folic acid myo-Inositol Menadion sodium bisulfite	2.00 0.50 0.10 1.00 1.50 1.00 2.00 0.01 1.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride Folic acid myo-Inositol Menadion sodium bisulfite Nikotinamid	2.00 0.50 0.10 1.00 1.50 1.00 2.00 0.01 1.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride Folic acid myo-Inositol Menadion sodium bisulfite Nikotinamid Pyridoxal x HCl	2.00 0.50 0.10 1.00 1.50 1.00 2.00 0.01 1.00 0.10
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride Folic acid myo-Inositol Menadion sodium bisulfite Nikotinamid Pyridoxal x HCl Riboflavin Thiamine x HCl	2.00 0.50 0.10 1.00 1.50 1.00 2.00 0.01 1.00 0.10 1.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride Folic acid myo-Inositol Menadion sodium bisulfite Nikotinamid Pyridoxal x HCl Riboflavin Thiamine x HCl DL-α-Tocopherol phospha-	2.00 0.50 0.10 1.00 1.50 1.00 2.00 0.01 1.00 0.10 1.00
Vitamins	L-Ascorbic acid D(+)-Biotin Calciferol D-Calcium pantothenate Choline chloride Folic acid myo-Inositol Menadion sodium bisulfite Nikotinamid Pyridoxal x HCl Riboflavin Thiamine x HCl	50.00 2.00 0.50 0.10 1.00 1.50 2.00 0.01 1.00 1.00 0.10 1.00 0.10 1.00 0.10

When 5,958.00 mg/L HEPES is included there is only 6,300.00 mg/L sodium chloride.

 $\left(1\right)$ usually on stock, $\left(2\right)$ minimum order 10 l, $\left(3\right)$ available upon request



Endopan

Background

Endothelial cells line the blood and lymphatic vessels and the internal cavities of the heart. They display a strongly flattened, polygonal form and mostly rest on a basal membrane. They adhere to each other by desmosomes and tight-junctions.

With a total cell number of about one trillion (10¹²), the endothelium is one of the biggest organs of the body and plays a key role in many physiological and patho-physiological processes (e.g. cell-based immune response, wound healing, inflammation, allergy, cardiovascular diseases, tumour growth). A huge number of soluble factors circulating in the blood or released by neighbouring cells, control proliferation or apoptosis of endothelial cells and the invasion and migration of leucocytes to the endothelium, thereby regulating the maintenance, degeneration, or regeneration of blood vessels

The endothelium constitutes a highly specialized organ that lines the vascular system and lymphatic channels in a complex network of arteries, veins, and microvessels which differ in size, structure, and function. The cultivation of endothelial cells from large vessels, predominantly from human umbilical vein, is a routine procedure in many laboratories, and this has contributed huge to the development of modern vascular biology. However, there is convincing evidence that microvascular endothelial cells display a number of important functional differences, compared to large vessel-derived

endothelial cells, with regard to their growth factor response and their regulation of adhesion molecule expression.

They serve as the barrier separating circulating blood from the extracellular matrix and interstitium in the body. Cells involved in the pathogenesis of tumor angiogenesis, wound healing, and acute or chronic inflammation are predominantly of micro-vascular origin. Several functions associated with the micro-vasculature in situ are expressed by micro-vascular endothelial cells in cell culture

Micro-vessels are not simply tubes but have also a second cellular component, the mural cell or pericyte. Little is known about later stages of vessel growth, including the addition of pericytes to the capillary and its influence on endothelial growth and function. In vivo, pericytes form an incomplete envelopment around the endothelial cells within the micro-vascular basement membrane of capillaries and post-capillary venules.

Evidence clearly indicates that differences exist between endothelial cells of the microvasculature and those that line large vessels. These include differences in secreted products, in the expression of cell adhesion molecules, and in cytokine-induced regulation of cell adhesion molecules. Thus, a precise delineation of the biology of microvascular endothelial cells is crucial to our understanding of such important processes as inflammation, tumor progression, cardiac microcirculation, and blood-brain barrier function.

Endopan 3 Large Vessel Endothelial Cell Medium

Endopan 3 ready-to-use ⁽¹⁾	500 ml	P04-00100
Endopan 3 kit with 9 supplements ⁽¹⁾	500 ml	P04-0010K

Composition

Endopan 3 ready-to-use is a specially developed medium for the in vitro culture of human endothelial cells containing all components necessary for optimal growth. It is designed for use in an incubator at 37° C with a 5% CO2 atmosphere.

Endopan 3 kit is provided with FBS and supplements in separate sterile packing. This will enable the user to prepare a medium for special application. For example, FBS, VEGF, FGF-2, or other components may be omitted from the complete medium for specific experimental settings.

Endopan MV Microvascular Endothelial Cell Medium

Endopan MV ready-to-use ⁽³⁾	500 ml	P04-00200 NEW
Endopan MV kit with 8 supplements(3)	500 ml	P04-0020K NEW

(1) usually on stock, (2) minimum order 10 l, (3) available upon reques



Endopan PRO Endothelial Progenitor Medium

Media

Background

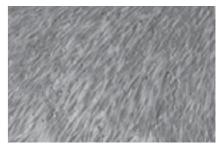
Endothelial cells line blood and lymphatic vessels and the internal cavities of the heart. They display a strongly flattened, polygonal form and mostly rest on a basal membrane. With a total number of about 10¹² cells, the endothelium is one of the biggest organs of the body and plays a key role in many physiological and pathophysiological processes. A number of factors control proliferation or apoptosis of endothelial cells, thereby regulating the maintenance, degeneration, or regeneration of blood vessels.

New blood vessel formation occurs via angiogenesis or vasculogenesis, a process thought to be restricted to embryonic development. In 1997, postnatal vasculogenesis has been proposed as an important mechanism for angiogenesis via blood or bone marrow derived circulating progenitor endothelial cells (PEC) (Asahara et al, Science 1997). Consequently, PECs have been extensively studied as a potential cell therapy for the repair of damaged blood vessels. Animal studies clearly demonstrated that administration of PECs partially rescued cardiovascular dysfunction or myocardial injury with evidence for PEC contribution to new vessel growth. In most studies, PECs are defined by cell surface expression of CD34, CD133, or VEGF-R2 (KDR). Because these molecules are also present on hematopoietic progenitors, relying only on surface markers can not exclude a contamination with hematopoietic linage cells. More recently, a PEC population has been identified which shows expression of endothelial as well as progenitor, but not hematopoietic cell markers (Ingram et al, Blood. 2004;104:2752). Importantly, these cells have been tested for a high proliferative potential in clonogenic assays and additionally characterized by formation of functional blood vessels in vivo (Yoder et al, Blood. 2007;109:1801).

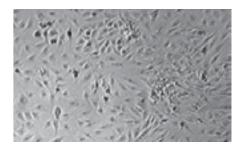
Composition

Endopan PRO ready-to-use is a complete medium specially developed for the in vitro culture of human progenitor endothelial cells (hPEC) containing all components necessary for optimal colony formation, clonogenic growth, and rapid proliferation.

Endopan PRO kit is provided with FBS growth supplement (pre-screened and tested for progenitor cells) and additional supplements in separate sterile packing. This will enable the user to prepare a medium for special application.



hPEC in Endopan PRO (P6)



hPEC colony (P1) with outgrowing cells in Endopan PRO

Endopan PRO ready-to-use ⁽³⁾	500 ml	P04-00700
Endopan PRO kit with 6 supplements(3)	500 ml	P04-0070K

1) usually on stock, (2) minimum order 10 l, (3) available upon request



Pantum

Description

Pantum are ready-to-use growth media which contain purified plasma proteins and lipids, such as serum albumin and cholesterol, specific growth factors, components of soybean extract, an iron transport protein and enriched trace elements. The new formulations result in stable cell growth under defined culture conditions. No addition of serum or growth factors is necessary.

Pantum 386 for epithelial cells

Pantum 386 is a modified formulation of DMEM and specially developed to optimize the growth of epithelial cell lines.

Pantum 586A for adherent cells

Pantum 586A is particularly suited for culture of adherent cells to stimulate their growth. It is a modification of Iscove's MEM.

Pantum L24 for lymphocytes

Pantum L24 is suited for culture of peripheral blood lymphocytes. Adult lymphocytes lack the ability to proliferate. Therefore, Pantum L24 contains a mitogen (phytohemagglutinin, PHA) specifically acting on the cell cycle. It is a modified formulation of RPMI 1640.

Pantum T64 for tumor cells

Pantum T64 is specially developed for culture of tumor cells to stimulate the growth of this cell type. It is a modification of RPMI 1640.

Pantum 386 ⁽³⁾	500 ml	P04-00386	NEW
Pantum 586A ⁽³⁾	500 ml	P04-00586	NEW
Pantum L24 ⁽³⁾	500 ml	P04-00024	NEW
Pantum T64 ⁽³⁾	500 ml	P04-00064	NEW

Hepatopan Human Hepatocyte Medium

Description

Like every human organ, the liver consists of a complicated compound of different cells with varying functions. Hepatocytes represent – in terms of figures – with 75 % of the total number of liver cells the most important component. The metabolism, this means the chemical transformation of almost all substances which are taken in by the body, takes place in the liver.

Composition

The hepatocyte medium from PAN-Biotech is supplied as a basal medium with four supplements (storage of the supplements at -20° C). The supplements have to be added to the medium before use. The medium does not contain fetal bovine serum.

Hepatopan with 4 supplements ⁽³⁾ 500 ml P04-00600	Hepatopan with 4 supplements(3)	500 ml	P04-00600
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Melanopan Melanocyte Medium

Description

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Melanocytes are embedded in the basal and spike cell layer of the epidermis. They produce the pigment melanin and take care of the protective function of the skin against UV damages. If the solar radiation is too strong, the melanocytes are damaged and can develop into tumour cells.

Composition

The melanocyte growth medium from PAN-Biotech is supplied as a basal medium with seven supplements (storage of the supplements at -20° C). The supplements have to be added to the medium before use. The medium does not contain fetal bovine serum.

Melanopan with 7 supplements(3)	500 ml	P04-740500	
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www.pan-biotech.com

1) usually on stock, (2) minimum order 10 l, (3) available upon request



Neuropan Neuronal Cell Medium

Neuropan Basal Medium

Neuropan basal medium supports the growth of hippocampus cells and many other neuronal cells of the central nervous system. A feeder layer of astrocytes is not required. Neuropan basal medium does not contain glutamate which should be added for the initial culture (25 $\mu M).$ Before use, Neuropan basal medium is supplemented with serum or for a serum-free culture with Neuropan 27 or NS21 Supplement.

Neuropan 27 is a concentrate for the serum-free cultivation of neural cells.

NS21 Supplement

To culture neurons in the absence of serum, defined supplements such as B27 are widely used. However, available supplements exhibit some variability in their capability to support neurons in culture. NS21 Supplement is a newly developed serum substitute for neuronal cultures of cells from the central and peripheral nervous system.

Neuropan-Basal Media (Basicmedia) ⁽²⁾	500 ml	P04-00900
Neuropan 27 supplement $20x^{(2)}$	100 ml 10 ml	P07-07100 P07-07010
Neuropan 27 supplement $50x^{(2)}$	100 ml 10 ml	P07-07200 P07-07210
Neuropan 27 supplement 20x without Antioxidant ⁽²⁾	100 ml 10 ml	P07-10100 P07-10010
Neuropan 2 supplement 100x ⁽²⁾	100 ml 10 ml	P07-11100 P07-11010
NS21 Supplement 50x sterile ⁽³⁾	10 ml	P07-20021
NS21 Supplement 50x non-sterile ⁽³⁾	10 ml	P07-20021 P07-20001

Stempan ES-Cell Medium

Description

Stem cells are non-specialized cells with the ability (potency) to develop into different organo-typic cell types (e. g. heart, nerve, blood, muscle and cartilage cells). Depending on their origin, they are divided into embryonic and adult stem cells.

Composition

For the cultivation of embryonic stem cells, PAN-Biotech has developed a complete ready-to-use medium. The medium contains fetal bovine serum.

Stempan DMEM ⁽²⁾ with L-Glutamine with 3.7 g/L NaHCO ₃ without LIF	500 ml	P08-50500
Stempan GMEM ⁽²⁾ with L-Glutamine with 2.75 g/L NaHCO ₃ without LIF	500 ml	P08-50600

EMEM Fibroblasts Fibroblast Medium

Description

Based on EMEM, this medium was supplemented with amino acids and vitamins and optimized for an improved growth of fibroblasts.

For the cultivation of fibroblasts, this medium has to be supplemented with 10% FBS before use.

EMEM Fibroblasts ⁽¹⁾	500 ml	P04-08049
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) usually on stock, (2) minimum order 10 l, (3) available upon reques



Amniopan Prenatal Cytogenetics Medium

Description

Amniopan is a complete ready-to-use medium intended for in vitro diagnostic use with a short term culture of human fetal cells from amniotic fluid or chorion villi biopsy (CVS) material for a standardized application in cytogenetic studies.

Amniopan is intended for in vitro use and has been designed for establishing cultures of human fetal cells from amniotic fluid or chorion villi biopsies (CVS), which then can be used in karyotyping, fluorescence in-situ hybridisation (FISH) or other cytogenetic procedures. The media formulation of Amniopan was optimized on human fetal cells from amniotic fluid and CVS, with special emphasis on fast attachment of cells to the cell culture substrate and efficient cell growth to facilitate rapid diagnostic findings.

Composition

Amniopan is supplied frozen as a complete medium, ready-to-use in a 100 ml format. It is based on alpha-MEM and contains antibiotics, L-glutamine, foetal bovine serum (FBS), hormones and growth factors.

Suitability

Amniopan is a complete medium (ready-to-use) for the cultivation of human fetal cells from amniotic fluid and chorion villi biopsy samples. It is suitable for a rapid expansion of amniotic cells in order to investigate chromosomal disorders. The number and quality of metaphases in Amniopan are significantly higher and independent of individual batches as compared to other media.

Amniopan ⁽¹⁾	100 ml	P04-70100
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Amniopan S2 Prenatal Cytogenetics Medium

Description

Amniopan S2 is a complete ready-to-use medium intended for in vitro diagnostic use with a short term culture of human fetal cells from amniotic fluid or chorion villi biopsy (CVS) material for a standardized application in cytogenetic studies.

Amniopan S2 is intended for in vitro use and has been designed for establishing cultures of human fetal cells from amniotic fluid or chorion villi biopsies (CVS), which then can be used in karyotyping, fluorescence in-situ hybridisation (FISH) or other cytogenetic procedures.

The media formulation of Amniopan S2 was further optimized on human fetal cells from amniotic fluid and CVS, with special emphasis on fast attachment of cells to the cell culture substrate and efficient cell growth to facilitate rapid diagnostic findings.

Composition

Amniopan S2 is supplied frozen as a complete medium, ready-to-use in a 100 ml format. It is based on alpha-MEM and contains antibiotics, L-glutamine, fetal bovine serum (FBS), hormones and an increased amount of growth factors.

Suitability

Amniopan S2 is a ready-to-use medium for the cultivation of human fetal cells from amniotic fluid and chorion villi biopsy samples. It is suitable for a rapid expansion of amniotic cells in order to investigate chromosomal disorders. The number and quality of metaphases in Amniopan S2 are significantly higher and independent of individual batches as compared to other media.

Amniopan S2 ⁽¹⁾	100 ml	P04-70101	
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1) usually on stock, (2) minimum order 10 l, (3) available upon reques



Marrowpan Marrow Cell Medium

Description

Marrowpan is a complete ready-to-use medium intended for in vitro diagnostic procedures with a short term culture of bone marrow and other hematopoietic cells for cytogenetic studies.

Marrowpan is intended for in vitro use and has been designed for establishing cultures of bone marrow and leukemic blood cells, which then can be used in karyotyping, fluorescence in-situ hybridisation (FISH) or other cytogenetic procedures.

Marrowpan can be used as a neutral medium to culture different haematopoietic cells (myeloid and lymphoid lineages) present in bone marrow or leukemic blood samples. Marrowpan is also used together with a mitogen specific for B or T lymphocytes where these particular lineages are being investigated.

Composition

Marrowpan is supplied frozen as a complete medium, ready-to-use in a 100 ml format. It is based on alpha-MEM and contains antibiotics, L-glutamine, fetal bovine serum (FBS), hormones and growth factors.

Suitability

Marrowpan is a complete medium (ready-to-use) for the cultivation of cells from peripheral blood or bone marrow. It is suitable for a rapid expansion of blood cells in order to investigate leukemic diseases (e.g. ALL, AML, CLL, CML, MPN, MDS). The number and quality of metaphases in Marrowpan are significantly higher and independent of individual batches as compared to serum-containing media.

Marrowpan ⁽¹⁾ 100 ml P04-70200

Marrowpan S2 Marrow Cell Medium

Description

Marrowpan S2 is a complete ready-to-use medium intended for in vitro diagnostic procedures with a short term culture of bone marrow and other hematopoietic cells for cytogenetic studies. Marrowpan S2 is intended for in vitro use and has been designed for establishing cultures of bone marrow and leukemic blood cells, which then can be used in karyotyping, fluorescence in-situ hybridisation (FISH) or other cytogenetic procedures.

Marrowpan S2 can be used as a neutral medium to culture different haematopoietic cells (myeloid and lymphoid lineages) present in bone marrow or leukemic blood samples. Marrowpan S2 is also used together with a mitogen specific for B or T lymphocytes where these particular lineages are being investigated.

Compositio

Marrowpan S2 is supplied frozen as a complete medium, ready-to-use in a 100 ml format. It is based on alpha-MEM and contains antibiotics, L-glutamine, Foetal Bovine Serum (FBS), hormones and an increased amount of growth factors.

Suitability

Marrowpan S2 is a ready-to-use medium for the cultivation of cells from peripheral blood or bone marrow. It is suitable for a rapid expansion of blood cells in order to investigate leukemic diseases (e.g. ALL, AML, CLL, CML, MPN, MDS). The number and quality of metaphases in Marrowpan S2 are significantly higher and independent of individual batches as compared to serum-containing media.

Marrowpan S2 ⁽¹⁾	100 ml	P04-70201
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1) usually on stock, (2) minimum order 10 l, (3) available upon request



Serum-free Stem Cell Media

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Introduction

Research and development in the field of stem cell biology has been tremendously advanced in the last decade. Today, some cell types are used in clinical studies or applications, and several more are close to being employed in cellular therapy. One important aspect for any application of stem and progenitor cells in patients is the isolation and expansion of these cells under defined conditions. For this purpose, the presence of FBS in cell cultures is undesirable. PAN-Biotech is offering a full range of serum-free media for stem and progenitor cells for the most important fields of research and development. Some of these stem cell media are free of animal-derived components, enabling the culture of cells in conditions close to clinical application.

In addition to hematopoietic stem and progenitor cells, also other types of stem and progenitor cells (e.g. mesenchymal stem cells, endothelial progenitor cells, and very small embryonic-like stem cells) circulate under steady-state conditions at detectable levels in peripheral blood, with their numbers increasing in response to stress, inflammation, tissue organ injury (e.g. myocardial infarction, stroke, or colitis), or mobilizing agents (e.g. colony-stimulating factors, G-CSF, GM-CSF).

colony-stimulating factors, G-CSF, GM-CSF). Human mesenchymal stem stells (hMSC) have gained attention as one of very few cell types used clinically for cell therapy and tissue engineering due to their immuno-modulatory as well as their regenerative potential. MSC can be isolated from various sources: e.g. bone marrow, adipose tissue, or human umbilical cord. MSC have the capability to differentiate in vitro into connective tissue cells such as adipocytes, chondrocytes, and osteoblasts. The umbilical cord has been the most popular source for easy to obtain stem cells. Hematopoietic stem cells harvested from cord blood have been successfully used for the treatment of diseases. Stem cell populations have also been reported in other compartments of the umbilical cord, amnion, sub-amnion, perivascular region, Wharton's jelly, umbilical blood vessel adventitia and endothelium. Some differences in stemness characteristics between compartments have been found. New blood vessel formation occurs via angiogenesis or vasculogenesis, a process thought to be restricted to embryonic development. In 1997, postnatal vasculogenesis has been proposed as an important mechanism for angiogenesis via blood or bone marrow-derived circulating progenitor endothelial cells (PEC) (Asahara et al. Science). From thereon, PECs have been extensively studied as potential cell therapy for the repair of damaged blood vessels. Animal studies clearly demonstrated that administration of PECs partially rescued cardiovascular dysfuntion or myocardial injury.

Embryonic stem cells (ESCs) derived from the inner cell mass of a blastocyst are pluripotent stem cells with unique properties of self-renewal. ESCs can divide indefinitely in vitro, while maintaining the capacity to generate all cell types of an adult organism. The unique identity of ESCs is governed by a network of transcriptional factors along with epigenetic factors. The epigenetic status of ESCs features an open chromatin structure with characteristic histone and DNA modification profiles.

Somatic cells can acquire ESC properties through nuclear reprogramming. Three major approaches, including somatic cell nuclear transfer, cell fusion, and forced introduction of defined transcription factors have been established to reprogram somatic cells to pluripotency. The latter approach was first reported by Yamanaka et al. in 2006, who demonstrated that the expression of combined transcription factors, Oct4, Sox2, Klf4 and c-Myc is reprogramming somatic cells into ESC-like cells, termed induced pluripotent stem cells (iPS cells). Since this initial report, the technology has attracted great attention and motivated numerous investigations because of its tremendous potential for regenerative medicine. IPS cells have been shown to be highly similar to ESCs, in terms of transcription programs, chromatin modification profiles and global chromatin configuration. Functionally, at least some of the iPS cells have a developmental potential equivalent to ESCs in many aspects, such as the expression of certain stem cell genes and proteins, chromatin methylation patterns, doubling time, embryoid body and teratoma formation, viable chimera formation, and potency and differentiability. The full extent of their relation to natural pluripotent stem cells is still being assessed. Already, iPSCs are a widely accepted advance in stem cell research, as they may allow researchers to obtain pluripotent stem cells for therapeutic application, without the controversial use of embryos. Because iPS cells can be developed from a patient's own somatic cells, it is believed that treatment of iPS cells would avoid immunogenic responses. IPS cells have become an alternative cell source for transplantation.

Stem and progenitor cells as well as induced pluripotent stem cells are thus attractive autologous or allogenic agents for the treatment of malignant and non-malignant hematopoietic and non-hematopoietic disorders.





PowerStem ESPro1 with LIF ⁽³⁾	100 ml Kit	P04-7701K
	500 ml Kit	P04-77010K
PowerStem ESPro1 without LIF(3)	100 ml Kit	P04-7751K
	500 ml Kit	P04-77510K
PowerStem ESPro2 with LIF ⁽³⁾	100 ml Kit	P04-7702K
	500 ml Kit	P04-77020K
PowerStem ESPro2 without LIF(3)	100 ml Kit	P04-7762K
	500 ml Kit	P04-77620K
PowerStem EST ⁽³⁾	100 ml Kit	P04-77210K
	500 ml Kit	P04-77250K
PowerStem HE1 ⁽³⁾	100 ml Kit	P04-7711K
	500 ml Kit	P04-77110K
PowerStem HE2 ⁽³⁾	100 ml Kit	P04-7712K
	500 ml Kit	P04-77120K
PowerStem iPS1 ⁽³⁾	100 ml Kit	P04-7713K
	500 ml Kit	P04-77130K
PowerStem iPS2 ⁽³⁾	100 ml Kit	P04-7714K
	500 ml Kit	P04-77140K
PowerStem MSC1 ⁽³⁾	100 ml Kit	P04-77310K
	500 ml Kit	P04-77350K
PowerStem HPSC ⁽³⁾	100 ml Kit	P04-77410K
	500 ml Kit	P04-77450K
PowerStem PEC1 ready-to-use ⁽³⁾	500 ml	P04-777500
PowerStem PEC1 kit ⁽³⁾	500 ml Kit	P04-77750K

www.pan-biotech.com

PowerStem ESPro1

PowerStem ESPro1 is an easy to use serum-free medium for cultivation of embryonic stem cells of mice (mES cells). These pluripotent cells are derived from blastocysts and they can be established to a permanent cell culture. After injection into blastocysts in chimeras, they can form all tissues, including germ cells. In PowerStem ESPro1, the mES cells largely maintain their undifferentiated state and can be integrated into the germ line.

Composition

PowerStem ESPro1 contains purified proteins, lipids, salts, amino acids, trace elements, attachment factors, hormones and growth factors in an optimized formulation. PowerStem ESPro1 is fully chemically defined and contains no peptones or hydrolysates.

Please note: Supplemented PowerStem ESPro1 contains LIF in a concentration of 10 µg/l. If higher levels of LIF are required for your experimental setting, please add additional LIF to the medium.

Suitability

Serum-free cultivation of embryonic stem cells of mice (mES cells), while maintaining the undifferentiated state. PowerStem ESPro1 is especially designed for the serum-free generation of knockout-mice from genetically modified mES cells. PowerStem ESPro1 has also been proven to support the serum-free cultivation and expansion of tumor progenitor cells.

Please note: For research use only, not for therapeutic or diagnostic use.

Special advantages

PowerStem ESPro1 allows the cultivation and expansion of mouse embryonic stem cells (mES cells) under serumfree conditions. It is fully defined in its composition and thus enables constant and comparable experimental conditions resulting in highly reproducible data. The mES cell culture can be established without the usual feeder layer (primary fibroblasts), cells show a high proliferation rate and largely retain an undifferentiated state. By adding specific differentiation factors, mES cells can differentiate in vitro to the desired cell types (e.g. nerve cells, muscle cells, endothelial cells, etc.). Following injection into blastocysts, they can form all tissues in chimeras. Therefore it is possible to generate animals whose genome has been manipulated previously in a cell culture (e.g. knock-out / knock-in mice).

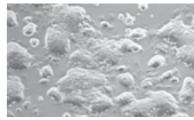
Please note: For differentiation studies LIF supplement must be omitted.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for PowerStem ESPro1. In addition, instructions for use can also be found at www.pan-biotech.com.



mES-cells in PowerStem ESPro1



JM8-cells in PowerStem ESPro1



mES-cells in medium with 10% FBS

PowerStem ESPro1	100 ml Kit	P04-7701K
with LIF ⁽³⁾	500 ml Kit	P04-77010K
PowerStem ESPro1	100 ml Kit	P04-7751K
without LIF ⁽³⁾	500 ml Kit	P04-77510K





PowerStem ESPro2

PowerStem ESPro2 is a serum-free medium for cultivation and expansion of embryonic stem cells of mice (mES cells). PowerStem ESPro2 is especially designed to proliferate and expand mES cells without differentiation. To differentiate the proliferated mES cells into different cell types the relevant protocols and differentiation factors can be used.

Composition

PowerStem ESPro2 contains purified proteins, lipids, salts, amino acids, trace elements, attachment factors, hormones and growth factors in an optimized formulation. PowerStem ESPro2 is fully chemically defined and contains no peptones or hydrolysates.

Please note: Supplemented PowerStem ESPro2 contains LIF in a concentration of $10\mu g/l$. If higher levels of LIF are required, please add additional LIF to the medium.

Suitability

PowerStem ESPro2 is especially designed for the serum-free cultivation of murine embryonic stem cells (mES cells), while maintaining the undifferentiated state. PowerStem ESPro2 is suitable for the serum-free generation of knockout-mice from genetically modified mES cells. PowerStem ESPro2 has also been proven to support the serum-free cultivation and expansion of tumor progenitor cells.

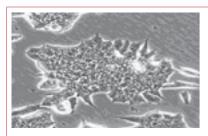
Please note: For research use only, not for therapeutic or diagnostic use

Special advantages

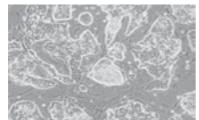
PowerStem ESPro2 allows the cultivation and expansion of mouse embryonic stem cells (mES cells) under serumfree conditions. It is fully defined in its composition and thus enables constant and comparable experimental conditions resulting in highly reproducible data. The mES cell culture can be established without the usual feeder layer (primary fibroblasts), cells show a high proliferation rate and largely retain an undifferentiated state. By adding specific differentiation factors, mES cells can differentiate in vitro to the desired cell types (e.g. nerve cells, muscle cells, endothelial cells, etc.).

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for PowerStem ESPro2. In addition, instructions for use can also be found at www.pan-biotech.com.



mES-cells in PowerStem ESPro2



mES-cells in PowerStem ESPro2

www.pan-biotech.com



ES-cells in medium with 10% FBS

PowerStem ESPro2	100 ml Kit	P04-7702K
with LIF(3)	500 ml Kit	P04-77020K
PowerStem ESPro2	100 ml Kit	P04-7762K
without LIF(3)	500 ml Kit	P04-77620K

(1) usually on stock, (2) minimum order 10 l, (3) available upon reques



PowerStem EST

PowerStem EST is a serum-free system for the cultivation and proliferation of undifferentiated mouse embryonic stem cells (mES cells) and their subsequent differentiation into beating myocardial cells (e.g. for the embryonic stem cell test EST). The EST has been formally validated by the European Centre for Validation of Alternative Methods (ECVAM) as an acceptable in vitro embryotoxicity assay. The in vitro embryonic stem cell test (EST) allows for categorisation of the embryo-toxic potential of chemicals and drug candidates. For the screening process of newly developed chemicals and pharmaceuticals, a prediction model was developed based on the inhibition of differentiation of murine embryonic stem cells into cardiomyocytes.

The application of the EST for chemical testing reduces time, testing costs and the amount of animal experimentation for embryo-toxicity tests.

Composition

PowerStem EST medium kit is composed of a complex basal medium containing salts, amino acids, vitamins, and micronutrients to which a serum-free supplement (PowerStem EST growth supplement) consisting of a mixture of proteins, growth factors and hormones is added immediately prior to use. For sustainment in undifferentiated condition and growth of ES cells, mouse leukemia inhibitory factor (mLIF, 1000 U/ml) is added to the supplemented basal medium (PowerStem EST LIF supplement). For differentiation into beating myocardial cells, a mix of differentiation factors (PowerStem EST differentiation supplement) is added to the supplemented basal medium (without mLIF).

Suitability

Cardiomyocytes differentiated from stem cells can be used for a multitude of purposes:

- Use in basic research for examining early development processes needed for functional cardiogenesis in vitro
- Testing chemicals and pharmaceutical ingredients for mutagenicity, cytotoxicity and embryotoxicity (embryonic stem cell test, EST)
- Screening of anti-angiogenetic substances
- Electrophysiological analyses for investigating cardio-active drugs
- Development of new active ingredients

PowerStem EST ⁽³⁾ 100 ml Kit P04-77210K 500 ml Kit P04-77250K	EST ⁽³⁾	100 1111 1111		
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The basal medium is used for both, proliferation and differentiation; defined factors are added according to the objective – sustainment and growth or differentiation of FS cells

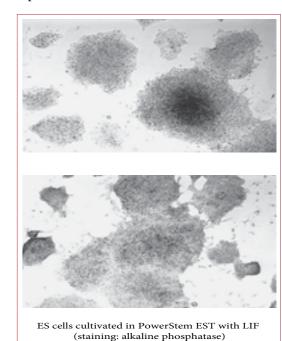
Special advantages

Traditionally, in vitro differentiation of mouse embryonic stem cells takes place using fetal bovine serum (FBS). It has been shown that the use of FBS is a limiting factor for successful differentiation of ES cells into cardiomyocytes. Some batches of FBS result in poor differentiation, while some batches may not allow differentiation at all. The search for suitable FBS batches and the dramatic variability makes the differentiation of ES cells with serum-containing media a time and money consuming exercise.

In contrast, it has been demonstrated that the number of differentiated ES cells is substantially increased under serum-free conditions, and the rate of differentiation is quite stable. The PowerStem EST medium kit successfully stimulates the expansion of undifferentiated ES-cells and promotes their subsequent differentiation into beating myocardial cells under serum-free conditions, resulting in highly comparable findings from standardized experiments.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for PowerStem EST. In addition, instructions for use can also be found at www.pan-biotech.com.



(1) usually on stock, (2) minimum order 10 l, (3) available upon reque



PowerStem HE1

PowerStem HE1 is a specialized serum-free medium for the cultivation and expansion of human embryonic stem cells (hES cells). Pluripotent human embryonic stem cells have the capacity to differentiate into all of the somatic cell types and therefore hold great promise for regenerative medicine. Even after long-term culture, cells maintained on Matrigel or Laminin retain a normal karyotype and a stable proliferating rate.

PowerStem HE1 basal medium and PowerStem HE1 growth supplement are guaranteed stable for 12 months when properly stored. PowerStem HE1 complete medium (basal + supplement) is stable for 1 month when stored in the dark at 2-8° C. We do not recommend using the complete medium beyond 1 month.

Composition

PowerStem HE1 contains purified proteins, lipids, salts, amino acids, trace elements, attachment factors, hormones and growth factors in an optimized formulation. PowerStem HE1 is fully defined and contains no peptones or hydrolysates.

Please note: PowerStem HE1 contains b-FGF in a concentration of 20 μ g/l. If higher b-FGF levels are required, please add additional b-FGF to the medium.

Suitability

Serum-free cultivation of human embryonic stem cells (hES cells), while maintaining an undifferentiated state.

Please note: For research use only, not for therapeutic or diagnostic use.

Special advantages

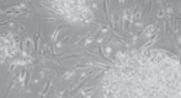
PowerStem HE1 allows the cultivation and expansion of hES cells under serum-free conditions. It is fully defined in its composition and thus enables constant and comparable experimental conditions resulting in highly reproducible data. The hES cells can be cultivated without the usual feeder layers (primary fibroblasts), they show a high proliferation rate and largely retain their undifferentiated state. By adding specific differentiation factors, hES cells can differentiate in vitro to the desired cell types (e.g. nerve cells, muscle cells, endothelial cells, etc.).

Instructions for use

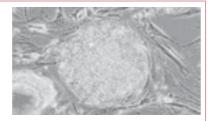
Detailed instructions will be provided with the accompanying datasheet for PowerStem HE1. In addition, instructions for use can also be found at www.pan-biotech.com.



hES-cells in PowerStem HE1



hES-cells colony in PowerStem HE1



hES-cells in medium with 10 % FBS

DavisanCtoma LIE1(3)	100 ml Kit	P04-7711K
PowerStem HE1 ⁽³⁾	500 ml Kit	P04-77110K

PowerStem HE2

PowerStem HE2 is a specialized serum-free medium for cultivation and expansion of human embryonic stem cells (hES cells). Pluripotent human embryonic stem cells have the capacity to differentiate into all of the somatic cell types and therefore hold great promise for regenerative medicine. Even after long-term culture, cells maintained on Matrigel or Laminin retain a normal karyotype and a stable proliferating rate.

PowerStem HE2 basal medium and PowerStem HE2 growth supplement are guaranteed stable for 12 months when properly stored. PowerStem HE2 complete medium (basal + supplement) is stable for 1 month when stored in the dark at 2-8° C. We do not recommend using the complete medium beyond 1 month.

Composition

PowerStem HE2 contains purified and recombinant proteins, lipids, salts, amino acids, trace elements, attachment factors, hormones and growth factors in an optimized fomulation. PowerStem HE2 is chemically defined and contains no animal-derived components.

Please note: PowerStem HE2 contains b-FGF in a concentration of 2 μ g/l. If higher b-FGF levels are required, please add additional b-FGF to the medium.

Suitabilit

Serum-free cultivation of human embryonic stem cells (hES cells), while maintaining an undifferentiated state.

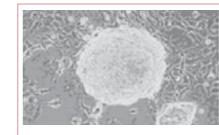
Please note: For research use only, not for therapeutic or diagnostic use.

Special advantages

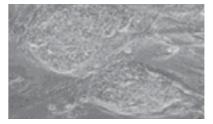
PowerStem HE2 allows the cultivation and expansion of hES cells under serum-free conditions. It is fully defined in its composition thus enabling constant and comparable experimental conditions resulting in highly reproducible data. The hES cells can be cultivated without the usual feeder layers (primary fibroblasts), they show a high proliferation rate and largely retain their undifferentiated state. By adding specific differentiation factors, hES cells can differentiate in vitro to the desired cell types (e.g. nerve cells, muscle cells, endothelial cells, etc.).

Instructions for use

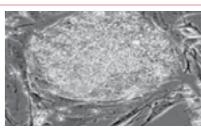
Detailed instructions will be provided with the accompanying datasheet for PowerStem HE2. In addition, instructions for use can also be found at www.pan-biotech.com.



hES-cells in PowerStem HE2



hES-cells in PowerStem HE2



hES-cells in medium with 10 % FBS

PowerStem HE2 ⁽³⁾	100 ml Kit	P04-7712K
	500 ml Kit	P04-77120K

(1) usually on stock, (2) minimum order 10 l, (3) available upon reques



1) usually on stock, (2) minimum order 10 l, (3) available upon request



PowerStem iPS1

PowerStem iPS1 is a specialized serum-free medium for the cultivation and expansion of human induced pluripotent stem cells (iPS cells). Induced pluripotent stem cells behave similar to human embryonic stem cells and have the capacity to differentiate into all of the somatic cell types and therefore hold great promise for regenerative medicine. Even after long-term culture (> 50 passages) iPS cells retain a normal karyotype and a stable proliferating rate.

PowerStem iPS1 basal medium and PowerStem iPS1 growth supplement are guaranteed stable for 12 months when properly stored. PowerStem iPS1 complete medium (basal + supplement) is stable for 2 weeks when stored in the dark at 2-8° C. We do not recommend using the complete supplemented medium beyond 2 weeks.

Composition

PowerStem iPS1 contains purified and recombinant proteins, lipids, salts, amino acids, trace elements, hormones and growth factors in an optimized formulation. PowerStem iPS1 is a defined medium and contains no animal- or human-derived substances (except human serum albumin (100 μ g/ml) as a stabilizing agent).

Please note: PowerStem iPS1 contains FGF-2 in a high concentration; it is not recommended to add additional FGF-2.

Serum-free cultivation of human induced pluripotent stem cells (iPS cells) under defined conditions, while maintaining an undifferentiated state.

Please note: For research use only, not for therapeutic or diagnostic use.

Special advantages

PowerStem iPS1 allows the cultivation and expansion of iPS cells under serum-free conditions. It is fully defined in its composition and thus enables constant and comparable experimental conditions resulting in highly reproducible data. The iPS cells can be cultivated without the usual feeder layer of primary fibroblasts, they show a high proliferation rate and largely retain their undifferentiated state. By adding specific differentiation factors, iPS cells can be differentiated in vitro to the desired cell types (e.g. nerve cells, muscle cells, endothelial cells, etc.).

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for PowerStem iPS1. In addition, instructions for use can also be found at www.pan-biotech.com

PowerStem iPS1(3)	100 ml Kit	P04-7713K
Powerstem iPST	500 ml Kit	P04-77130K

PowerStem iPS2

PowerStem iPS2 is a chemically defined serum-free medium for cultivation and expansion of human induced pluripotent stem cells (iPS cells). Induced pluripotent stem cells behave similar to human embryonic stem cells and have the capacity to differentiate into all of the somatic cell types and therefore hold great promise for regenerative medicine. Even after long-term culture (> 50 passages) iPS cells retain a normal karyotype and a stable proliferating rate.

PowerStem iPS2 basal medium and PowerStem iPS2 growth supplement are guaranteed stable for 12 months when properly stored. PowerStem iPS2 complete medium (basal + supplement) is stable for 1 week when stored in the dark at 2-8° C. We do not recommend using the complete supplemented medium beyond 1 week.

Composition

PowerStem iPS2 contains lipids, salts, amino acids, trace elements, hormones and recombinant growth factors in an optimized formulation. PowerStem iPS2 is chemically defined and contains no animal- or human-derived

Please note: PowerStem iPS2 contains a high concentration FGF-2; it is not recommended to supplement with additional FGF-2.

100 ml Kit P04-7714K PowerStem iPS2(3) 500 ml Kit P04-77140K

Serum-free cultivation of induced pluripotent stem cells (iPS cells), while maintaining an undifferentiated state.

Please note: For research use only, not for therapeutic or diagnostic use.

Special advantages

PowerStem iPS2 allows the cultivation and expansion of iPS cells under serum-free conditions. It is fully defined in its composition thus enabling constant and comparable experimental conditions resulting in highly reproducible data. The iPS cells can be cultivated without the usual feeder layer of primary fibroblasts, they show a high proliferation rate and largely retain their undifferentiated state. By adding specific differentiation factors, iPS cells can be differentiated in vitro to the desired cell types (e.g. nerve cells, muscle cells, endothelial cells, etc.).

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for PowerStem iPS2. In addition, instructions for use can also be found at www.pan-biotech.com





PowerStem MSC1

PowerStem MSC1 is an easy to use xeno-free medium without animal derived components (ADCF) for cultivation and proliferation of human mesenchymal stem cells (hMSC). PowerStem MSC1 is especially designed for the proliferation of human mesenchymal stem cells without differentiation. PowerStem MSC1 supports long-term growth of MSC and preserves their multi-lineage potential. In addition, MSC cultured in PowerStem MSC1 expands faster and shows a significant reduction in hematopoietic cell contamination at early passages compared to serum-based media. To differentiate the proliferated MSC into different cells types the relevant protocols and differentiation factors should be used.

Both the PowerStem MSC1 basal medium and PowerStem MSC1 growth supplement are guaranteed stable for 6 months when properly stored. PowerStem MSC1 complete medium (basal + supplement) is stable for 1 month when stored in the dark at 2-8° C. We do not recommend using the complete medium beyond one month. Do not freeze complete PowerStem MSC1 medium.

Composition

PowerStem MSC1 contains salts, amino acids, trace elements, hormones, growth factors, and enriched human proteins and lipids in an optimized formulation. PowerStem MSC1 is free of animal derived components (ADCF, xenofree) and contains no undefined peptones or hydrolysates.

Suitabili

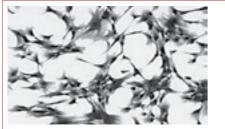
Serum-free cultivation of human mesenchymal stem cells (hMSC) while maintaining the undifferentiated state and multi-lineage potential. Please note: For research use only, not for therapeutic or diagnostic use.

Special advantages

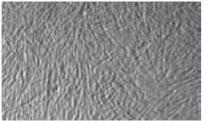
PowerStem MSC1 allows the cultivation of human mesenchymal stem cells under xeno-free conditions. It is free of animal or human serum and thus enables constant and comparable experimental conditions resulting in highly reproducible data. PowerStem MSC1 is completely free of animal components (ADCF, xeno-free) and thus suitable for a research approach in regenerative medicine and tissue engineering. By adding specific differentiation factors, MSC can differentiate in vitro to the desired cell types (bone, cartilage, adipose tissue etc.).

Instructions for use

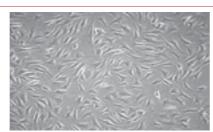
Detailed instructions will be provided with the accompanying datasheet for PowerStem MSC1. In addition, instructions for use can also be found at www.pan-biotech.com. For more instructions please see instruction manual for isolation and culture of hMSC.



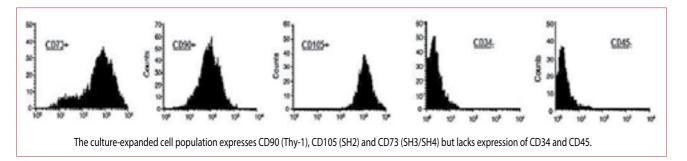
Sub-confluent hMSC in PowerStem MSC1



Confluent hMSC in PowerStem MSC1



hMSC in medium with 10% FBS



PowerStem $MSC1^{(3)}$

100 ml Kit 500 ml Kit P04-77310K P04-77350K

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



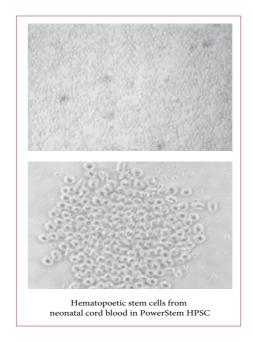
PowerStem HPSC

PowerStem HPSC is a specialized serum-free medium for the cultivation and expansion of human hematopoietic stem cells (HPSC) and cells of myeloid lineage in suspension culture. Hematopoietic stem cells are CD34+, which are the earliest hematopoietic stem cells identifiable in bone marrow, peripheral blood and neonatal cord blood. By adding one or more differentiation factors or changing culturing conditions, HPSC can be induced to differentiate into different types of hematopoietic lineage cells.

PowerStem HPSC basal medium, PowerStem HPSC growth supplement and PowerStem HPSC cytokine supplement are guaranteed stable for 12 months when properly stored. PowerStem HPSC complete medium (basal + supplements) is stable for 3 months when stored in the dark at 2-8° C. We do not recommend using the complete medium beyond 3 months.

Composition

PowerStem HPSC contains purified proteins, lipids, salts, amino acids, trace elements, attachment factors, hormones and growth factors in an optimized formulation. PowerStem HPSC is fully defined and contains no FBS.



Suitabilit

Serum-free cultivation and expansion of human hematopoietic CD34+ stem cells from bone marrow, peripheral blood and neonatal cord blood.

Please note: For research use only, not for therapeutic or diagnostic use.

Special advantages

PowerStem HPSC allows the cultivation and expansion of human hematopoietic CD34+ stem cells and cells of myeloid lineage under serum-free conditions. It is fully defined in its composition and thus enables constant and comparable experimental conditions with easily reproduceible results. The hematopoietic stem cells can be cultivated without stromal cells, they show a high proliferation rate and largely retain their undifferentiated state. By adding specific differentiation factors, hematopoietic cells can be differentiated in vitro to different types of hematopoietic lineage cells.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for PowerStem HPSC. In addition, instructions for use can also be found at www.pan-biotech.com.

References

Horschitz S et al. (2010) Generation of neuronal cells from human peripheral blood mononuclear cells. Neuro Report 21:185.

PowerStem HPSC ³⁾	100 ml Kit 500 ml Kit	P04-77410K P04-77450K	
------------------------------	--------------------------	--------------------------	--

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



PowerStem PEC1

Endothelial cells line blood vessels and the internal cavities of the heart. They display a flattened, polygonal form and adhere to each other by desmosomes and tight-junctions. With a total number of about 10^{12} cells, the endothelium is one of the biggest organs of the body and plays a key role in many physiological and pathophysiological processes. A number of factors control proliferation and apoptosis of endothelial cells, thereby regulating maintenance, degeneration, or regeneration of blood vessels.

New blood vessel formation occurs via angiogenesis or vasculogenesis, a process restricted to embryonic development. In 1997, postnatal vasculogenesis has been proposed as an important mechanism for angiogenesis via blood or bone marrow derived circulating progenitor endothelial cells (PEC) (Asahara et al. Science 1997). PEC have been extensively studied as potential cell therapy for the repair of damaged blood vessels. Animal studies clearly demonstrated that administration of PEC partially rescued cardiovascular dysfuntion or myocardial injury with evidence for PEC contribution to new vessel growth.

While controversy exists as to the identity of endothelial cell progenitors, recently a PEC population has been identified which shows expression of typical endothelial as well as progenitor markers (Ingram et al. Blood. 2004;104:2752-2760). Importantly, these cells have been tested for a high proliferative potential in clonogenic assays and characterized by formation of functional blood vessels in vivo (Yoder et al. Blood. 2007;109:1801-1809).

With endothelial cell progenitors rapidly moving into the field of interest for vascular tissue engineering with potential therapeutic application, the presence of whole animal serum or animal-derived components in culture media is undesirable for a cell therapeutic approach.

Description

PowerStem PEC1 ready-to-use (P04-777500) is a specially developed medium for a serum- and xenofree in vitro culture of human progenitor endothelial cells (hPEC) containing all components necessary for optimal colony formation, clonogenic growth, and rapid proliferation. It is designed for use in an incubator at 37° C with a 5% CO₂ atmosphere.

PowerStem PEC1 kit (P04-77750K) is provided with supplements (pre-screened and tested for progenitor cells) in separate sterile packing. This will enable the user to prepare a medium for special application. For example, VEGF, FGF-2, or other components may be omitted from the complete medium for specific experimental settings. Please note that such a formulation will not promote optimal cell growth. Therefore, this composition can not be used for routine long-term culture of PEC. Please make sure that sterility is not compromised when adding individual components to prepare complete medium. The medium should be carefully but thoroughly mixed after addition of all components to assure a homogeneous solution. Store basal or complete medium at 2 – 8° C and store supplements at -20° C. Expiry: 6 months.

Basal medium (w/o supplements) or complete/ready-to-use medium should not be frozen!

Please note: For research use only, not for therapeutic or diagnostic use.

Instructions for use

Detailed instructions will be provided with the accompanying datasheet for PowerStem PEC1. In addition, instructions for use can also be found at www.pan-biotech.com.

References

www.pan-biotech.com

- a) Asahara T et al. (1997) Isolation of putative progenitor endothelial cells for angiogenesis. Science 275:964
- b) Ingram DA et al. (2004) Identification of a novel hierarchy of endothelial progenitor cells using human peripheral and umbilical cord blood. Blood 104:2752
- c) Yoder MC et al. (2007) Redefining endothelial progenitor cells via clonal analysis and hematopoietic stem/progenitor cell principals. Blood 109:1801

 PowerStem PEC1 ready-to-use⁽³⁾
 500 ml
 P04-777500

 PowerStem PEC1 kit
 500 ml Kit
 P04-77750K

(1) usually on stock, (2) minimum order 10 l, (3) available upon request





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Description

Maximum efficiency without loss of quality

PAN-Biotech reagents are tested according to the highest possible quality standards. All liquid reagents are dissolved according to in-house specifications, sterilized

and filtered at 0.2 $\mu m.$ Before final release, the reagents undergo extensive quality tests e. g. sterility, pH value, osmo, endotoxin.

Different Media Supplements

HAT supplement (50x) ⁽¹⁾	100 ml	P07-02100
HT supplement (50x) ⁽¹⁾	100 ml	P07-01100
Hepes buffer 1M ⁽¹⁾	100 ml 500 ml	P05-01100 P05-01500
Hepes - Sodium salt ⁽¹⁾	100 g 500 g	P05-01100P P05-01500P
Sodium bicarbonate 7.5 % ⁽¹⁾	100 ml	P04-44100
Sodium pyruvate 100 mM ⁽¹⁾	100 ml	P04-43100
ITS solution I (100x) ⁽¹⁾	5 ml 10 ml	P07-03100 P07-03110
ITS solution II (100x) ⁽¹⁾	5 ml 10 ml	P07-03200 P07-03210
ITS solution IV (100x) ⁽³⁾ (w: Linoleic acid, BSA)	5 ml 10 ml	P07-03400 P07-03410
Insulin human rec. 10 mg/ml solution ⁽¹⁾	10 ml	P07-04300
Insulin human recombinant ⁽¹⁾	100 mg	P07-04200
Sterile Water for cell culture ⁽¹⁾	500 ml 1 L 20 L	P04-991500 P04-991000 P04-992000
β-Mercapthoethanol 50 mM in PBS ⁽¹⁾	20 ml 100 ml	P07-05020 P07-05100
Tryptose phosphate (50x) ⁽³⁾ 130 g/l Tryptose phosphate in distilled water	100 ml	P10-031100
Pluronic F-68 10 % ⁽¹⁾	100 ml	P08-02100
Human Transferrin apo ⁽¹⁾	100 mg 500 mg 1 g	P06-21100 P06-21500 P06-21000
Demecolcin solution 10 μg/ml ⁽¹⁾	10 ml	P07-91010
Sodium chloride solution 0.9 % ⁽¹⁾	500 ml	P05-39500

1) usually on stock, (2) minimum order 10 l, (3) available upon request



Buffered Salt Solutions

Dulbecco's Phosphate Buffered Salt Solution

Composition

	Components	mg/L
Inorga-	Potassium chloride	200.00
nic Salts	Potassium dihydrogen	200.00
	phosphate	
	Sodium chloride	8000.00
	di-Sodium hydrogen phosphate	1150.00
	anhydrous	
	Calcium chloride x 2H2O	133.00
	Magnesium chloride x 6H2O	100.00

Liquid Salt Solution

DPBS⁽¹⁾

without Ca and Mg

DPBS ⁽¹⁾ without Ca and Mg	500 ml	P04-36500
	1 L 2,5 L	P04-361000 P04-3625C
	5 L	P04-3650C
	10 L	P04-360010B
DPBS (10x) ⁽¹⁾ without Ca and Mg	500 ml	P04-53500
DPBS non-sterile ⁽¹⁾ without Ca and Mg	2,5 L	P04-362500
DPBS ⁽¹⁾		
with Ca and Mg	500 ml	P04-35500
DPBS (10x) ⁽²⁾ with Ca and Mg	500 ml	P04-37500
Powder		

Earl's Buffered Salt Solution

Composition

	Components	mg/L
Inorga-	Potassium chloride	400.00
nic Salts	Sodium chloride	680.00
	Sodium dihydrogen phosphate	140.00
	x H ₂ O	
	Calcium chloride x 2H2O	264.92
	Magnesium chloride x 6H2O	139.57
Other	D(+)-Glucose anhydrous	1000.00
Compo-	Phenol red	10.00
nents		

Liquid Salt Solution

EBSS ⁽²⁾	500 ml	P04-30500
EBSS ⁽²⁾ without Phenol red	500 ml	P04-39500
EBSS ⁽²⁾ without Ca and Mg with 2.2 g/l NaHCO ₃	500 ml	P04-31500
EBSS (10x) ⁽³⁾	500 ml	P04-38500
EBSS (10x) ⁽³⁾ without Ca and Mg without Phenol red	500 ml	P04-47500
Powder		
EBSS ⁽¹⁾	10 L 50 L	P04-30010P P04-30050P

Buffered Salt Solutions

Hank's Balanced Salt Solution

Composition

	Components	mg/L
Inorga-	Potassium chloride	400.00
nic	Sodium chloride	8000.00
Salts	di-Sodium hydrogen phosphate anhydrous	47.88
	Calcium chloride x 2H2O	185.44
	Magnesium sulphate dried	139.52
	Potassium dihydrogen phosphate	60.00
Other	D(+)-Glucose anhydrous	1000.00
Compo- nents	Phenol red	10.00

Hank's Balanced Salt Solution

Liquid Salt Solution		
HBSS ⁽²⁾	100 ml	P04-32100
with 0.35 g/l NaHCO ₃	500 ml	P04-32500
HBSS (10x) ⁽²⁾	100 ml	P04-49100
without NaHCO ₃	500 ml	P04-49500
HBSS ⁽¹⁾		
without Ca and Mg	100 ml	P04-33100
with 0.35 g/l NaHCO ₃	500 ml	P04-33500
HBSS (10x) ⁽¹⁾		
without Ca and Mg	100 ml	P04-50100
without 0.35 g/l NaHCO₃	500 ml	P04-50500
HBSS ⁽¹⁾		
without Phenol red	100 ml	P04-32105
with 0.35 g/l NaHCO ₃	500 ml	P04-32505
HBSS (10x) ⁽²⁾		
without Phenol red	100 ml	P04-49105
without NaHCO ₃	500 ml	P04-49505
HBSS ⁽¹⁾		
without Ca and Mg		
without Phenol red	100 ml	P04-34100
with 0.35 g/l NaHCO₃	500 ml	P04-34500
	1 L	P04-341000
HBSS (10x) ⁽²⁾		
without Ca and Mg		
without Phenol red	100 ml	P04-50105
without 0.35 g/l NaHCO ₃	500 ml	P04-50505
- 1		

Powder

www.pan-biotech.com

 $\begin{array}{cccc} HBSS^{(1)} & 10 \ L & P04-32010P \\ without \ NaHCO_3 & 50 \ L & P04-32050P \\ \\ HBSS^{(1)} & & & \\ without \ Ca \ and \ Mg & 10 \ L & P04-33010P \\ without \ NaHCO_3 & 50 \ L & P04-33050P \\ \end{array}$

(1) usually on stock, (2) minimum order 10 l, (3) available upon request





P04-36050P

Buffered Salt Solutions

Puck's Salt Solution A

Composition

	Components	mg/L
Inorga-	Potassium chloride	400.00
nic Salts	Sodium chloride	8000.00
Other	D(+)-Glucose anhydrous	1000.00
Compo-		
nents		

Liquid Salt Solution

 $\begin{array}{ccc} Puck's \ Salt \ Solution \ A^{(2)} & 100 \ ml & P04-51100 \\ & 500 \ ml & P04-51500 \end{array}$

Gey's Balanced Salt Solution Composition

	Components	mg/L
Inorga-	Potassium chloride	370.00
nic Salts	Sodium chloride	7000.00
	di-Sodium hydrogen phosphate	120.00
	Calcium chloride x2H2O	225.10
	Magnesium chloride x 6 H2O	210.00
	Magnesium sulphate anhydrous	34.20
	Potassium dihydrogen phosphate nahydrous	30.00
Other	D(+)-Glucose anhydrous	1000.00
Compo-	·	
nents		

Liquid Salt Solution

GBSS $^{(1)}$ 500 ml P04-48500 with 2.27 g/l NaHCO $_3$

Powder

GBSS⁽¹⁾ 10 L P04-48010P without NaHCO₃ 50 L P04-48050P

Tryode's Salt Solution

Composition

	Components	mg/L
Inorga-	Potassium chloride	200.00
nic Salts	Sodium chloride	8000.00
	Magnesium chloride anhydrous	100.00
	Calcium chloride anhydrous	200.00
	Sodium phosphate monobasic	50.00
Other	D(+)-Glucose anhydrous	1000.00
Compo-	-	
nents		

Liquid Salt Solution

Tryode's Salt Solution⁽²⁾ 500 ml P04-54500

Powder

Tryode's Salt Solution⁽¹⁾ 10 L P04-54010P without NaHCO₃ 50 L P04-54050P

Amino Acids and Vitamins

Amino Acids

BME solution (50x), without L-Glutamine $^{(3)}$	100 ml	P08-2000
L-Glutamine 200 mM ⁽¹⁾	50 ml 100 ml	P04-80050 P04-80100
Stable L-Glutamine 200 mM ⁽¹⁾ (L-Alanyl-L-Glutamine)	50 ml 100 ml	P04-82050 P04-82100
L-Glutamine ⁽¹⁾ Powder	25 g 100 g 500 g	P04-80025P P04-80100P P04-80500P
Stable Glutamine Powder ⁽¹⁾ (L-Alanyl-L-Glutamine)	10 g	P04-82010P
MEM (50x) without L-Glutamine(1)	100 ml	P08-30100
MEM (50x) with L-Glutamine ⁽³⁾	100 ml	P08-31100
MEM NEAA (100x) ⁽¹⁾	100 ml	P08-32100

Vitamins

BME vitamins ⁽³⁾	100 ml	P08-40100
MEM (100x) vitamine solution ⁽¹⁾	100 ml	P08-41100

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(1) usually on stock, (2) minimum order 10 l, (3) available upon request





(1) usually on stock, (2) minimum order 10 l, (3) available upon request

Antibiotics and Antifungal Drugs

50 x 1 ml	P06-01001
ΕO Γ1	
50 x 5 ml	P06-01005
50 ml	P06-01050
100 ml	P06-01100
50 mg	P06-01050P
100 mg	P06-01100P
25 mg	P06-01225P
50 mg	P06-01250P
10 g	P06-02010P
25 g	P06-02025P
50 x 1 ml	P06-03001
50 x 5 ml	P06-03005
50 ml	P06-03050
100 ml	P06-03100
50 x 1 ml	P06-13001
50 x 5 ml	P06-13005
50 ml	P06-13050
100 ml	P06-13100
1 g	P06-03001P
10 g	P06-03010P
25 g	P06-03025P
20 ml	P06-08020
100 ml	P06-08100
50 mg	P06-080050P
l g	P06-080100P
50 x 1 ml	P06-04001
50 x 5 ml	P06-04005
50 ml	P06-04050
100 ml	P06-04100
50 x 1 ml	P06-14001
50 x 5 ml	P06-14005
50 ml	P06-14050
100 ml	P06-14100
50 x 1 ml	P06-15001
50 x 5 ml	P06-15005
50 ml	P06-15050
100 ml	P06-15100
10 g	P06-04010P
50 g	P06-04050P
50 ml	P06-05050
100 ml	P06-05100
100 ml	P06-02100 NEW
50 ml	P06-06050
100 ml	P06-06100
10 g	P06-06010P
ū	P06-06025P
100 g	P06-06100P
	Da : 0=000
100 ml	P06-07800
	100 ml 50 mg 100 mg 25 mg 50 mg 10 g 25 g 50 x 1 ml 50 x 5 ml 50 ml 100 ml 50 x 1 ml 50 x 5 ml 50 ml 100 ml 10 g 25 g 20 ml 100 ml 50 x 1 ml 50 x 5 ml 50 mg 1 g 50 x 1 ml 50 x 5 ml 50 ml 100 ml 50 x 5 ml 50 ml 100 ml 50 x 5 ml 50 ml 100 ml 100 ml 50 x 1 ml 50 x 5 ml 50 ml 100 ml 50 x 1 ml 50 x 5 ml 50 ml 100 ml 50 x 1 ml 50 x 5 ml 50 ml 100 ml 50 x 1 ml 50 x 5 ml 50 ml 100 ml 50 x 1 ml 50 x 5 ml 50 ml 100 ml 50 x 5 ml 50 ml 100 ml

Page 100

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Antibiotics and Antifungal Drugs

Panaticin 420 50 mg/ml(3)	20 ml	P06-16020
Paneticin 420 50 mg/ml ⁽³⁾	100 ml	P06-16100
Paneticin G418 50 mg/ml ⁽³⁾	20 ml	P06-16220
Paneticin 420 Powder ⁽¹⁾	1 g 5 g 10 g	P06-16001P P06-16005P P06-16010P
Penicillin/Streptomycin ⁽¹⁾ 10,000 Units Penicillin/ml 10 mg Streptomycin/ml	50 x 1 ml 50 x 5 ml 50 ml 100 ml	P06-07001 P06-07005 P06-07050 P06-07100
Penicillin/Streptomycin/Amphotericin B Mix ⁽¹⁾ 10,000 Units Penicillin/ml 10 mg Streptomycin/ml 25 μg Amphotericin B/ml in 0.85 % saline	50 x 1 ml 50 x 5 ml 50 ml 100 ml	P06-07301 P06-07305 P06-07350 P06-07300
Penicillin G potassium salt Powder ⁽¹⁾	25 g 100 g	P06-08025P P06-08100P
Streptomycin sulphate Powder ⁽¹⁾	25 g 50 g 100 g	P06-11025P P06-11050P P06-11100P
Polymycin B sulphate 10,000 Units/ml ⁽³⁾	50 ml	P06-09050
Tiamulin 1 mg/ml ⁽³⁾	50 x 1 ml 50 x 5 ml 50 ml 100 ml	P06-12001 P06-12005 P06-12050 P06-12100
Zeocin 100 mg/ml ⁽³⁾	10 ml	P06-28010

Mycorase

Description

Mycorase has been developed to remove a broad range of different strains of mycoplasma in most cell types. Mycorase is easy to use and does not affect eukaryotic cell proliferation. It is a highly effective antibiotic solution for safe eradication of mycoplasma contamination.

Special advantages

- Ready-to-use solution
- Effective removal of mycoplasma
- No effect on cell proliferation
- Broad range of action
- Permanent cure for most cell types

Mycorase ⁽¹⁾	100 ml	P06-02100 NEW	
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) usually on stock, (2) minimum order 10 l, (3) available upon request



www.pan-biotech.com

Enzymes for Cell Dissociation

Collagenase type I

Natural balance of enzyme activity. Recommended for cell preparation from epithelial lung tissue, tissue of the urprarenal gland and adipose tissue. Store at $2-8^{\circ}$ C.

Collagenase type II

With especially high activity of clostripain and trypsin. Recommended for cell preparation from liver tissue, bone tissue, cardiac tissue, thyroid gland tissue and salivary gland tissue. Store at $2 - 8^{\circ}$ C.

Collagenase type III

Normal collagenase activity with a minimum of proteolytic activity. Especially recommended for breast tissue. Store at $2-8^{\circ}$ C.

Collagenase type IV

Selected low tryptic activity at high collagenase activity and normal clostripain level. Recommended for cell preparation from the pancreatic island. Store at $2-8^{\circ}$ C.

Collagenase type I (Worthington - USA orgin)	100 mg 1 g	LS0004194 LS0004196
Collagenase type II (Worthington - USA orgin)	100 mg 1 g	LS0004174 LS0004176
Collagenase type III (Worthington - USA orgin)	100 mg 1 g	LS0004180 LS0004182
Collagenase type IV (Worthington - USA orgin)	100 mg 1 g	LS0004186 LS0004188

Accutase

Description

Accutase is a ready-to-use cell detachment solution made of collagenolytic and proteolytic enzymes. It is used for routine detachment of adherent cells from tissue culture plates and flasks. A multitude of cell types has been successfully sub-cultured with Accutase, including fibroblasts, endothelial, vascular smooth muscle cells, as well as hepatocytes, embryonal stem cells, and many immortalized cell lines such as adherent CHO and BHK cells, HEK 293, L929, HeLa, 3T3, and others. Accutase is free of mammalian or bacterial products.

Composition

Accutase enzymes (activity > 500 U/ml) in DPBS w/o Ca/Mg with 0.5 mM EDTA and phenol red.

Suitability

Accutase can be used as a direct replacement of trypsin for cell dissociation.

Special advantages

Neutralizing of the Accutase enzymes is not required for routine cell culture passaging. The product is active at room temperature, no pre-warming required or recommended. Gentle detachment of cells for analysis of cell surface markers, transfection procedures, migration or proliferation assays, flow cytometry, and routine cell passage.

Accutase ⁽¹⁾	100 ml	P10-21100	NEW	
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(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Enzymes for Cell Dissociation

Trypsin and others

Application

For dissociation of tissue and cell monolayer cultures.

Shelf life

Solution 24 months, powder 36 months. The shelf life commences with date of production. Our trypsin is tested negative for mycoplasma.

Storage

Solution at -20° C, powder at 2 - 8° C.

Other sizes and custom formulation

Please ask for other sizes or special formulations of trypsin; in most cases we can provide a solution.

Trypsin 0.25 %/EDTA 0.02 % in PBS ⁽¹⁾	100 ml	P10-019100
without Ca and Mg with Phenol red	500 ml	P10-019500
Trypsin 0.25 %/EDTA 0.02 % in PBS ⁽¹⁾	100 ml	P10-020100
without Ca and Mg	500 ml	P10-020500
Trypsin 0.25 % in PBS ⁽¹⁾	100 ml	P10-021100
without Ca and Mg	500 ml	P10-021500
(10x) Trypsin 2.5 % in PBS ⁽²⁾	100 ml	P10-022100
without Ca and Mg	500 ml	P10-022500
Trypsin 0.05 %/EDTA 0.02 % in PBS ⁽¹⁾	100 ml	P10-023100
without Ca and Mg	500 ml	P10-023500
Trypsin 0.05 %/EDTA 0.02 % in PBS ⁽¹⁾	100 ml	P10-0231SP
without Ca and Mg with Phenol red	500 ml	P10-0235SP
(10x) Trypsin 0.5 %/EDTA 0.2 % in PBS ⁽²⁾	100 ml	P10-024100
without Ca and Mg	500 ml	P10-024500
Trypsin 0.5 %/EDTA 0.01 % in PBS ⁽²⁾	100 ml	P10-027100
without Ca and Mg	500 ml	P10-027500
Trypsin 0.25 %/1 mM EDTA 4 Na in PBS ⁽¹⁾	100 1	D10 020100
without Ca and Mg	100 ml	P10-028100
Trypsin 0.25 %/1 mM EDTA in HBSS ⁽¹⁾	100 ml	P10-029100
without Ca and Mg with Phenol red	500 ml	P10-029500
Trypsin 0.05 %/EDTA 4 Na 0.02 % in HBSS ⁽²⁾	100 ml	P10-040100
wih Phenol red	500 ml	P10-040500
Trypsin special solution (for ES-cells)(1)	100 ml	P10-100100
Trypsin Inhibitor 1 mg/ml ⁽¹⁾	100 ml	P10-033100
	25 g	P10-025025P
Trypsin powder (1:250) porcine origin ⁽¹⁾	100 g	P10-025100P
	250 g	P10-025500P
EDTA 1 % in DDC without Co and Mg(2)	100 ml	P10-026100
EDTA 1 % in PBS without Ca and Mg ⁽²⁾	500 ml	P10-026500
Dispase II neutral proteins, grade II ⁽¹⁾	100 ml	P10-032100
Dispase purified neutral protease (3)	10 mg	LS0002100

) usually on stock, (2) minimum order 10 l, (3) available upon reques



Attachment Factors

Collagen A

Application

For dissociation of tissue and cell monolayer

Storage

Solution at -20° C (frozen), powder at 2 – 8° C

Shelf life

Solution 24 months, powder 36 months. The shelf life commences with date of production. Our trypsin is tested negative for mycoplasma.

Acid-soluble collagen from bovine placenta

- Add an equal volume of sterile PBS to the collagen
- Add 1 ml per 10 cm² of culture flask and incubate at 35 37° C for 30 min
- Remove solution and wash 1x with PBS; use culture flasks immediately

In monolayer culture, normal human and murine liver cells were successfully grown for a period of up to one week, provided that the culture flasks were coated with collagen. Cell growth rates can often be improved by surface coating with attachment factors such as fibronectin, collagen, gelatine or poly-lysine. With a collagen coating, survival time of e.g. hepatocytes can be extended from one week for up to four weeks.

Storage: 2 – 8° C

Collagen A	1 x (6 x 5 ml)	P06-20030

Collagen R (type I)

0.2 % sterile solution

Type 1 rat tail collagen; 2 mg/ml in 0.1 % acetic acid. Excellent substrate for the culture of hepatocytes, fibroblasts and epithelial cells.

0.4 % sterile solution

Type 1 rat tail collagen; 4 mg/ml in 0.1 % acetic acid. Excellent substrate for the culture of hepatocytes, fibroblasts and epithelial cells.

Collagen R 0.2 % sterile solution	20 ml 100 ml	P06-20166 P06-20100
Collagen R 0.4 % sterile solution	20 ml	P06-20020

Gelatine Solution

Description

The gelatine solution is used for coating cell culture dishes. It is applied in adherent cell cultures working with e.g. endothelial cells or ES-cells.

Gelatine solution 0.1 % in PBS ⁽¹⁾	500 ml	P06-20410
Gelatine solution 2 % in PBS ⁽²⁾	100 ml	P06-25200

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Attachment Factors

Laminin Mouse

Description

This highly purified preparation of mouse Laminin I increases cell adhesion, migration, growth, and differentiation. It is composed of 111 chains with a total MW of 800 kD and is used for the coating of culture dishes.

Source:

Murine Engelbreth-Holm-Swarm (EHS) tumor

Storage Buffer:

Dulbecco's Modified Eagle's Medium with 10 µg/ml gentamycin sulfate

Storage

Store at -20° C or at -80° C in a manual defrost freezer **Purity:**

Purity > 90 % by SDS-PAGE

Specifications

Functional assays

• Supports the formation of neuronal filaments of NG108-15 cells in a neurite outgrowth assay

Sterility testing

- No bacterial or fungal growth detected after incubation at 37° C for 14 days following USP XXIV, Chapter 71 sterility testing
- No mycoplasma contamination detected by PCR
- Endotoxin concentrations < 20 EU/ml by LAL assay

Coating procedure

The recommended working concentration is 0.05 - $10~\mu g/$ cm² of growth surface (0.05 - $10~\mu g/ml$) depending on cell type.

a. Thaw stock solution on ice for several hours. Place plates on ice and pre-chill pipette tips. Distribute the solution to completely cover the bottom of the wells.

b. The following table gives suggested volume required per well:

Plate Type	Volume Laminin per Well
6 wells (or 35 mm dish)	1 ml
24 wells	200 µl
48 wells	50 μl
96 wells	20 μl

c. Incubate the plates at 37° C for 1 hour. In the laminar flow hood, remove excess liquid from the wells of the tissue culture plate.

Rinse the wells once with tissue culture medium and then add your cells.

Laminin from mouse	1 mg	P06-20501
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Fibronectin

Description

Fibronectin is a large glycoprotein widely distributed in soluble form in the plasma and body fluids. Many cell types synthesize fibronectin. There is also an insoluble form of fibronectin in tissues. Plasma fibronectin is not identical to cellular fibronectin but equally effective in supporting cell attachment. Fibronectin promotes the attachment and spreading of many adherent cells on plastic, but also mediates binding to other extracellular matrix components such as e.g. collagen.

Preparation

Fibronectin is purified from human plasma; donors tested negative for anti-HIV antibodies and HBs antigen.

Reconstitution

Dissolve contents in sterile water. Prepare a 1 mg/ml solution by gently warming the vial to 37° C; do not agitate, this may cause precipitation.

Recommended amount for coating

For coating of cell culture vessels 1-5 µg/cm2 is used.

Storag

Lyophylisate can be stored at 2-8° C; solution stored at -20° C in aliquots.

Fibronectin	5 mg	2705005
Fibronectin solution	1 mg/ml	2705001S



Attachment Factors

Bovine Serum Albumin

Description

Albumins serve as additive proteins for tissue cultures. They are the main protein component in serum and are added to cell culture media to increase the stability of cell membranes and to bind possibly toxic elements.

Advantages

- High purity > 99 %
- Minimal lot to lot variation
- Stringent control of raw materials
- US origin
- Full documentation
- Special products for unique applications

Storage

Powder and preserved solutions at 2 - 8 °C. Preservative free solution at -20 °C

Daving Common Albumain (DCA) Frantian V(I)	50 g	P06-1391050 P06-1391100
Bovine Serum Albumin (BSA) Fraction V ⁽¹⁾	100 g 500 g	P06-1391100 P06-1391500
Bovine Serum Albumin (BSA) Protease free ⁽¹⁾	100 g 500 g	P06-139210 P06-139250 NEW
Bovine Serum Albumin (BSA) Low Endotoxin ⁽¹⁾	10 g 50 g	P06-139310 P06-139350 NEW
Bovine Serum Albumin (BSA) Fatty acid free ⁽¹⁾	10 g 50 g	P06-139410 P06-139450 NEW
Bovine Serum Albumin (BSA) ⁽¹⁾ Microbiological grade	10 g 50 g	P06-0849010 P06-0849050
Bovine Serum Albumin (BSA) 30 % Solution, (1) Low salt	100 ml 500 ml	P06-138110 P06-138150 NEW
Bovine Serum Albumin (BSA) 30 % Solution, (1) High salt	100 ml 500 ml	P06-138210 P06-138250 NEW
Bovine Serum Albumin (BSA) 30 % Solution, (1) High polymer	100 ml 500 ml	P06-138310 P06-138350 NEW
Bovine Serum Albumin (BSA) 30 % Solution, (1) Low salt, Preservative free, Protease free	100 ml 500 ml	P06-138410 P06-138450 NEW

Other sizes are available upon request.

Human Serum Albumin

Description

Human Serum Albumin is a high quality product suitable for different applications.

Storage

Human Serum Albumin can be stored at 2 - 8 °C.

Human Serum Albumin (HSA) ⁽³⁾	25 g 50 g	P06-26025 P06-26050
	50 g	1 00-20030

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



Separating Solutions

Pancoll

Description

In many cases the isolation of cells is the first step for gene expression studies or in diagnostic procedures. Besides biological separation techniques physical separation methods are most commonly used. These methods use physical differences such as size and electrical charge of the particles to be separated. For this purpose so-called separating solutions (= centrifugation media) are used.

These media have to comply with the following criteria:

- Form a density gradient over the desired range
- Desired pH value and desired osmolality easily adjustable
- The solutions should not be too viscous at high density
- Do not cause any functional or morphological changes in biological materials
- Do not penetrate biological membranes

Our Pancoll separating solutions contain a polysaccharide with a molecular weight of 400,000 daltons; this hydrophilic polymer allows for production of aqueous solutions for cell separation with a density of up to 1.2 g/ml. PAN-Biotech offers a variety of ready-to-use products with a density of 1.063 g/ml up to 1.091 g/ml for a very wide range of cell separation applications.

Storage: 2° C to ambient temperature

When properly stored, separating solutions are stable for at least 36 months. The storage period starts with the manufacturing date.

Pancoll human, density 1.077 g/ml ⁽¹⁾	100 ml 500 ml	P04-60100 P04-60500
Pancoll mouse, density 1.086 g/ml ⁽²⁾	100 ml 500 ml	P04-64100 P04-64500
Pancoll rat, density 1.091 g/ml ⁽²⁾	100 ml 500 ml	P04-65100 P04-65500
Pancoll animal, density 1.077 g/ml ⁽²⁾	100 ml 500 ml	P04-63100 P04-63500
Pancoll monocytes, density 1.068 g/ml ⁽³⁾	100 ml 500 ml	P04-68100 P04-68500
Pancoll platelets, density 1.063 g/ml ⁽³⁾	100 ml 500 ml	P04-67100 P04-67500

) usually on stock, (2) minimum order 10 l, (3) available upon request



Separating Solutions Pre-Filled

Description

Pancoll separating solutions from PAN-Biotech are made from a neutral, highly cross-linked, hydrophilic polymer of sucrose with an average molecular weight of 400,000 daltons. Pancoll is suited for separation of lymphocytes and other cell types.

The ready-to-use solutions are available in 500 ml bottles (see page 105) as well as in pre-filled ready-to-use tubes with a separating membrane.

Stability

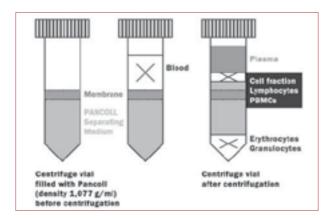
Pancoll is stable for 36 months at 2 °C to 20 °C. Protect from light!

Typical results with Pancoll

Lymphocytes	60 ± 20 % 95 ± 5 % > 90 %	yield of lymphocytes from original blood samples of the lymphocyte fraction are mononuclear leukocytes live cells (trypan blue-exclusion)
Other cells	3 ± 2 % 5 ± 2 % < 0,5 %	granolocytes erythrocytes total number of platelets of the original blood sample

Method of seperation

For lymphocyte separation blood is used which has been defibrinated or treated with anticoagulants (Heparin, EDTA, Citrate), and which is diluted with the same volume of a physiological saline solution. Then the Pancoll solution is carefully covered with a layer of diluted blood in a centrifuge vial, without mixing the phases. After a short centrifugation step (e.g. 800-1000x g for 20-30 minutes) at room temperature the lymphocytes, together with monocytes and platelets, can be harvested from the white blood cells layer between the plasma sample layer and the Pancoll. The separated cells are then washed twice in physiological saline solution to purify the lymphocytes by removing platelets. During centrifugation the cells of the blood sample migrate to the Pancoll layer where they get into contact with the polysaccharide contained in Pancoll. The red blood cells are aggregated by this substance at room temperature immediately. Aggregation causes an increase of the sedimentation rate of the red blood cells which aggregate together with the granulocytes as a sediment at the bottom of the centrifuge vial. Lymphocytes, monocytes and platelets are not so dense and can not enter and pass through the Pancoll layer. These cells are concentrated as white blood cell layer above the Pancoll layer and therefore can be harvested easily by careful pipetting. In subsequent centrifugation steps the lymphocytes are washed to remove remaining platelets, serum and Pancoll. As a result of this process a highly purified suspension of viable lymphocytes and monocytes (PBMC) is obtained.



Pancoll human, density 1.077 g/ml ⁽¹⁾	25 x 50 ml 50 x 10 ml	P04-60125 P04-60225
Pancoll animal, density 1.077 g/ml ⁽³⁾	50 x 10 ml	P04-63225

PAN SL-S Product Line

Endothelial cell biology has been greatly advanced by studying cultured vascular endothelial cells in vitro. Besides the understanding of many physiologic and pathologic processes, a multitude of basic cell signalling processes has been elucidated by using endothelial cells in culture. Traditionally, complete endothelial growth media contain animal serum. The advance of so-called low-serum media for endothelial cells has improved the quality of experimental data acquired in recent years. However, endothelial cells may synthesize substances which can not be detected due to their low quantity or masking effects from serum. In the past, cellular signalling pathways in endothelial cells have not been decipherable experimentally because even low concentrations of serum present in traditional media induce an undefined and undesired stimulation of cell surface receptors or intracellular signalling which only may become evident under serum-free conditions. As endothelial cells move into the field of interest for vascular tissue engineering with potential therapeutic application, the presence of whole animal serum is undesirable for such applications in the future.

All products described below are intended for use in a serum-free Endothelial Cell Culture System. Endothelial cells from different sources may be employed. For convenient use in this system, PAN-Biotech offers endothelial cells from human umbilical cord strictly isolated und cultured under animal serum-free conditions. This exclusive cell culture system is optimized for the maintenance and expansion of endothelial cells under serum-free conditions. Information about the composition, suitability, special advantages, and instructions are given for each individual product. For more information on SERUM-FREE Endothelial Cell Culture System from PAN-Biotech, please see accompanying data sheets.

Panexin SL-S is a genuine serum substitute which can fully replace FBS in otherwise completely supplemented endothelial cell culture media.

SL-S Trypsin/EDTA has been specifically designed for use in serum-free cell culture systems. A relatively low activity of trypsin results in gentle detachment of endothelial cells; a low phenol red concentration acts as a convenient indicator while providing mild conditions for the cells.

SL-S Medium is a working medium for rinsing of culture vessels after trypsin reaction to ensure a complete harvest of cells or for short time incubation of endothelial cells in growth factor free conditions. This medium is not suited for growth or long term culture of endothelial cells.

SL-S Trypsin Inhibitor is optimized to stop trypsin reaction and simultaneously providing ideal conditions for endothelial cells to recover from trypsin activity.

SL-S Collagen has been developed as a ready-touse solution for the coating of new culture vessels in endothelial subculture.

SL-S Cryopan is a serum-free medium for cryoconservation of endothelial cells resulting in high rates of recovery of viable cells after thawing.

Panexin SL-S Serum Substitute for HUVEC cultures ⁽³⁾	25 ml	P04-90065S
SL-S Trypsin/EDTA ⁽³⁾	50 ml	P10-0231SF
SL-S Trypsin-Inhibitor ⁽³⁾	50 ml	P10-0331SF
SL-S Medium (Working Medium)(3)	500 ml	P04-300500
SL-S Collagen 0.01 % ⁽³⁾	25 ml	P06-20650
SL-S Cryopan ⁽³⁾	25 ml	P07-94050

(1) usually on stock, (2) minimum order 10 l, (3) available upon request



(1) usually on stock, (2) minimum order 10 l, (3) available upon request



DMSO

Description

DMSO (Dimethylsulfoxide) is a colourless organic solvent which enters the cell and distributes inside the cell to prevent the formation of damaging ice crystals during the freezing procedure.

Dimethylsulfoxide (DMSO)	100 ml	P60-15840100
Dimethylsulfoxide (DMSO) for cell culture	100 ml	P60-36720100

Freezing Medium

Description

Our freezing medium is recommended for the cryoconservation of cells. The medium is based on DMEM, supplemented with a mix of fetal bovine serum and DMSO. This composition guarantees a high survival rate and excellent cell growth after thawing.

Freezing medium ⁽¹⁾	50 ml	P07-90050

Cryopan

Description

Cryopan I is a serum-free freezing medium for the cryoconservation of cells in a nitrogen storage tank or an ultra-low temperature freezer (< -150° C). It contains DMSO.

Freezing procedure

- Refrigerate freezing medium, culture medium and freezing tubes!
- Trypsinize cells, transfer the cells into the culture medium and centrifuge. Discard the supernatant and resuspend in culture medium
- Cell count should be adjusted to 1 5 x 10⁶/ml.
 The cells should be carefully resuspended to avoid clustering
- Spin down and resuspend the cells in an appropriate volume of cool freezing medium by pipetting only once or twice. Distribute 1 ml cell suspension per freezing tube
- To achieve a defined freezing rate of about 1° C per minute, manual freezing devices or computercontrolled freezing may be used

Ideal freezing rate: decrease of 1° C per minute

Carraman I(3)	10 ml	P07-92010
Cryopan 167	50 ml	P07-92050

1) usually on stock, (2) minimum order 10 l, (3) available upon request



Disinfectants

Barrycidal 36 spray bottle	50 ml	360050
Barrycidal 36 dispenser bottle	500 ml	360400
Barrycidal 36 spray bottle	11	361000
Barrycidal 36 can	51	365000
Barrycidal 36 can	101	360000
Barrycidal dispenser box	100 cloths	360101
Barrydin can	51	465000
Desipure can	101	660000
Spray head for 500 ml bottles		700500
Spray head for 1 l bottles		701000
Dosage pump for 500 ml bottles		710500
Wall-mounted dispenser for 500 ml bottles E24 (short handle) ELS24 (long handle)		720500 720501
Dosage pump for cans		730010
Tap for cans		740010



Disinfectants

Barrycidal* 36 is a modern disinfectant which fulfills the newest technical standards. It has a broad spectrum of efficacy and can be used in many aspects of daily life. With the new composition all areas of disinfection and hygiene are covered.

Barrycidal

Characteristics

Ready-to-use solution

Barrycidal® 36 disinfectant is a ready-to-use solution, suitable for prophylaxis of hospital-acquired infections in all areas of hospitals as well as for disinfection measures in food industry, diairies, soft drink industry, etc. Barrycidal[®] 36 is composed of a synergistic mixture of selected organic nitrogen compounds. It is effective against the whole spectrum of bacteria, yeasts, fungi, and viruses. Barrycidal® 36 is free of aldehydes, phenolic derivatives, chlorine and peroxides.

Special advantages of the ready-to-use disinfectant solution Barrycidal:

- Alcohol-free
- Non allergenic
- Poisonous category free
- Odourless
- High degree of biological degradability
- Good compatibility for all materials
- No irritation of skin or mucous membranes
- Rapid onset of action with long lasting effect
- Odour binding
- Stainless cleaning
- Large spectrum of efficacy: bacteria, yeast, fungi, viruses (e.g. Hepatitis B, HIV, Rota virus
- Free of mercury, aldehydes, phenol derivates, chlorine or peroxides

Indications and area of application

Cleaning and disinfection of all kinds of surfaces and objects in one step, especially in areas sensitive for smell, as well as for hand disinfection.

- Medical practice and hospitals
- Public baths
- Fire departements
- Health care
- Asylum
- Cosmetics, Kindergarten
- Laboratory, incubators, centrifuges
- Food industry
- Public facilities
- Podiatry, police
- Ambulance, solarium, sauna, hospital, schools
- Animal shelter
- Veterinarian
- Foot disinfection
- Athlete's foot prophylaxis
- Shoe desinfection

Surface disinfection (Hospitalismus prophylaxis and in general practice, bactericidal, fungicidal):

- undiluted/60 min
- HBV/HIV: undiluted/30 min
- Athlete's foot prophylaxis undiluted/15 min

Application

Surface disinfection

- Dosage: undiluted/60 min
- Spray disinfection: The surfaces to be disinfected should be completely wet by spraying. Let dry, no rinsing necessary, unless for surfaces, which come into contact with food

Composition

100 g contain:

0.0975 g n-Octyl-dimethyl-benzylammonium chloride 0.0300 g Benzethonium chloride 0.0025 g Methyl-benzethonium chloride with some more cleaning and disinfectant substances like propandiol, triethanolamine, etc.

Physico-chemical Properties

Appearance	clear, colourless solution
pH-Value (20° C)	8.0 ± 0.5
Density (20° C)	1.046 ± 0.020
Stability	5 years

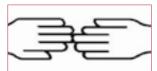
Available sizes

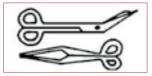
50 ml spray bottle 500 ml round bottle 1000 ml spray bottle 5 liter can

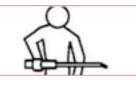
10 liter can

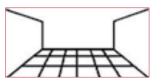
Classified as non-poisonous and non-toxic solution

BAG E1227/T73512 (Swiss) DGHM-listed (Germany) OEGHMP-listed (Austria)











Disinfectants

Barrydin

Concentrate for instrument disinfection free of aldehydes and phenols

Characteristics

Barrydin is a disinfectant based on a synergistic mixture of quaternary ammonium compounds, guanidinium derivatives and alcylpolyamines. It is also free of aldehydes, particulary formaldehyde, phenol derivatives, chlorine, alcohol and similar. It is characterized by a broad spectrum of activity, short contact time, excellent cleaning properties, neutral smell and compatibility of materials. In addition, protein fixation is avoided due to aldehyde-free formulation.

Range of application

Short term disinfection and cleaning in one step. Cleaning and disinfection of all kinds of surgical instruments in all clinical departments and medical practices incl. instruments for micro-invasive surgery (MIS), anesthesia material and flexible and rigid endoscopes.

Composition

100 g Barrydin contain:

3.75 g Cocos-propylen-diamin guanidine di-acetate 5.63 g Didecyl-oxyethyl-methyl-ammonium propionate

Physico-chemical properties

Appearance:	clear, blue-green solution
pH-Value (20° C):	concentrate: 9.7 2 % aqueous solution 10.4
Conductivity of the concentrate:	10 mS x cm ⁻¹
Density (20° C):	0.995

Microbiology

- Bactericidal (incl. TbB, mycobacterium terrae) and Fungicidal
- Virus inactivating (HBV, HIV, Adenovirus, Papovavirus, Poliovirus)
- Sporocidal (qualitative suspension testing)

Dosage

Instrument disinfection (incl. M. tuberculosis):

1.0 % for 60 min 2.0 % for 30 min

Short time disinfection:

3% for 15 min

HBV/HIV: 1% 60 min – 2% 15 min Adeno virus 2% 60 min – 4% 30 min Papova virus 1% 60 min - 2% 30 min

Polio virus 50° C: 1% 10 min

Application

Prepare a working dilution in the right concentration!

Instrument disinfection

Immediately after use place open instruments in the working solution. All areas to be cleaned have to be fully submerged in disinfectant solution. Close container if possible. Replace solution for heavy contamination. Wash instruments carefully with running water after appropriate contact time and allow drying. Suitable for cycling techniques and ultrasonic devices.

Registrations and listings

SFOHP = Swiss Federal Office of Health Public Bern E1230/T91445 Poison-classification: No. 4

Disinfectant listed by DGHM (Germany certificate)

Available size

5 litre PE-can

Advantages

- Cleaning in one step
- Very economical
- Short term disinfection Aldehyde and phenol free
- Extremely low concentration
- Very broad spectrum of activity
- Good compatibility for all instruments





Disinfectants

Desipure

Disinfectant cleaning concentrate

Characteristics

Desipure C-100 is a disinfectant concentrate with intensive cleaning properties for all purposes of surface disinfection and cleaning in all areas of hospitals for hospital-acquired infections as well as for disinfection measures in food industry, kitchen, household and so on. Desipure C-100 contains as active ingredient a surface active organic nitrogen compound. It is effective on the whole spectrum of bacteria, incl. Salmonella, yeasts and fungi.

Desipure C-100 is free of aldehydes, particularly formaldehyde, phenolic derivatives, chlorine and peroxides.

Indications

Cleaning and disinfection of all kinds of surfaces and objects in one step, particularly in areas sensitive for unpleasant odour.

Composition

100 g Desipure C-100 contain

- 9.8 g N,N-Didecyl-N-methyl-poly (oxyethy) ammoniumpropionate
- 12.0 g ethoxylated fatty alcohols glycol derivatives

Physico-chemical properties

Appearance	clear, yellowish solution
pH-Value (20° C):	7.0 ± 1.0
Density (20° C):	0.995
Stability:	5 years

Microbiology

Bactericidal

Fungicidal

Virus inactivating (HBV, HIV, Rotaviruses)

Dosage

Surface disinfection for hospitalism prophylaxis:

1.0 % for 1 h

0.5 % for 4 h

Application - Surface disinfection

- Wiping disinfection (two-bucket-method) or similar
- Out of dosage apparatuses
- Cleaning machines
- Spraying with suitable equipment

Registrations and Listings

BAG/SFOHP E1229 / T78288 Poison-classification no.4 Disinfectant listed of DGHM (Germany)

Available sizes

10 litre cans

Dosage pumps for 10 litre-cans

Discharge taps for 10 litre cans

Advantages

- Neutral odour
- Virus inactivating
- Good biological degradability
- Broad action spectrum
- Highly economical
- Rapid surface disinfection
- Aldehyde and formaldehyde free
- Cleaning in one process step





Biologicals

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Chemokines	120

www.pan-biotech.com

Introduction

Cytokines, growth factors, and chemokines from PAN-Biotech

Cytokines are becoming increasingly important in cell biology. Sugar-containing proteins have regulating functions for the growth and differentiation of body cells. Many cytokines also play an important role in immunological reactions, and they are then generally referred to as mediators.

Cytokines, which primarily trigger and/or regulate the proliferation and differentiation of target cells, are known as growth factors. These proteins that are transferred as signals from one cell to another, and therefore relay information, play a role mainly in the development of multi-cellular organisms. Growth factors are either secreted i.e. released by the cells into the environment or they are membrane-bound. They function when recognised by a receptor on the surface of the target cell. Only cells that carry the specific receptor for the respective growth factor (= ligand) can respond to the signal.

As soon as the factor binds to its ligand, a change in conformation results in the generation of an intracellular signal. By further signal transfer, this will cause genes to be activated or inactivated.

PAN-Biotech offers a high-quality range of cytokines, growth factors as well as chemokines, chemotactical cytokines ("chemoattractant cytokines"), which can be secreted by many cell types e.g. from phagocytes and dendritic cells but also from tissue cells.

Chemokines can attract and activate leukocytes. They therefore play an important role as mediators in regulating targeted leukocyte migration and the inflammation processes triggered as a result.

Please ask for any chemokine or growth factor which is not listed. We may be able to provide.





Human Cytokines and Growth Factors

Short Term	Description	Cat.no.	Size
Acrp30 HEK	Adiponectin Globular rec.	CB-2800007	2 μg
		CB-2800008	10 μg
Acrp30	Adiponectin rec.	CB-2800001	5 μg
		CB-2800002	25 μg
Acrp30 Tri	Adiponectin Trimeric Form rec.	CB-2800010	2 μg
DMD 4	P. M. I. di di di	CB-2800011	10 μg
BMP-4	Bone Morphogenetic protein-4 rec.	P-3610002	2 μg
BMP-6	Bone Morphogenetic protein-6 rec.	CB-1113008	2 μg
BMP-7	Bone Morphogenetic protein-7 rec.	CB-1113011 CB-1113012	2 μg 10 μg
BDNF	Brain-Derived Neurotrophic Factor rec.	CB-1115000	2 μg
		CB-1115001	10 μg
		CB-1115002	1 mg
CT-1	Cardiotrophin-1 rec.	CB-1115006	2 μg
CNTF	Ciliary Neurotrophic Factor rec.	CB-1515001	20 μg
EGF	Epidermal Growth Factor rec.	CB-1101001	100 μg
		CB-1101002	500 μg
		CB-1101003	1 mg
EPO-α	Erythropoietin-alpha rec.	CB-2015001	50 μg
FGF-1	Fibroblast Growth Factor-acidic rec.	CB-1102010	10 μg
		CB-1102011	50 μg
FGF-2	Fibroblast Growth Factor-basic rec.	CB-1102024	10 μg
		CB-1102021 CB-1102023	50 μg
Flt3	Flt3-Ligand rec.	CB-1102023	1 mg
1113	Pito-Ligand rec.	CB-1119000 CB-1119001	2 μg 10 μg
GDNF	Glial-Drived Neurotrophic Factor rec.	CB-1116001	10 μg
GMCSF/IL3	gm-csf/IL-3 Fusion Protein (PIXY321) rec.	CB-2110005	10 μg
GMCSF	Granulocyte Macrophage-Colony	CB-2110000	
GMCSI	Stimulating Factor rec.	CB-2110000 CB-2110002	2 μg 10 μg
	Community Fuctor Foot	CB-2110003	1 mg
GCSF	Granulocyte-Colony Stimulating Factor rec.	CB-2110100	2 μg
	, , ,	CB-2110101	10 μg
HGF Sf9	Hepatocyte Growth Factor Sf9 rec.	CB-1108002	2 μg
		CB-1108010	10 μg
		CB-1108100	1 mg
HGF CHO	Hepatocyte Growth Factor, CHO, rec.	CB-1108003	2 μg
		CB-1108011	10 μg
IGF-1	Insulin Like Growth Factor-1 rec.	CB-1104112	20 μg
ICD 2	T. It till C. It P. I. 5	CB-1104113	100 μg
IGF-2	Insulin Like Growth Factor-2 rec.	CB-1104201	10 μg
To out is:	In outling to a	CB-1104202	50 μg
Insulin	Insulin rec.	P-2701002 P-2701001	25 mg 250 mg
		1-2/01001	250 mg

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Human Cytokines and Growth Factors

Short Term	Description	Cat.no.	Size
IFN-α 1	Interferon-alpha 1 rec.	CB-2120100 CB-2120101	2 μg 10 μg
IFN-α 2a	Interferon-alpha 2a rec.	CB-2120110 CB-2120112	20 μg 100 μg
IFN-β 1a	Interferon-beta 1a rec.	P-2360011 P-2360012 P-2360013	2 μg 10 μg 1 mg
IFN-β 1b	Interferon-beta 1b rec.	CB-2120121	10 μg
IFN-γ	Interferon-gamma rec.	P-2060020 P-2060100 P-2061000	20 μg 100 μg 1 mg
IL-1β	Interleukin-1 beta rec.	CB-2130120 CB-2130121	2 μg 10 μg
IL-1β His	Interleukin-1 beta, His Tag rec.	CB-2130123	5 μg
IL-10	Interleukin-10 rec.	CB-2131000 CB-2131001	2 μg 10 μg
IL-12	Interleukin-12 rec.	CB-2131201	10 μg
IL-15	Interleukin-15 rec.	CB-2131500 CB-2131501	2 μg 10 μg
IL-2	Interleukin-2 rec.	CB-2130203 CB-2130202	10 μg 50 μg
IL-3	Interleukin-3 rec.	CB-2130300 CB-2130301	2 μg 10 μg
IL-4	Interleukin-4 rec.	CB-2130405 CB-2130407	2 μg 10 μg
IL-5	Interleukin-5 rec.	CB-2130501	10 μg
IL-6	Interleukin-6 rec.	CB-2130600 CB-2130603	5 μg 20 μg
KGF	Keratinocye Growth Factor rec.	CB-1105001	10 μg
Leptin	Leptin rec.	CB-1300058	200 μg
LIF	Lif rec.	CB-1106001	10 μg
β-NGF	Nerve Growth Factor beta 2 rec.	CB-1117001M CB-1117001	5 μg 20 μg
NRG1	Neuregulin-1/Heregulin-b2 rec.	CB-4070010	10 μg
NT-3	Neurotrophin-3 rec.	CB-1125032	10 μg





Human Cytokines and Growth Factors

Short Term	Description	Cat.no.	Size
PDGF-AA	Platelet Derived Growth Factor-AA rec.	CB-3410010	10 μg
		CB-3410011	1 mg
PDGF-AB	Platelet Derived Growth Factor-AB rec.	CB-1109301	10 μg
PDGF-BB	Platelet Derived Growth Factor-BB rec.	CB-1109200	2 μg
		CB-1109201	10 μg
Resistin	Resistin rec.	CB-1300118	5 μg
SCF	Stem Cell Factor rec.	CB-1110000	2 μg
		CB-1110001	10 μg
		CB-1110002	1 mg
TPO	Thrombopoietin rec.	CB-1127000	2 μg
TRAIL	TNF-Related Apoptosis Inducing Ligand/Apo2L rec.	CB-1127100	10 μg
TGF-β 1	Transforming Growth Factor-Beta 1 rec.	CB-1111131	1 μg
		CB-1111122	5 μg
		CB-1111123	100 µg
TGF-β 3	Transforming Growth Factor-Beta 3 rec.	CB-1111151	2 μg
		CB-1111153	10 μg
rHuTNFR	Tumor Necrosis Factor Receptor Fusion Protein rec.	CB-1111162	1 mg
TNF-α	Tumor Necrosis Factor-alpha rec.	CB-1112011	10 μg
		CB-1112012	50 μg
VEGF (121)	Vascular Endothelial Growth Factor (121) rec.	CB-1114002	10 μg
VEGF	Vascular Endothelial Growth Factor rec.	CB-1114100	2 μg
		CB-1114102	10 μg
VEGF-C	Vascular Endothelial Growth Factor Related Protein rec.	CB-1114011	10 μg
VEGF CHO	Vascular Endothelial Growth Factor, CHO rec.	CB-1114013	2 μg

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Other Species Cytokines and Growth Factors

Short Term	Description	Cat.no.	Size
mβ-NGF	Murine beta Nerve Growth Factor rec.	CB-1117007 CB-1117008	5 μg 20 μg
mEGF	Murine Epidermal Growth Factor rec.	CB-1214120 CB-1214121	100 μg 500 μg
mFGF-2	Murine Fibroblast Growth Factor-basic rec.	P-3860001 P-3860002	10 μg 50 μg
mFlt3	Murine Flt3-Ligand rec.	CB-2250001	10 μg
mGMCSF	Murine Granulocyte Macrophage-Colony Stimulating Factor rec.	CB-2210000 CB-2210001 CB-2210002	2 μg 10 μg 1 mg
mGCSF	Murine Granulocyte-Colony Stimulating Factor rec.	CB-1200000	2 μg
mIFN-γ	Murine Interferon-gamma rec.	CB-2230030 CB-2230031	20 μg 100 μg
mIL-1α	Murine Interleukin-1 alpha rec.	CB-2230111	10 μg
mIL-1β	Murine Interleukin-1 beta rec.	CB-2230120 CB-2230121	2 μg 10 μg
mIL-12	Murine Interleukin-12 rec.	CB-2231202	0.1 mg
mIL-2	Murine Interleukin-2 rec.	CB-2230220 CB-2230221	5 μg 20 μg
mIL-3	Murine Interleukin3- rec.	CB-2230300 CB-2230301 CB-2230302	2 μg 10 μg 1 mg
mIL-4	Murine Interleukin-4 rec.	CB-2230403	2 μg
mIL-6	Murine Interleukin-6 rec.	CB-2230600 CB-2230601 CB-2230602	2 μg 10 μg 1 mg
mMCSF	Murine Macrophage Colony Stimulating Factor rec.	P-4390002 P-4390010	2 μg 10 μg
mSCF	Murine Stem Cell Factor rec.	CB-1210000	2 μg
mTNF-α	Murine Tumor Necrosis Factor-alpha rec.	CB-1212011M CB-1212011	5 μg 20 μg
mVEGF	Murine Vascular Endothelial Growth Factor rec.	CB-1214000 CB-1214001	2 μg 10μg
bECGS	ECGS (from bovine hypothalamus)	CB-11000050	50 mg
oPrl	Ovine Prolactin rec.	CB-2310015 CB-2310016	10 μg 50 μg
pGMCSF	Procine Granulocyte Macrophage-Colony Stiumlating Factor rec.	CB-2330000	2 μg
pIL-10	Porcine Interleukine-10 rec.	CB-2331001	10 μg
rIFN-γ	Rat Interferon-gamma rec.	CB-2420031 CB-2420032	20 μg 100 μg
rIL-1a	Rat Interleukin-1 alpha rec.	CB-2430122	1 mg
rTNF-α	Rat Tumor Necrosis Factor-alpha rec.	CB-1412011	20 μg





Chemokines

Short Term	Description	Cat.no.	Size
I-309	Human I-309 (CCL1) rec.	CB-1232701	10 μg
I-TAC	Human I-TAC (CXCL11) rec.	P-3340002	20 μg
Lymphotactin	Human Lymphotactin (XCL1) rec.	CB-1122001	10 μg
MCP-2	Human Monocyte Chemotactic Protein-2 (CCL8) rec.	CB-1407003	2 μg
SDF-1α	Human Stromal Cell-Derived Factor-1 alpha (CXCL12) rec.	CB-1118005	10 μg
mMCP-1	Murine Monocyte Chemotactic Protein-1 (CCL2) rec.	CB-2232000 CB-2232002	2 μg 1 mg
mSDF-1α	Murine Stromal Cell-Derived Factor-1 alpha (CXCL12) rec.	CB-1118008	10 μg





Molecular Biology

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Polymerases

Taq DNA Polymerase

The Taq DNA Polymerase is a thermo-stable DNA polymerase complex exactly following the original procedure for the isolation of DNA polymerases.

Storage and dilution buffer

20~mM Tris-HCI (pH 8.0), 100~mM KCL, 0.1~mM EDTA, 1~mM DTT, 50% glycerol, 0.5% Nonidet P40 and 0.5% Tween 20.

Unit definition

One unit is defined as the amount of enzyme that incorporates 10 nmoles of dNTP's into acid-insoluble fraction in 30 minutes at 72° C under the standard assay conditions: 25 mM TAPS (tris-(hydrooxymethyl)-methyl-amino-propansulfonic acid, sodium salt) pH 9,3 (at 25° C), 50 mM KCl, 2 mM 50 mM MgCl₂, 1 mM beta-mercaptoethanol, 200 μ M each dATP, dGTP, dTTP, 100 μ M dCTP (a mix of cold and P32-labelled), 12,5 μ g activated salmon sperm DNA, in a final volume of 50 μ l.

Supplied buffers (alternatively with complete or incomplete buffer)

- 10x PCR buffer with MgCl₂: 100 mM Tris-HCI (pH 9.0 at 25° C), 500 mM KCl 15 mM MgCl₂, 1.0% Triton X-100
- 10x PCR buffer without MgCl₂:
 100 mM Tris-HCI (pH 9.0 at 25° C), 500 mM KCl
 1.0% Triton X-100
- Magnesium stock solution:
 25 mM MgCl₂

Stabilit

The enzyme is stable for more than 12 months if stored at -20° C. The enzyme is also stable for some days at temperatures above 20° C.

Associated activities

Endonuclease and exonuclease activities were not detectable after 4 hours incubation of 1 μg native lambda DNA and 0.22 μg of EroR I-digested lambda DNA at 72° C in the presence of 15 - 20 units of Taq DNA Polymerases.

Properties and application

The Taq DNA Polymerase is a thermostable DNA polymerase from T. aquaticus of high purity with good fidelity and high processivity.

Taq DNA Polymerase with buffer and MgCI2	250 units	MB-30010250
Taq DNA Polymerase with buffer and MgCI ₂ with Phenol red	250 units	MB-30020250





Polymerases

PANScript DNA Polymerase

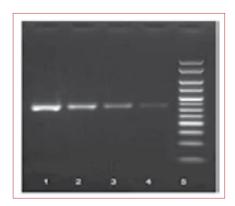
Features and applications

- Consistent results
- Premium Taq polymerase suited for a wide range of applications
- Processes fragments of up to 5Kb
- Leaves "A" overhang
- Available as ready-to-use 2x reaction mixes (PAN Mix and PAN Mix red)
- Routine PCR applications
- Products suitable for TA cloning

PANScript is widely used by molecular biologists that have come to depend upon the robust performance of this reagent.

PANScript is a highly purified thermostable DNA polymerase offering very high yield over a wide range of PCR templates, and is the ideal choice for most assays. PanScript is a robust preparation and consistently delivers high yields with minimal background. PANScript possesses 5' to 3' exonuclease activity and leaves an "A" overhang such that the PCR product is suitable for effective integration into TA cloning vectors.

PANScript is supplied with 10x NH₄-based reaction buffer, which provides optimal conditions for most experiments. Additional MgCl₂ is provided to allow reaction conditions to be adjusted to suit the template. The specificity and performance of PANScript can be further improved with the use of 2x PAN Mate Additive (Cat No. PAN737041), which is designed for GC- or ATrich DNA, dirty templates or sequences with a high level of secondary structure.



High performance with PANScript A175 bp fragment was amplified from pG EM 3z f(+) using PANScrip DNA-Polymerase.Lane 1-4: 10-fold serial dilution of template. (starting concentration 25 ng/ μ l) Lane 5: PANLadder V

PANScript DNA Polymerase is purified from Thermus aquaticus.

PCR Reaction Conditions (for a 50 µl volume)

10x NH₄ Buffer 5 μl 50 mM MgCl₂ Solution 1.5 - 4.0 μl 100 mM dNTP Mix (see below) 0.5 - 1.0 μl Template and Primers as required PANScript 0.5 - 1.0 μl Water (ddH₂O) up to 50 μl 100 mM dNTP Mix is available as a separate product (Cat No: PAN73028)

Denature: 94 – 96° C

Elongate: 70 – 72° C (allowing 15 - 30 seconds/Kb)

This data is intended for use as a guide only; conditions will vary from reaction to reaction and may need optimization.

Reagent specifications

10x NH₄ Reaction Buffer:

 $160 \text{ mM} \text{ (NH}_4)2\text{SO}_4$, 670mM Tris-HCl (pH 8.8 at 25° C), 0.1% stabilizer

MgCl₂ Stock Solution:

50 mM MgCl₂ (suggested final concentration 1.5 mM - 4 mM).

Storage buffer

 $20~\mathrm{mM}$ Tris-HCl, pH 7.5, $100~\mathrm{mM}$ NaCl, 0.1 mM EDTA, $2~\mathrm{mM}$ DTT, 50% Glycerol and stabilizers.

Storage conditions

PANScript can be stored for 12 months at -20° C.

Associated activities

Endonuclease and exonuclease activities were not detectable after 2 and 1 hour incubations, respectively, of 1 μ g lambda DNA and 0.22 μ g of EcoR I-digested lambda DNA at 72° C in the presence of 15 - 20 units of PANScript DNA polymerase.

Unit definition

One unit is defined as the amount of enzyme that incorporates 10 nmoles of dNTPs into acid-insoluble form in 30 minutes at 72° C.

PANScript DNA Polymerase 500 units MB-1100500



Polymerases

PANScript red DNA Polymerase

Features and applications

- Easy visual recognition
- Direct loading onto agarose gels
- Same high performance as PANScript DNA Polymerase
- Leaves "A" overhang
- Available as a ready-to-use 2x reaction mix (PAN Mix Red)
- Routine PCR assays
- Products suitable for TA cloning
- High throughput applications

PANScript red DNA Polymerase is a formulation of our regular PANScript DNA Polymerase, which contains a non-toxic and non-hazardous red dye. The red dye provides an easy and quick identification of reactions to which the enzyme has been added, and facilitates the confirmation of complete mixing. When the reaction is complete, a sample of the reaction mix can be loaded directly onto the agarose gel without the need for loading buffer, since the mix is of sufficiently high density to sink to the bottom of the gel. The red dye migrates towards the positive electrode, thereby providing a means to monitor the progress of the electrophoresis.

The presence of the dye has no effect on routine enzymatic manipulations, although rare exceptions may occur. In order to produce a reaction of sufficient density to allow for the direct loading of a sample onto a gel, we recommend using a minimum of 1.5 Units per 50 μ l reaction.

The specificity and performance of PANScript red can be further improved with the use of 2x PAN Mate Additive (Cat No. PAN737041), which is designed for GC or ATrich DNA, dirty templates or sequences with a high level of secondary structure.

PANScript DNA Polymerase is purified from Thermus aquaticus.

PCR Reaction Conditions (for a 50 µl volume)

10x NH₄ Buffer	5 μl
50 mM MgCl2 Solution	1.5 - 4.0 μl
100 mM dNTP Mix (see below)	0.5 - 1.0 μl
Template and Primers	as required
PANScript red	1.5 - 2.5 μl
Water (ddH2O)	up to 50 μl
100 mM dNTP Mix is available as a separate product (Cat No:	
PAN73028)	

Denature: 94° – 96° C

Elongate: 70° – 72° C (allowing 15 - 30 seconds/Kb)

This data is intended for use as a guide only; conditions will vary from reaction to reaction and may need optimization.

Reagent specifications

10x NH₄ Reaction Buffer: 160 mM (NH₄)₂SO₄, 670 mM Tris-HCl (pH 8.8 at 25° C), 0.1% stabilizer

MgCl2 Stock Solution: 50 mM MgCl2 (suggested final concentration 1.5 mM - 4 mM).

Storage buffer

20~mM Tris-HCl, pH 7.5, 100~mM NaCl, 0.1 mM EDTA, 2~mM DTT, 50% Glycerol and stabilizers and inert dye. Storage Conditions

PANScript red can be stored for 12 months at -20° C.

Associated activities

Endonuclease and exonuclease activities were not detectable after 2 and 1 hour incubations, respectively, of 1 μg lambda DNA and 0.22 μg of EcoR I-digested lambda DNA at 72° in the presence of 15 - 20 units of PANScript red DNA polymerase.

Unit definition

One unit is defined as the amount of enzyme that incorporates 10 nmoles of dNTPs into acid-insoluble form in 30 minutes at 72° C.

PANScript red DNA Polymerase	500 units	MB-1100600
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- Outstanding and robust performance
- For PCR assays requiring hot-start
- Excellent yield in quantitative assays
- Convenient set up at room temperature
- Leaves "A" overhang
- Available as ready-to-go versions PAN Hot Mix and PAN Hot Mix red
- Highly suited to real-time assays
- Products suitable for TA cloning

PAN Hot Start is a heat-activated thermostable DNA polymerase isolated from a novel organism. PAN Hot Start provides improved specificity as compared to standard polymerases and can eliminate the presence of non-specifics, such as primer-dimers and mis-primed products. PAN Hot Start is inactive at room temperature and therefore, prior to PCR cycling, requires activation by heat treatment for 10 minutes. Subsequently, the reaction can be handled according to the user's existing protocols for thermostable DNA polymerases.

Specificity and performance of PAN Hot Start can be further improved with the use of 2x PAN Mate Additive, which is designed for GC- or AT-rich DNA, "dirty" templates or sequences with a high level of secondary structure.

PCR Reaction Conditions (for a 50 µl volume)

10x PAN Hot Start Buffer 50 mM MgCl₂ 1.5 - 4.0 µl 100 mM dNTP Mix (see below) 0.5 - 1.0 µl Template and Primers as required PAN Hot Start 0.2 - 1.0 µl Water (ddH₂O) up to 50 µl 100 mM dNTP Mix is available as a separate product (Cat No: PAN73028) Activate: pre-heating step at 95° C for 10 minutes Denature: 94° – 96° C Extension: 72° C (allowing 15 - 30 seconds/Kb) This data is intended for use as a guide only; conditions will vary from reaction to reaction and may need optimization.

Reagent specifications

10x PAN Hot Start Buffer:160mM (NH₄)₂SO₄, 1M Tris-HCl pH 8.3 and enhancers

Storage Conditions

PAN Hot Start DNA Polymerase can be stored for 12 months at -20° C.

Storage and Dilution Buffer

20 mM Tris-HCl, pH 7.5, 100 mM NaCl, 0.1mM EDTA, 2 mM DTT, 50% Glycerol, and stabilizers.

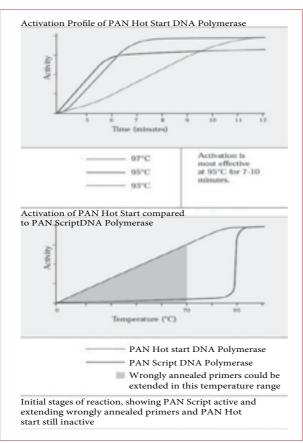
Associated activities

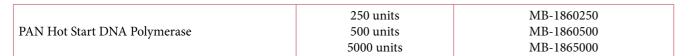
Endonuclease and exonuclease activities were not detectable after 4 hours of incubation of 1 µg of pBR322 plasmid DNA and 0.5 µg Hind III-digested lambda phage DNA at 72° C in the presence of 20 u of PAN Hot Start.

Unit definition

One unit is defined as the amount of enzyme that

10nmoles of dNTPs into acid-insoluble form in 30 minutes at 72° C.





Polymerases

PowerScript DNA Polymerase short range

Features and applications

- Ideal for problematic templates that fail with standard Tag DNA Polymerases
- Ideal for fragments up to 5 Kb in length
- Higher fidelity than Taq
- For high fidelity PCR
- Suitable for both TA and blunt-end cloning

Powerscript short DNA Polymerase is a high-performance proprietary complex of enzymes specifically designed for difficult/problematic PCR applications requiring high processivity with fidelity that would normally fail with standard Taq Polymerases.

PowerScript short DNA Polymerase is recommended for short genomic DNA fragments of up to 3 Kb, or up to 5 Kb on Lambda DNA.

Components	250 Units	500 Units
PowerScript short range	62.5 μl	125 μl
10x OptiBuffer	1.2 ml	2 x 1.2 ml
50 mM MgCI ₂ Solution	1.2 ml	1.2 ml
5x Hi-Spec Additive	1.2 ml	1.2 ml

Reagent specifications

5x Hi-Spec Additive is a specificity enhancer. If necessary, re-dissolve Hi-Spec by heating to 70° C and vortexing.

Storage buffer

20 mM Tris-HCl, pH 7.5, 100 mM NaCl, 0.1mM EDTA, 2 mM DTT, 50% Glycerol, and stabilizers.

Storage conditions

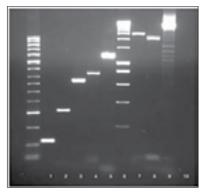
PowerScript short DNA Polymerase can be stored for 12 months at -20° C.

Associated activities

Endonuclease and exonuclease activities were not detectable after 4 hours of incubation of 1 µg of pBR322 plasmid DNA and $0.5\,\mu g$ Hind III-digested Lambda DNA at 72° C in the presence of 20 units of PowerScript short.

Unit definition

One unit is defined as the amount that incorporates 10nmoles of dNTPs into acid-precipitable form in 30 minutes at 72° C.



High specificity with problematic templates using PowerScript short.

A range of fragments from human genes were amplified, varying in length and GC content.

Lane 1: PANLadder II

119 bp and 43% GC product amplified from the human glucocerebrosidase gene

321 bp and 37% GC product amplified from Angiotensin receptor II gene 626 bp and 56% GC product amplified

from the Rhodopsin gene

762 bp and 33% GC product amplified

from the ß-Globin gene 1200 bp and 54% GC product amplified Lane 6

from the alpha-1-antitrypsin gene Lane 7: PANLadder I

2256 bp and 52% GC product amplified

from the p53 gene

2000 bp and 32% GC product amplified from the Angiotensin receptor II gene

Lane 10: 6000 bp and 51% GC product amplified from the alpha-1-antitrypsin gene

Dayyon Carint DNA Dalymanasa shart ranga	250 units	PAN721064
PowerScript DNA-Polymerase short range	500 units	PAN721065



Features and applications

- Ideal for problematic templates that fail with standard Taq DNA Polymerases
- Ideal for fragments 2 20 Kb in length
- Higher fidelity than Taq
- Available as a ready-to-use 2x reaction mix
- For high fidelity PCR
- Suitable for both TA and blunt-end cloning

PowerScript DNA Polymerase long range is a high-performance proprietary complex of enzymes specifically designed for difficult/ problematic PCR applications requiring high processivity with fidelity that would normally fail with standard Taq polymerases.

PowerScript DNA Polymerase long range is recommended for long Genomic DNA fragments of between 2 - 20 Kb, or up to 30 Kb Lambda DNA fragments. With Lambda DNA as template, the best performance is achieved in the 2 - 20 Kb range. PowerScript long is our original widely used PowerScript formulation.

Components	250 Units	500 Units
PowerScript long range 10x OptiBuffer 50 mM MgCI ₂ Solution 5x Hi-Spec Additive	62.5 μl 1.2 ml 1.2 ml 1.2 ml	125 μl 2 x 1.2 ml 1.2 ml 1.2 ml
3x 111 Spec ridditive	1.2 1111	1.2 1111

Reagent specifications

5x Hi-Spec Additive is a specificity enhancer. If necessary, re-dissolve Hi-Spec by heating to 70° C and vortexing.

Storage buffer

20 mM Tris-HCl, pH 7.5, 100mM NaCl, 0.1mM EDTA, 2 mM DTT, 50% Glycerol, and stabilizers.

Storage conditions

PowerScript short DNA Polymerase can be stored for 12 months at -20° C.

Associated activities

Endonuclease and exonuclease activities were not detectable after 4 hours of incubation of 1 μg of pBR322 plasmid DNA and 0.5 μg Hind III-digested Lambda DNA at 72° C in the Presence of 20 units of PowerScript long.

Unit definition

One unit is defined as the amount that incorporates 10nmoles of dNTPs into acid-precipitable form in 30 minutes at 72° C.



Long range PCR with PowerScript long. PowerScript long is a polymerase ideally suited to the amplification of long DNA fragments. A 20 Kb fragment of Lambda DNA was amplified using PowerScript long DNA Polymerase.

Lane 1: PANLadder I (top band = 10 Kb)

Lane 2: Amplification of 20 Kb Lambda DNA fragment

PowerScript DNA Polymerase long range	250 units	MB-1120250
PowerScript DNA Polymerase long range	500 units	MB-1120500

Molecular Biology Reagents

Proteinase K

Features and applications

- Broad-spectrum serine protease
- Active under denaturing conditions
- Stable at high temperatures
- Molecular biology grade
- Available as powder and stabilized stock solution
- Inactivation of RNases/DNases during nucleic acid extraction
- Protein modification
- General protein digestion
- Determination of enzyme localization

Proteinase K is an enzyme used to digest most proteins in molecular-biological techniques. The enzyme may be used at 56° C for up to 4 hours, or 37° C for overnight incubations. Proteinase K solution is stabilized with a specially formulated buffer, and can be used directly from the freezer.

Recommendations for use

- Dissolve to 20 mg/ml in 50 mM Tris-HCl, 2 mM calcium acetate, pH 8.0
- Proteinase K may be used at 56° C for up to 4 hours, or 37° C for overnight incubations
- Proteinase K has an optimal pH of 7.5 12.0
- To remove common contaminants from nucleic acid preparations use at a working concentration of 5 $\mu g/m^{1}$

Storage conditions

Proteinase K can be stored for 12 months at -20° C.

Contaminants

RNase Activity: No detectable ribonuclease activity detected with MS2RNA after 6 hour incubation at 37° C DNase Activity: No detectable nicking activity detected with pBR322 after 6 hour incubation at 37° C

Unit definition

One unit is defined as the amount of enzyme that will liberate 1.0 µmol of tyrosine per minute at 37° C, pH 7.5.

Proteinase K	100 mg	MB-4300002

PAN Ladder I

Features

- 14 bands from 200 bp 10 000 bp
- Accurate quantitation
- Easy identification and orientation
- Ready-to-use format

PANLadder™ I is a popular ready-to-use molecular weight marker, especially designed for easy DNA quantification and size determination. This ready-to-use format reduces handling steps and saves time; simply transfer HyperLadder I from the vial to the gel.

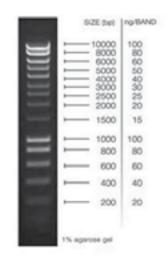
PANLadder™ I produces a pattern of 14 regularly spaced bands, ranging from 200 to 10,000 bp. To allow easy identification and orientation, the 1000 and 10,000bp bands have the highest intensity.

When the standard loading volume of 5 μ l per lane (720 ng of DNA) is being used, each band corresponds to a precise amount of DNA.

A 5x sample loading buffer is supplied for your convenience. Under no circumstances should it be used to dilute/ load ladder.

Storage conditions

PANLadder[™] I can be stored at -20° C until first use and thereafter at 2-8° C for up to 6 months. Avoid multiple freeze/thaw cycles.



DANE 11 T	200 lanes	PAN733025
PAN Ladder I	500 lanes	PAN733026





PAN DNA Clean

Features and applications

- Column-free PCR clean-up
- Post-PCR recovery of up to 98%
- Cost-effective, simple and rapid protocol
- Products are suitable for immediate downstream applications
- PCR clean-up
- Removes primers, primer-dimers, dNTPs and restriction enzymes
- DNA or dsRNA purification or concentration

PAN DNA Clean is a novel, inexpensive solution, which provides a column-free method for nucleic-acid purification. Using a simple and rapid procedure, PAN DNA Clean can be used to purify or concentrate DNA or dsRNA from PCR reactions or any enzymatic digests. This method is easy to follow, combining convenience, speed and excellent recovery rates.

Simple, flexible and column-free protocol

PAN DNA Clean removes proteins (such as restriction enzymes, polymerases, etc.), primers, primer-dimers and dNTPs. A very straightforward protocol allows the precipitation of nucleic acids ≥75 bp without the need for organic solvents, glass milk or expensive spin-columns. Unlike many column-based methods, PAN DNA Clean maximizes recovery with nucleic acid solutions, whether of low, medium or high concentration. PAN DNA Clean purifies nucleic acid without the use of chaotropic salts (which often contribute to denaturation of the DNA duplex). PAN DNA Clean enables the researcher to resuspend the cleaned-up nucleic acids in any buffer and volume of choice, thus permitting the purification process to be tailored specifically to suit the experiment.

Optimized nucleic acid recovery

PAN DNA Clean has been tailored to maximize the amount of nucleic acid recovered after purification, providing up to 98% recovery of the original sample for immediate downstream applications, such as cloning and sequencing. PAN DNA Clean exhibits great versatility, achieving unsurpassed recovery rates, independently of the amount of nucleic acid or its concentration.

Storage conditions

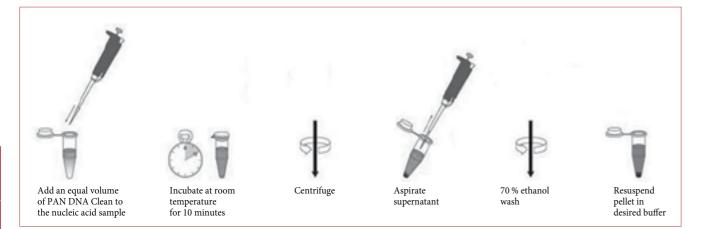
PAN DNA Clean solution can be stored at room temperature for 12 months. Do not freeze. Avoid exposure to light.



Lane 1: PCR mix before cleaning, 125 bp fragment Lane 2: PCR mix treated with PAN DNA Clean 1:1 PCR mix treated with PAN DNA Clean 1:2

Purification with PCR Clean up Kit (Silica membrane)

Lane 5: PANLadder I



1 x 5 ml PAN737042 PAN DNA Clean 2 x 12,5 ml PAN737046



Molecular Biology Reagents

Molecular Biology

Agarose (molecular grade)

Features

- DNase/RNase-free
- Excellent value and clarity
- High gel strength
- DNA/RNA electrophoresis
- Ideal for separating nucleic acids of a wide range of sizes, especially large fragments (> 10 Kb)

Agarose (DNase/RNase-free) is an extremely pure, high molecular biology grade agarose powder that has been extensively tested for RNase contamination. Agarose provides high resolution of DNA and RNA separated by electrophoresis and offers consistent resolution from lot

Storage conditions

Cool, dry place

Analytical specifications

Appearance:	White crystals or
	powder
Gel strength of 1.5% (w/v) gel:	> 1220g/cm ²
Fusion point:	88 - 90° C
Gelling temperature:	37 - 39° C
EEO:	0.05 - 0.1
Moisture:	< 7%
Sulphate:	< 0.06%
DNase and RNase:	Absent





Molecular Biology Reagents

dNTP Sets

Features and applications

- Ultra-pure: > 99% tris-phosphate by HPLC
- Extended shelf-life of 24 months at -20° C
- Free from PCR inhibitors
- DNase, RNase and Nickase free

Suitable for a wide variety of applications such as:

- Standard and long range PCR assays
- cDNA synthesis
- qPCR
- Microarrays
- DNA sequencing
- DHPLC
- Labelling

A set of ready-to-use molecular grade dNTP solutions consisting of 4 separate 100mM solutions of dATP, dGTP, dCTP and dTTP. For use in DNA polymerization reactions, DNA labelling and sequencing processes. Dependable PCR grade. All dNTPs are supplied as Lithium salts in purified water at pH 7.5. Lithium salts have greater resistance to repeated freezing and thawing cycles than Sodium salts, and Lithium salt dNTP preparations remain sterile over the entire shelf life due to the bacterio-static activity of Lithium towards various microorganisms.

Storage conditions

dNTP Set can be stored for 24 months at -20° C. Avoid multiple freeze/thaw cycles. For long-term storage, aliquoting is recommended.

Characteristics	dATP	dCTP	dGTP	dTTP
Product	dATP Lithium 100 mM Solution	dCTP Lithium 100 mM Solution	dGTP Lithium 100 mM Solution	dTTP Lithium 100 mM Solution
Nomenclature	2'-deoxyadenosine- 5'-triphosphate	2'-deoxyadenosine- 5'-triphosphate	2'-deoxyadenosine- 5'-triphosphate	2'-deoxyadenosine- 5'-triphosphate
Formula	$C_{10}H_{12}N_5O_{12}P_3Li_4$	$C_9H_{12}N_3O_{13}P_3Li_4$	$C_{10}H_{12}N_5O_{13}P_3Li_4$	$C_{10} H_{13} N_2 O_{14} P_3 Li_4$
Molecular Weight	514.9 g/mol	490.9 g/mol	530.9 g/mol	505.9 g/mol
λmax pH 7.0	259 nm	272 nm	252 nm	267 nm
ε at λmax @ pH7.0	15.4 E x mmol ⁻¹ x cm ⁻¹	9.1 E x mmol ⁻¹ x cm ⁻¹	13.7 E x mmol ⁻¹ x cm ⁻¹	9.6 E x mmol ⁻¹ x cm ⁻¹
A_{250}/A_{260}	0.78 ± 0.03	0.82 ± 0.03	1.16 ± 0.05	0.65 ± 0.03
A_{280}/A_{260}	0.15 ± 0.02	0.98 ± 0.03	0.66 ± 0.03	0.73 ± 0.02
Concentration	$100\text{mM} \pm 2\%$	$100\text{mM} \pm 2\%$	100mM ± 2%	100mM ± 2%
Appearance	Clear Colorless Solution	Clear Colorless Solution	Clear Colorless Solution	Clear Colorless Solution
pH of Solution	7.5	7.5	7.5	7.5
dNTP (HPLC Area)	≥ 99 %	≥ 99 %	≥ 99 %	≥ 99 %
dNDP (HPLC Area)	< 1 %	< 1 %	< 1 %	< 1 %
DNases, RNases,				
Nicking Activity	Negative	Negative	Negative	Negative
Storage	at -20 ° C	at -20 ° C	at -20 ° C	at -20 ° C
Stability	≤ 24 months	≤ 24 months	≤ 24 months	≤ 24 month

dNTP set (dA+dC+dG+dT)	100 mM 4 x 250 μl	PAN739025
	100 mM 4 x (4 x 250 μl)	PAN739026

Molecular Biology Reagents

dNTP Mix

Features and applications

- Convenient, pre-optimized and pre-mixed
- Ultra-pure: > 99% tris-phosphate by HPLC
- Extended shelf-life of 24 months at -20° C
- Free from PCR inhibitors
- DNase, RNase and Nickase free

Suitable for a wide variety of applications such as:

- Standard and long range PCR assays
- cDNA synthesis
- qPCR
- Microarrays
- DNA sequencing
- **DHPLC**
- Labeling

A ready-to-use molecular grade dNTP Mix containing dATP, dCTP, dGTP and dTTP at pH 7.5 as Lithium salts in purified water. The mix is designed to save handson time for researchers and minimize the possibility of contamination. For use in DNA polymerization reactions, DNA labeling and sequencing processes. Dependable PCR grade. Lithium salts have greater resistance to repeated freezing and thawing cycles than Sodium salts, and Lithium salt dNTP preparations remain sterile over the entire shelflife due to the bacteriostatic activity of Lithium towards various microorganisms.

dNTP Mix Reaction Guidelines

100 mM Mix contains 25 mM of each dNTP

Reaction Volume Master Mix Reactions 50 μl 0.5 μl 1000

40 mM Mix contains 10 mM of each dNTP

Reaction Volume Master Mix Reactions 50 μl 1.25 μl 400

This is a guide only, for long-range applications adjust accordingly.

Storage conditions

dNTP Mix can be stored for 24 months at -20° C. Avoid multiple freeze/thaw cycles. For long-term storage, aliquoting is recommended.

Typical analysis

Lithium salts, > 99% deoxynucleoside triphosphates (HPLC, area %), < 1% deoxynucleoside monophosphates and deoxynucleoside diphosphates.

The dNTPs are > 99% pure by HPLC and are free of DNase, RNase, Protease, phosphatase and nicking activity.

ANTED Miss (AA + AC + AC + AT)	20 μmol	40 mM	500 μl	PAN739043
dNTP Mix (dA+dC+dG+dT)	50 µmol	100 mM	500 μl	PAN739028







Chapter 8 PANsys

PANsys

PANsys 3000 135-136

PANsys 3000

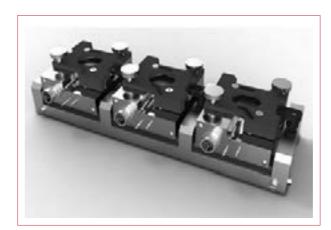
Fully automated cell culture systems

Introduction

PANsys 3000 is a highly automated cell culture system that enable a significant reduction in manual processes as well as a complete control and documentation of all important cell culture parameters. Apart from an automated control and supervision of these cell culture parameters, the system offers an integrated and automated evaluation of the cells in culture, determining growth rate, metabolism, and morphology.

The specially developed, patented cell culture chambers are the core of the cell culture units. In the cell culture chambers, cells are automatically supplied with appropriate media according to the selected culture parameters.

The optics with precision control and operation are equipped with a high-resolution CCD microscope (transmitted light, phase contrast and optional 6-channel fluorescence). This microscope records images of freely selectable points in the cell culture according to specified time intervals. These images are automatically saved by software and can be evaluated online. The automated cell-culture system PANsys 3000 is a nearly universally applicable tool for a highly efficient cell culture. Miscellaneous series of tests have proven the suitability of the system for the cultivation of a multitude of cells and cell lines under widely varying conditions.



PANsys 3000 features

Closed supply system with automated regulation of all necessary substances (CO₂, O₂, media, nutrients, temperature, activating substances or test drugs).

Automated cell-culture system where various cell types, cell lines and media can be cultivated under controlled conditions.

Up to six separate cell culture chambers (multi-chamber system) with individual equipment of each chamber and automatic adjustment of pre-selected cell culture parameters.

Continuous surveillance and documentation of all relevant cell culture parameters (temperature, CO₂, O₂, pH) with simultaneous microscopic monitoring of cell morphology and growth rate.

Life cell imaging: continuous video-microscopic monitoring of the cells with storage, documentation and analysis. Morphological changes and growth behaviour can be quickly detected and evaluated.

High-level microscopy system with a phase contrast and multi-channel fluorescence microscope for detailed and complex microscopy applications. Individual adaptation of optical characteristics (filters, channels, etc.)

Saving and documentation of all relevant recorded data of cell cultivation, including cell culture parameters and microscope data. Automated evaluation and analysis of all culture parameters and microscope images with powerful, modern software tools.





PANsys 3000

Different PANsys 3000 versions offered

PANsys 3000 Expert

Expert version with bright-field and phase contrast-microscope, fluidics pumps and incubation-chambers, including software package for system data control and image-analysis

PANsys 3000 Professional

Professional version with bright-field and phase contrast-microscope, optimized fluidics pumps and special chambers for flow-experiments, including software package for system data control and image analysis

PANsys 3000 HighEnd

High-end version with bright-field, phase contrast and 6-channel fluorescence microscope, sophisticated fluidics features, special cell culture chambers, high-end software package for system data control and image analysis

Optional add-on modules

O₂-pH Measurement

Special measuring unit for the detection of O₂ and pH in each cell culture chamber

Gas-Mixing Module Special gas-mixing module for an individual mixing of incubation gas (O₂-N₂-CO₂), suitable for special hypoxia- or hyperoxia experiments Cell Impedance Measuring

Lab-on-a-chip to analyze impedance, O₂, pH, ECAR and OCR in each cell culture chamber

Raman Spectroscopy

Special Raman spectroscopy unit for a fast and non-invasive detection of cellular pathways and cellular differentiation, label-free with minimal cellular interactions

Areas of application

- Stem cell research
- Development of serum-free media
- Cloning and differentiation studies
- Optimization of cell culture media
- Investigation of cell-cell interaction
- Drug development and substance screening
- Tissue cultivation and tissue engineering
- Cell cycle analysis
- Cellular cytotoxicity tests to reduce animal testing
- Endothelial flow experiments with shear-stress
- Simulation and optimization of production processes
- Customized applications

PANsys 3000 offers unique and extraordinary possibilities for in vitro cultivation and research on the most diverse cell lines under in vivo-like conditions. It reduces the gap between in vitro and in vivo applications through a powerful cell culture and life-cell imaging system with easy operability. PANsys 3000 enables the design of complex, artificial organ models, in order to determine specific cell behaviour and influencing factors, environmental conditions, or activating substances. PANsys 3000 improves developmental quality through the vastly detailed analysis and documentation of all data related to cell culture, including microscopic imaging. PANsys 3000 reduces your development costs through automation and a significant reduction of manual processes with much less resource consumption. PANsys 3000 reduces your development times through a parallelization of cell culture processes.



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