

# PeproTech Cell Culture Media **PeproGrow**<sup>TM</sup>

Redefining Media Quality Utilizing PeproTech Cytokine Products



www.peprotech.com

# **Table of Contents**

PeproTech Cell Culture Media Products
PeproGrow <sup>™</sup> hMSC Medium
PeproGrow <sup>™</sup> hESC Medium8
PeproGrow <sup>™</sup> Endothelial Media
PeproGrow <sup>™</sup> EPC Kit11
PeproGrow™ MacroV Kit11
PeproGrow <sup>™</sup> MicroV Kit11
PeproGrow <sup>™</sup> HEK293 Medium14
PeproGrow <sup>™</sup> CHO Medium16
PeproGrow-1 Serum-Free Cell Culture Supplement Kit 18
PeproGrow <sup>™</sup> Media Products Chart
Cell Culture Glossary

PeproTech products are for research use only and are not for therapeutic or diagnostic use. Copyright © 2021 Peprotech, Inc. All Rights Reserved.

PeproTech is a privately owned biotechnology company focused on the development and manufacturing of high quality cytokine products for the life science and cell therapy markets. As a leading researcher and manufacturer of cytokine products, PeproTech is pleased to offer media that contain our recombinant proteins and/or growth factors. These media formulations have been engineered by PeproTech's scientists and collaborators to provide the basic nutrients to grow and support several specific cell types.





### The following Cell Culture Media Product information is available on our website at www.peprotech.com:

- Complete List of Cell Culture Media Products\*
- Affordable and Competitive Product Prices
- Certificates of Analysis
- Instruction Manuals
- Frequently Asked Questions (FAQs)

\*Some media kit components are also available for individual purchase.

# PeproGrow<sup>™</sup> hMSC Medium

## Maintenance Medium for Human Mesenchymal Stem Cells



- Complete Media
- Superior growth rates
- Xeno-free, phenol red-free
- Phenotypic integrity
- Maintained multipotency
- Affordable and competitive pricing
- Works well with adipose tissue-derived, bone marrow-derived, umbilical cord-derived, placental-derived, and urine-derived MSCs
- Developed in collaboration with American CryoStem Corporation

PeproGrow<sup>™</sup> hMSC (Mesenchymal Stem Cell) Medium is a xeno-free, human serum-containing, phenol red-free complete media formulation originally designed for the *in vitro* expansion of adipose-derived human mesenchymal stem cells (ADMSCs) in the multipotent state. This media formulation has been shown to be suitable for the sustained growth of adipose tissue-derived, bone marrow-derived, umbilical cord-derived, placental-derived, and urine-derived MSCs in both adherent and suspension culture. For optimal results, culturing should be conducted on a surface coated with PeproTech's Animal-Free Human Vitronectin Matrix as a surface-coating reagent; however, other suitable extracellular matrix (ECM) proteins, such as fibronectin or vitronectin, can be used. PeproGrow<sup>™</sup> hMSC Medium was designed and developed by PeproTech in collaboration with American CryoStem Corporation, and is supplied as a 500mL bottle of PeproGrow<sup>™</sup> hMSC Basal Medium (Catalog# BM-XF-HMSC-500) containing a human serum component, and a separate, lyophilized vial of animal-free PeproGrow<sup>™</sup> hMSC Growth Factor Supplement (Catalog# GF-XF-HMSC-500). The addition of the separate, lyophilized growth factor supplement to the basal medium results in a complete medium containing all growth factors and supplements necessary for optimal expansion of human mesenchymal stem cells in culture.

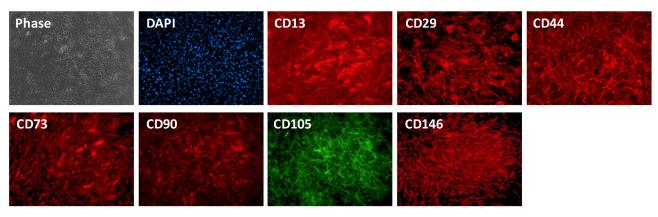


PeproGrow <sup>™</sup> hMSC Medium 500mL Kit	Catalog #XF-HMSC-500	
Basal Medium	BM-XF-HMSC-500	500mL
Growth Factor Component	GF-XF-HMSC-500	

### **PeproGrow<sup>™</sup> hMSC Medium Figures and Descriptions**

#### Figure 1

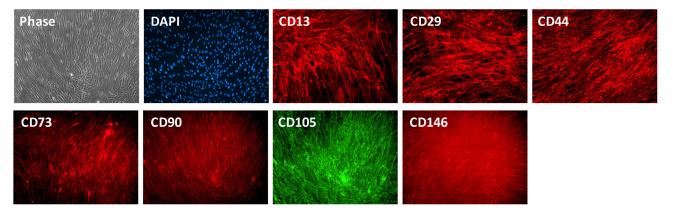
Fluorescent immunostaining of adipose tissue-derived mesenchymal stem cells (ADMSCs) grown in PeproGrow<sup>™</sup> hMSC Medium.



Cells were grown for 4 days on Animal-Free Human Vitronectin Matrix-coated chamber slides at 37°C in an atmosphere of 5% O<sub>2</sub> and 5% CO<sub>2</sub>. Following growth, the cells were rinsed once with PBS to remove debris and dead cells, then fixed for 20 minutes at RT with freshly-prepared 4% paraformaldehyde in PBS/HEPES, pH 7.4. Following fixation, cells were rinsed 3 times with PBS, blocked/permeabilized with 1x PERM/WASH (BD), and subjected to primary antibody staining (BioGems) at recommended concentrations overnight at 4°C. The following day, samples were washed 3 times with PBS, and incubated with secondary antibodies (ThermoFisher/Molecular Probes) diluted in 1:4 PERM/Wash:PBS as follows: all goat anti-rabbit antibodies were used at 1:1000, whereas goat anti-mouse isotypes were used at 1:500 dilutions. The samples were then washed 1 time with PBS containing 500 ng/ml DAPI, twice more in PBS, alone, and once in DI water to remove salts prior to mounting/cover-slipping in Aqua-Poly Mount (Polysciences) with 12 mm glass coverslips. The slides were allowed to dry overnight at RT prior to imaging using an Olympus IX71 Inverted microscope equipped with an inverted cooled CCD camera. Images were captured using iVision software (12 bit), pseudo-colored (DAPI = blue, AF488 = green, AF594 = red), and then exported to 8-bit format. In order to ensure specificity, all image capture times were adjusted to, or below, that of the isotype controls that revealed minimal background staining levels. As shown, PDMSCs grown in PeproGrow<sup>™</sup> hMSC Medium were strongly positive for CD13, CD29, CD44, CD73, CD90, CD105, and CD146.

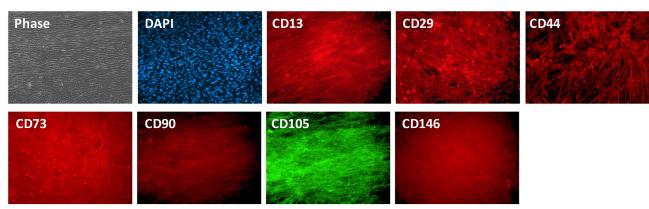
#### Figure 2

Fluorescent immunostaining of bone marrow-derived mesenchymal stem cells (BDMSCs) grown in PeproGrow™ hMSC Medium.



Cells were grown for 4 days on Animal-Free Human Vitronectin Matrix-coated chamber slides at 37°C in an atmosphere of 5% O<sub>2</sub> and 5% CO<sub>2</sub>. Following growth, the cells were rinsed once with PBS to remove debris and dead cells, then fixed for 20 minutes at RT with freshly-prepared 4% paraformaldehyde in PBS/HEPES, pH 7.4. Following fixation, cells were rinsed 3 times with PBS, blocked/permeabilized with 1x PERM/WASH (BD), and subjected to primary antibody staining (BioGems) at recommended concentrations overnight at 4°C. The following day, samples were washed 3 times with PBS, and incubated with secondary antibodies (ThermoFisher/Molecular Probes) diluted in 1:4 PERM/Wash:PBS as follows: all goat anti-rabbit antibodies were used at 1:1000, whereas goat anti-mouse isotypes were used at 1:500 dilutions. The samples were then washed 1 time with PBS containing 500 ng/ml DAPI, twice more in PBS, alone, and once in DI water to remove salts prior to mounting/cover-slipping in Aqua-Poly Mount (Polysciences) with 12 mm glass coverslips. The slides were allowed to dry overnight at RT prior to imaging using an Olympus IX71 Inverted microscope equipped with an inverted cooled CCD camera. Images were captured using iVision software (12 bit), pseudo-colored (DAPI = blue, AF488 = green, AF594 = red), and then exported to 8-bit format. In order to ensure specificity, all image capture times were adjusted to, or below, that of the isotype controls that revealed minimal background staining levels. As shown, PDMSCs grown in PeproGrow™ hMSC Medium were strongly positive for CD13, CD29, CD44, CD73, CD90, CD105, and CD146.

#### Figure 3



Fluorescent immunostaining of placental-derived mesenchymal stem cells (PDMSCs) grown in PeproGrow<sup>™</sup> hMSC Medium.

Cells were grown for 4 days on Animal-Free Human Vitronectin Matrix-coated chamber slides at 37°C in an atmosphere of 5% O<sub>2</sub> and 5% CO<sub>2</sub>. Following growth, the cells were rinsed once with PBS to remove debris and dead cells, then fixed for 20 minutes at RT with freshly-prepared 4% paraformaldehyde in PBS/HEPES, pH 7.4. Following fixation, cells were rinsed 3 times with PBS, blocked/permeabilized with 1x PERM/WASH (BD), and subjected to primary antibody staining (BioGems) at recommended concentrations overnight at 4°C. The following day, samples were washed 3 times with PBS, and incubated with secondary antibodies (ThermoFisher/Molecular Probes) diluted in 1:4 PERM/Wash:PBS as follows: all goat anti-rabbit antibodies were used at 1:1000, whereas goat anti-mouse isotypes were used at 1:500 dilutions. The samples were then washed 1 time with PBS containing 500 ng/ml DAPI, twice more in PBS, alone, and once in DI water to remove salts prior to mounting/cover-slipping in Aqua-Poly Mount (Polysciences) with 12 mm glass coverslips. The slides were allowed to dry overnight at RT prior to imaging using an Olympus IX71 Inverted microscope equipped with an inverted cooled CCD camera. Images were captured using iVision software (12 bit), pseudo-colored (DAPI = blue, AF488 = green, AF594 = red), and then exported to 8-bit format. In order to ensure specificity, all image capture times were adjusted to, or below, that of the isotype controls that revealed minimal background staining levels. As shown, PDMSCs grown in PeproGrow<sup>TM</sup> hMSC Medium were strongly positive for CD13, CD29, CD44, CD73, CD90, CD105, and CD146.

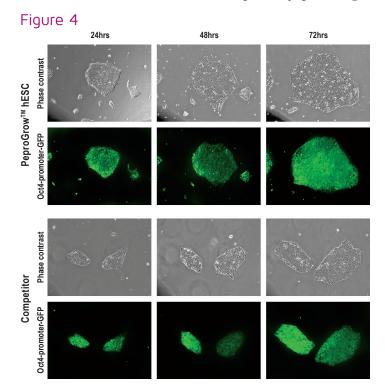
# PeproGrow<sup>™</sup> hESC Medium

## Maintenance Medium for hESCs and hiPSCs



- Phenol red-free and insulin-free
- Complete and chemically-defined
- High plating efficiency
- High quality recombinant growth factors from the leading manufacturer
- Developed in collaboration with and used in the Rutgers Stem Cell Training Course

PeproGrow<sup>™</sup> hESC Medium is a serum- and phenol red-free medium of a complete, chemically-defined formulation designed for feeder-free maintenance and expansion of both human embryonic stem cells (hESCs) and induced pluripotent stem cells (iPSCs) using Corning Matrigel® as a surface-coating matrix. This medium is intended for the culturing of hESCs and iPSCs in the undifferentiated, pluripotent state (SSEA4+/Oct4+), and demonstrates less than 15% spontaneous differentiation as indicated by flow cytometry. The proprietary formulation of the medium includes relevant growth factors, such as FGF2 (FGF-basic), but does not contain the insulin found in the majority of other hESC/iPSC media currently available on the market. PeproGrow<sup>™</sup> hESC Medium, which was designed and developed by PeproTech in collaboration with the Stem Cell Training Course at Rutgers University, is supplied as a 100mL, or 500mL, bottle of basal medium and a separate, lyophilized growth factor component.

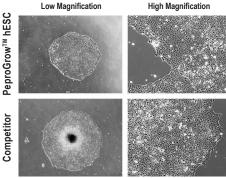


## The growth of H1 hESCs expressing turboGFP-NEO under the control of the Oct4 promoter.

H1 hESCs modified by lentivirus to contain the Oct4 promoter driving turboGFP, were cultured in PeproGrow<sup>™</sup> hESC Medium, were passaged using dispase and plated in PeproGrow<sup>™</sup> hESC Medium containing 2 µM Y-27632 onto Corning Matrigel® coated 6-well dishes. Cultures were fed daily and photographed 24, 48, and 72 hours post split. Green fluorescence represents maintenance of pluripotency as indicated by Oct4 promoter activity. PeproGrow<sup>™</sup> hESC Medium maintains cells in the pluripotent state like the competitor media and allows the cells to plate out better (faster confluency).

#### **PeproGrow<sup>TM</sup> hESC Medium Kit**

#### Figure 5

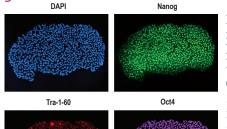


#### Cell Morphology of hESCs cultured in PeproGrow<sup>™</sup> hESC Medium.

This medium contains a unique blend of cytokines that at first may alter the cellular or colony appearance, making them more flat, and elastic.

however this morphological change appears less dramatic as the cell density increases, and over several passages. Stem cells will form standard tight circular colonies (left), and have expected morphology with large nucleus and small cytoplasm (right).

## Figure 6



#### Immunostaining of iPSCs cultured in PeproGrow<sup>™</sup> hESC Medium.

Cells were plated on 24-well dishes, paraformaldehyde fixed and then stained with DAPI and other indicated antibodies, which indicate cell pluripotency.

### **Specifications**

рН 7.35-7.40	
pm 7.55-7.40	
Osmolality 340-350 r	nOsm

HESC-500	
BM-HESC-500	500mL
GF-HESC-500	Vial for 500mL Basal Medium
HESC-100	
BM-HESC-100	100mL
GF-HESC-100	Vial for 100mL Basal Medium
-	BM-HESC-500       GF-HESC-500 <b>HESC-100</b> BM-HESC-100

# PeproGrow<sup>™</sup> Endothelial Media

## Maintenance Media for Endothelial Cells

PeproGrow<sup>™</sup> EPC, PeproGrow<sup>™</sup> MacroV, and PeproGrow<sup>™</sup> MicroV



- Complete media
- Antibiotic-free, antimycotic-free, antifungal-free, and phenol red-free
- Maintain outstanding endothelial cell morphology and function
- Increased activity of endothelial nitric oxide synthase (eNOS)

PeproTech offers three separate endothelial cell culture media formulations developed for the *in vitro* cultivation of: endothelial progenitor cells (EPCs; PeproGrow<sup>™</sup> EPC) derived from bone marrow or peripheral blood; endothelial cells from large vessels (PeproGrow<sup>™</sup> MacroV); and endothelial cells from small vessels (PeproGrow<sup>™</sup> MicroV). These media formulations maintain outstanding endothelial cell morphology and function, and increase the activity of endothelial nitric oxide synthase (eNOS), which account for a specific, crucial marker for endothelial cells. By doing this, the media provide an optimal cell culture environment for macrovascular and microvascular endothelial cells, as well as for EPCs; growing cells at rates that exceed commercially available media.

PeproTech's endothelial cell culture media kit is supplied as a 500mL bottle of basal medium and a separate growth supplement bottle that contains various essential growth factors and components for endothelial cell growth. Adding the growth supplement to the basal medium results in the complete culture medium. PeproTech's endothelial media do not contain antibiotics, antimycotics, antifungals, or phenol red, as these components can cause cell stress and masking effects that may reduce complete medium shelf life and influence experimental results.



### **Media Products**

PeproGrow <sup>TM</sup> EPC Kit (ENDO-BM & GS-EPC)	Catalog #700-EPC	
Basal Medium	ENDO-BM	500mL
Growth Supplement EPC	GS-EPC*	75mL

PeproGrow <sup>™</sup> MacroV Kit (ENDO-BM & GS-MacroV)	Catalog #700-MacroV	
Basal Medium	ENDO-BM	500mL
Growth Supplement MacroV	GS-MacroV*	25mL

#### **PeproGrow<sup>TM</sup> MicroV Kit (ENDO-BM & GS-MicroV)**

	_	
Basal Medium	ENDO-BM	500mL
Growth Supplement MicroV	GS-MicroV*	35mL

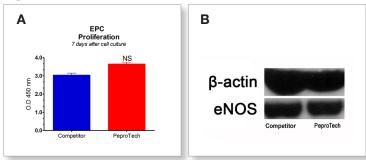
Catalog #700-MicroV

\* Contains FBS

## PeproGrow<sup>™</sup> Endothelial Media (continued)

### PeproGrow<sup>™</sup> Endothelial Media Figures and Descriptions

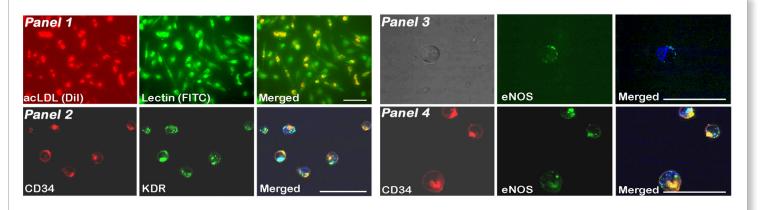
#### Figure 7



#### Endothelial Progenitor Cell, EPC Proliferation.

EPCs were seeded onto fibronectin-coated plates, and incubated for 7 days in the PeproGrow<sup>TM</sup> EPC Kit and a competitor's medium. Figure 7(A) represents the proliferative ability of EPCs assessed 7 days after cell cultivation using the XTT assay according to the manufacturer protocol. The proliferative ability of EPCs was expressed as the average optical density (O.D.) calculated using a plate reader for two independent assays run in triplicate. Figure 7(B) represents a standard Western Blot assay.

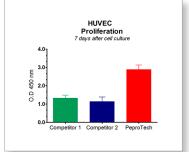
#### Figure 8



#### Endothelial Progenitor Cell, EPC Characterization.

EPCs were cultured for 7 days. Each description correlates to images from left to right: Panel 1: Acetylated LDL uptake by adherent spindle-shaped EPCs, FITC-conjugated lectin UEA-1 binding to the surface of EPCs, and positive double-stained (merged image) EPCs for acetylated LDL uptake and lectin binding. Panel 2: Immunofluorescence detection of CD34 antigen (red), KDR (green) on the surface of EPCs, and merged image. Panel 3: Immunofluorescence detection of eNOS on a single non-stained EPC (green). Panel 4: Immunofluorescence detection of CD34 antigen on the EPCs surface (red), eNOS (green), and merged image. The EPCs nuclei were stained with the blue fluorescent DNA dye DRAQ5<sup>TM</sup>.

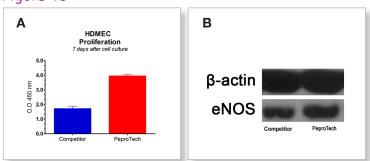
#### Figure 9



#### Macrovascular Endothelial Cell, HUVEC Proliferation.

HUVECs were seeded onto fibronectin-coated plates and incubated for 7 days in the PeproGrow<sup>TM</sup> MacroV Kit, and two competitors' media. The proliferative ability of HUVECs was assessed 7 days after cell cultivation using the XTT assay following the manufacturer's protocol. The proliferative ability of HUVECs was expressed as the average of optical intensity (O.D.) calculated using a plate reader from two independent assays run in triplicate.

#### Figure 10



#### Microvascular Endothelial Cell, HDMEC Proliferation.

HDMECs were seeded onto fibronectin-coated plates and incubated for 7 days in the PeproGrow<sup>TM</sup> MicroV Kit and a competitor's medium. Figure 10(A) represents the proliferative ability of HDMECs assessed 7 days after cell cultivation using the XTT assay following the manufacturer's protocol. The proliferative ability of HDMECs was expressed as the average of optical density (O.D.) calculated using a plate reader from two independent assays run in triplicate. Figure 10(B) represents a standard Western Blot assay.

### **Recommended Cell Types**

#### PeproGrow<sup>™</sup> EPC is recommended for Endothelial Progenitor Cells:

Human Endothelial Progenitor Cells (hEPCs)

PeproGrow™ MacroV is recommended for Macrovascular Endothelial Cells:

Human Umbilical Vein Endothelial Cells (HUVECs) Human Umbilical Artery Endothelial Cells (HUAECs) Human Aortic Endothelial Cells (HAoECs)

Human Pulmonary Artery Endothelial Cells (HPAECs)

Human Saphenous Vein Endothelial Cells (HSaVECs)

#### PeproGrow<sup>™</sup> MicroV is recommended for Microvascular Endothelial Cells:

Human Coronary Artery Endothelial Cells (HCAECs) Human Pancreatic Microvascular Endothelial Cells (HPaMECs) Human Dermal Microvascular Endothelial Cells (HDMECs) Human Pulmonary Microvascular Endothelial Cells (HPMECs) Human Dermal Lymphatic Endothelial Cells (HDLECs) Human Brain Microvascular Endothelial Cells (HBMECs)

### **Specifications**

Sterility	Absence of bacteria, mycoplasma and fungi	
pH	7.4-7.8	
Osmolality	260-280 mOsm	
Endotoxin Testing	<0.5 EU/mL	

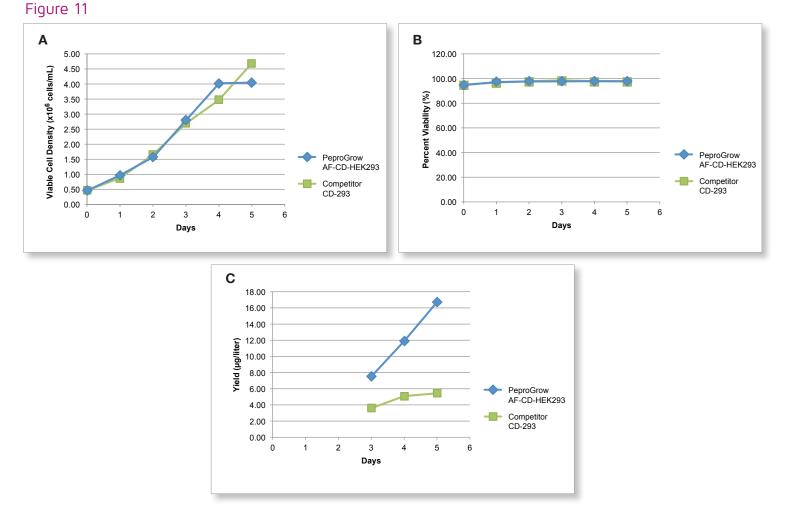
# PeproGrow<sup>™</sup> HEK293 Medium

## Maintenance Medium for HEK293 Cells



- Animal component-free, serum-free, protein-free, chemically-defined medium
- Complete medium containing L-alanyl-L-Glutamine for direct product use
- High recombinant protein expression

PeproGrow<sup>™</sup> HEK293 Medium is an animal component-free, protein-free, serum-free, chemically-defined, complete medium formulation for the *in vitro* cultivation of HEK293 cells (Thermo Fisher Scientific FreeStyle<sup>™</sup> 293-F cells, catalog number R790-07). This medium is intended for recombinant protein expression in suspension culture, which is recommended for a 5-day batch culture with a seeding density of 0.6 x10<sup>6</sup> cells/mL. This ready-to-use medium contains L-alanyl-L-Glutamine, amino acids, vitamins, and salts. An adaptation process is not required for Thermo Fisher Scientific FreeStyle<sup>™</sup> 293-F cells.



Figures 11 (A, B, C) illustrate HEK293 BMP-6 expression in PeproGrow<sup>™</sup> HEK293 Medium and competitor CD-293 medium.

#### **PeproGrow<sup>TM</sup> HEK293 Media**

PeproGrow <sup>™</sup> HEK293 Medium	Catalog #AF-CD-HEK293	1 L

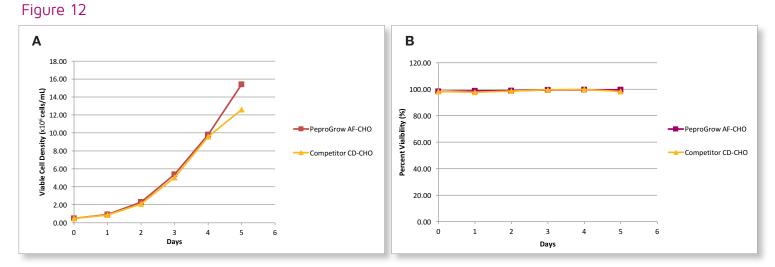
# PeproGrow<sup>™</sup> CHO Medium

## Maintenance Medium for CHO-S cell lines



- Animal component-free, serum-free, and protein-free
- Complete medium containing L-alanyl-L-Glutamine for direct product use
- High recombinant protein expression

PeproTech offers PeproGrow<sup>™</sup> AF-CHO Medium, for the *in vitro* cultivation of Chinese Hamster Ovary-S cells (Thermo Fisher Scientific catalog numbers 11619-012, R800-07, or A11557-01). This medium is intended for recombinant protein expression in suspension culture. PeproGrow<sup>™</sup> AF-CHO is an animal component-free, serum-free, protein-free, complete medium formulation. This ready-to-use medium contains L-alanyl-L-Glutamine, amino acids, vitamins, salts and non-animal-derived hydrolysates, which are recommended for a 5-day batch culture with a seeding density of 0.5 x10<sup>6</sup> cells/ mL. An adaptation process is not required for Thermo Fisher Scientific CHO-S cells.



Figures 12 (A, B) illustrate CHO-S cell density and percent viability of C1 Inhibitor-producing cells in PeproGrow<sup>™</sup> AF-CHO Medium and competitor CD-CHO Medium. Cell expression varies by protein product.

#### Figure 13



The western blot illustrates CHO-S cell culture sample production of C1 Inhibitor post 5-day culture.

lane 1: PeproGrow<sup>TM</sup> AF-CHO

lane 2: Commercial CD-CHO

#### **PeproGrow**<sup>TM</sup> CHO Media

PeproGrow<sup>TM</sup> AF-CHO Medium

## PeproGrow-1 Serum-Free Cell Culture Supplement Kit

Serum-free cell culture supplement kit for adherent HEK293, HeLa, and A549 cells



- Serum-free, animal-free, and protein-free medium supplement
- Chemically-defined medium supplement

PeproGrow-1 (catalog number 700-C100) is a serum-free cell culture media supplement formulation designed to sustain the growth of adherent mammalian cell lines, and has been tested with HEK293, HeLa, and A549 cells. This kit may potentially improve the culturing conditions of other adherent cells, however suitability for cells other than those pretested is not guaranteed. PeproGrow-1 is an animal component-free, protein-free, chemically-defined formulation. This kit is intended to be used with DMEM/F12 basal media (Thermo Fisher Scientific catalog #10565, or catalog #31331 for customers located outside the USA) and contains enough material to supplement 10L of media.

Note: DMEM/F12 media can contain phenol red, or phenol red can be added at the customer's discretion. DMEM/F12 media should not contain HEPES. Another vendor's DMEM/F12 can be purchased, however, PeproTech only tests PeproGrow-1 (Catalog # 700-C100) using Thermo Fisher Scientific DMEM/F12.

### PeproGrow-1 Serum-Free Cell Culture Supplement Kit Components:

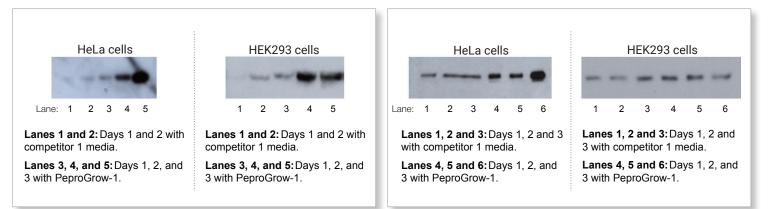
#### • Serum Replacement Solution (Catalog #SR-100):

This 100x serum replacement solution contains non-animal-derived, chemically-defined salts, designed to replace serum in HEK293, HeLa, and A549 cell culture media.

#### • Lipid Mixture Solution (Catalog #LM-200):

This 200x lipid mixture solution contains non-animal-derived fatty acids and lipids, designed to improve cell growth in serum-free media.

#### Figure 14



Western blot for Human TLR-3 production using PeproGrow-1 Serum-Free Cell Culture Supplement Kit Western blot for Human sDLL-1 production using PeproGrow-1 Serum-Free Cell Culture Supplement Kit

PeproGrow-1 Kit (SR-100 & LM-200)	Catalog #700-C100	
Serum Replacement Solution	SR-100	
Lipid Mixture Solution	LM-200	

# **PeproGrow™** Media Products Chart

Media	Cell Type(s)	Culture Type	Complete	Chemically- Defined
<b>PeproGrow-1 Kit</b> (LM-200 & SR-100) Lipid Mixture Solution Serum Replacement Solution	HeLa, HEK293, A549	Adherent		$\checkmark$
CHO Medium PeproGrow™ AF-CHO	Thermo Fisher Scientific CHO-S cell lines (Catalog numbers 11619-012, R800-07, A11557-01)	Suspension		
HEK293 Medium PeproGrow™ HEK293	Thermo Fisher Scientific FreeStyle™ 293-F cells (Catalog number R790-07)	Suspension		$\checkmark$
Human ESC Media PeproGrow™ hESC	Human ESCs and Human iPSCs	Adherent		$\checkmark$
Animal-Free Human Vitronectin Matrix PBS+ Kolliphor P 188 Cell Passaging/Non-Enzymatic Detachment Buffer PBS+HEPES+EDTA Human MSC Medium				-
PeproGrow™ hMSC Companion Products for Human hMSC Media	Human Mesenchymal Stem Cells	Adherent & Suspension	V	
Animal-Free Human Vitronectin Matrix and Buffer Ki Animal-Free Human Vitronectin Matrix PBS + Kolliphor P 188	t			
<b>Endothelial Media</b> <b>PeproGrow™ EPC Kit</b> (ENDO-BM & GS-EPC) Basal Medium Growth Supplement EPC	Human Endothelial Progenitor Cells	Adherent		
<b>PeproGrow™ MacroV Kit</b> (ENDO-BM & GS-MacroV) Basal Medium Growth Supplement MacroV	Human Macrovascular Endothelial Cells	Adherent		
PeproGrow <sup>™</sup> MicroV Kit (ENDO-BM & GS-MicroV) Basal Medium Growth Supplement MicroV	Human Microvascular Endothelial Cells	Adherent		

\*Select media products have individual components available for purchase.

Animal-Free	Xeno-Free	Serum-Free	Protein-Free	Catalog Number	Size	Additional Note(s)
$\checkmark$		$\checkmark$	$\checkmark$	<b>700-C100*</b> LM-200 SR-100	Kit 55mL 100mL	This product is a Serum-Free Cell Culture Supplement Kit to be used in conjunction with DMEM/F-12 basal media (Thermo Fisher Scientific Catalog #10565, or use Catalog #31331 for customers located outside the USA).
				AF-CHO	1L	Ready-to-use. Contains non-animal-derived hydrolysates.
		$\checkmark$	$\checkmark$	AF-CD-HEK293	1L	Ready-to-use.
		$\checkmark$		HESC-500 HESC-100	500mL 100mL	Phenol red-free. Supplied with a lyophilized growth factor component.

		AF-VMB-220	<b>Kit</b> 500µg 220mL	
		CPD-125	<b>Kit</b> 125mL	
		XF-HMSC-500	500mL	Phenol red-free. Supplied with a lyophilized growth factor component. This basal medium contains human serum. This basal medium has additional companion products available for purchase.

AF-VMB-220	<b>Kit</b> 500µg 220mL	
<b>700-EPC*</b> ENDO-BM GS-EPC	Kit 500mL 75mL	Growth Supplement EPC supplied as a frozen bottle.
<b>700-MacroV*</b> ENDO-BM GS-MacroV	Kit 500mL 25mL	Growth Supplement MacroV supplied as a frozen bottle.
<b>700-MicroV*</b> ENDO-BM GS-MicroV	Kit 500mL 35mL	Growth Supplement MicroV supplied as a frozen bottle.

## **Cell Culture Glossary**

Α

Adherent Cell Culture: Cells that form a single layer on an artificial substrate system in a medium, such as in a T-flask or a roller bottle.

Adult Stem Cell: A stem cell in the adult system that is in an undifferentiated state and maintains the ability to differentiate, generally into a cell type from its tissue of origin.

Animal-Free Medium: The medium does not contain animal-derived components.

Aseptic Technique: Procedures that are performed under sterile conditions. This prevents the introduction of fungi, bacteria, viruses, mycoplasma or other microorganisms into cell culture or other laboratory culturing conditions. These procedures may also prevent cell cross-contamination.

Basal Medium: An incomplete simple cell culturing medium that does not contain supplemental nutrients or growth factors.

**Blastocyst**: This is the early stage of an embryo composed of a fluid-filled cavity, an inner cell mass, and an outer cell mass. Embryonic stem cells are derived from the inner cell mass, while the outer cell mass forms the trophoblast.

Cellular Differentiation: The process that occurs when a cell becomes specialized for a particular cell
type and its function(s).

Chemically-Defined Medium: The chemicals contained in the medium are known and quantifiable. This medium does not contain yeast, animal or plant protein hydrolysates.

Complete Medium: All the necessary components/nutrients for the cell type are present for culturing.

**Defined Medium**: This is also known as chemically-defined medium. All the chemicals and quantities of the medium content are known.

Growth Supplement: This is an additional supplement that PeproTech supplies with some media products. The researcher adds this to the medium upon use, resulting in a complete medium, which contains growth factors/ proteins necessary for the particular cell type's growth.

Human Embryonic Stem Cell: A pluripotent stem cell that is derived from the inner cell mass of a blastocyst and can differentiate into different cell types.

Induced Pluripotent Stem Cell (iPSC): A stem cell that has been reprogrammed from an adult cell and has the ability to differentiate into other cell types.

Medium/Media (plural form of medium): The liquid solution used to grow and cultivate a particular cell type.

Pluripotent Stem Cell: A stem cell that is derived from the inner cell mass of the blastocyst and can differentiate into different cell types.

**Progenitor Cell**: A cell that can differentiate, but cannot renew itself. Generally, a progenitor cell is at a further stage of differentiation.

Protein-Free Medium: The medium does not contain protein, such as insulin, transferrin, albumin, and other protein growth factors.

Serum-Free Medium: The medium does not contain serum.

Suspension Cell Culture: Cells that are free floating in a medium, such as in a shaker flask on an orbital shaker, or a stirred tank bioreactor.

Undefined Medium: The medium may contain ingredient(s), such as a yeast extract, and animal or plant protein hydrolysates, in which the mixture of the chemical reagents is in unknown proportions.

Xeno-Free Medium: By some definitions xeno-free medium does not contain animal products, i.e., all the components are of non-animal origin. A xeno-free product may contain human-derived reagents.

# OUR QUALITY YOUR DEDICATION



5 Cedarbrook Drive Cranbury, NJ 08512

Ph: 800.436.9910 Fax: 609.497.0321 info@peprotech.com www.peprotech.com