Datasheet

SERAplex NTA

Defined Serum Substitute for Adherent Cells

Product	Description	Catalogue-No.	Size
SERAplex NTA	**Defined serum substitute for adherent cells**	**ST04-95070** **ST04-95700** **ST04-95750**	**50 ml** **100 ml** **500 ml**

Product description

SERAplex NTA is a ready-to-use, chemically defined serum substitute for the cultivation of adherent cells under serum-free conditions or to significantly reduce the amount of serum in cell culture. It supports the growth of many adherent cell types in an optimum manner without any extra handling compared to serum.

Storage conditions

Storage: -20°C (in the dark)
Stability: 2 years from date of production
Size: 50 ml, 100 ml, 500 ml, other sizes on request

Composition

SERAplex NTA contains purified proteins, lipids, salts, amino acids, trace elements, attachment factors and hormones in an optimized formulation. SERAplex NTA contains no growth factors, undefined hydrolysates or peptones.

Suitability

SERAplex NTA is suitable for the cultivation of a variety of adherent cells under serum-free culture conditions (please see figure 1) or to reduce the necessary FBS amount in cell culture.

Effect of SERAplex NTA in different cell lines

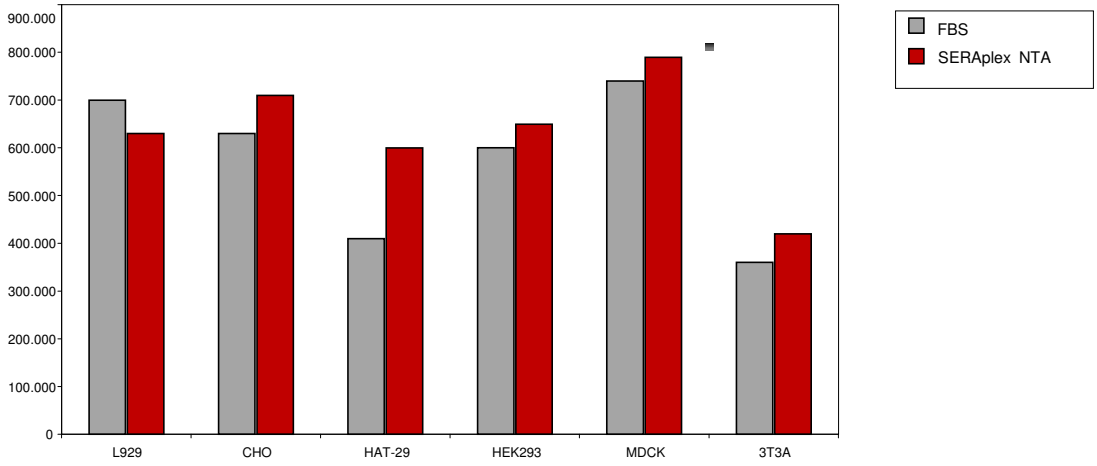

Fig. 1: Efficiency and growth stimulation of SERAplex NTA compared to FBS (10% each in DMEM/F12) at 7 day culture in different cell types

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Special advantages
SERAplex NTA is designed to replace or to reduce serum in the cell culture in a very simple manner. In most cases there is no need to change the basal medium. As SERAplex NTA is fully chemically defined and contains no peptones or hydrolysates, lot testing is no more necessary. It also allows high reproducibility and a simplified downstream process. SERAplex NTA contains no growth factors and enables defined proliferation and differentiation of stem cells. Characterization studies of growth factors will obtain more reproducible and clearer results. SERAplex NTA is also useful to develop sensitive cell-based *in vitro* tests and coculture procedures. For cell lines which require specific growth factors these should be added in a concentration as previously used.

Instructions for use
SERAplex NTA can be stored and used in the same manner as serum.

- Thaw SERAplex at maximum 37 °C. Please avoid repeated freeze-thaw cycles!
- To replace serum: Use the same basal medium and the same concentration of SERAplex as FBS. The performance can be further improved by optimizing the concentration of SERAplex or modifying/changing the basal medium[a] (Table 1)
- To reduce serum concentration: Use the same basal medium and add the same amount of SERAplex as the reduced amount of serum, until the minimal necessary concentration of FBS is found (1 to 2.5 % in most cases). The performance can be further improved by optimizing the the concentration of SERAplex or modifying/changing the basal medium[a] (also see adaptation instruction and table 1).
- Recommended inoculation cell density: 5.000 – 20.000 cells/cm^2.

- Solve cells as usual from the cell culture vessel (e.g. 0.25% trypsin inhibitor, Cat.No. ST10-033100 or Accutase®, Cat.No. ST10-21100). Once the cells have become round and detach from the surface inactivate trypsin with trypsin inhibitor (Cat.No. ST10-034100): Simply resuspend cells in about 1 ml trypsin inhibitor solution for every ml of trypsin solution used for dissociation. Note that Accutase® does not need to be inhibited.

Depending on the cell type, some differences in morphology or proliferation rate may be observed with varying standard media. Most applications were performed with DMEM and DMEM/F12 for adherent cells. Make sure that L-glutamine is present in sufficient quantity. The optimal SERAplex NTA concentration should be determined for each cell line. Tests can be started at a SERAplex NTA concentration of 10%, as with most cells the best results were obtained at this concentration.

Please note: For more demanding cells an adaptation to SERAplex NTA may be necessary.

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Adaptation instructions for SERAplex NTA

Precondition for a successful transition are vital cells (trypan blue exclusion staining), which should be harvested in the logarithmic growth phase. If 10% FBS was used in the original protocol,

Step 1: 7.5 % FBS + 2.5 % SERAplex

- Seed cells at 5 x 10^3 – 20 x 10^3 cells/cm^2.
- Observe cells under a microscope, at about 90 % confluence passage the cells for another 2-3 passages.

If normal growth is obtained transfer cells into:

Step 2: 5 % FBS + 5 % SERAplex

- Seed cells at 5 x 10^3 – 20 x 10^3 cells/cm^2 .
- Observe cells under a microscope, at about 90 % confluence passage the cells for another 2-3 passages.

If normal growth is obtained transfer cells into:

Step 3: 2.5 % FBS + 7.5 % SERAplex

- Seed cells at 5 x 10^3 – 20 x 10^3 cells/cm^2 .
- Observe cells under a microscope, at about 90 % confluence passage the cells for another 2-3 passages.

If normal growth is obtained transfer cells into:

Step 4: 1 % FBS + 9 % SERAplex

- Seed cells at 5x10^4 – 10x10^5 cells/ml.
- Observe cells under a microscope, at about 90 % confluence passage the cells for another 2-3 passages.

If normal growth is obtained transfer cells into:

Step 5: 10 % SERAplex

- Seed cells at 5 x 10^3 – 20 x 10^3 cells/cm^2.

Observe cells under a microscope.

For some cells an adaptation to serum-free conditions is difficult to reach or even impossible.

The following measures may help to facilitate a successful adaptation:

- Reseeding with a higher cell amount (about 2x to 4x of the usual cell density).
- Addition of growth factors (if known, which factors have a positive effect on the relevant cells).
- Coating the culture dishes or flasks with attachment factors (e.g. fibronectin, laminin, collagen, gelatine, etc).
- Change the basal medium. Note: A change of the basal medium to a richer or more complex formulation may be all that is needed to achieve growth in serum free condition.

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Table 1: Comparison of Cell Growth in 10% SERAplex NTA in different Basal Media versus Cell Growth in 10% FBS in different Basal Media

Cell Line	Origin	Basal Medium	Percentage of Growth 10% SERAplex NTA	Percentage of Growth 10% FBS
HEK 293 T	Renal cells, human embryonic	DMEM/F12 alpha-MEM DMEM	105% 76% 62%	100%
MDCK	Renal cells, canine	DMEM/F12 McCoy's 5A alpha-MEM	102% 91% 106%	100%
MDBK	Renal cells, bovine	RPMI 1640 McCoy's 5A DMEM	122% 135% 131%	100%
L 929	Fibroblasts, mouse	DMEM RPMI 1640 Ham's F-12	97% 78% 128%	100%
HT-29	Colon Carcinoma, human	IMDM DMEM/F12 alpha-MEM	108% 98% 96%	100%
HeLa S3	Cervix carcinoma epithel, human	Glasgow MEM IMDM EMEM	106% 72% 100%	100%
CHO	Ovarial cells epithel, Chinese hamster	DMEM/F12 IMDM alpha-MEM	106% 97% 82%	100%
CHO-Luc	Ovarial cells epithel, Chinese hamster, transfected	IMDM DMEM alpha-MEM	86% 97% 84%	100%
3T3A	Fibroblasts, mouse	RPMI 1640 McCoy's 5A DMEM/F12	98% 72% 97%	100%
MCF-7	Mammary carcinoma, human	Ham's F-12 DMEM/F12 RPMI 1640	292% 176% 214%	100%
RAW 264.7	Macrophages, mouse	McCoy's 5A DMEM/F12 alpha-MEM	40% 67% 38%	100%

Technical support

For technical support, questions or remarks please contact your local PAN-Seratech partner or the technical department of PAN-Seratech via email (info@pan-seratech.com) or phone +49-8543-601630.

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[a] As a basal medium, standard media such as RPMI 1640, DMEM (high or low glucose), DMEM/F12, IMDM etc. can be used. Make sure that L-glutamine is present in sufficient quantity (supplement L-glutamine as needed).