Product Information

**Stemline® Hematopoietic Stem Cell Expansion Medium**
without antibiotics or cytokines

Catalog Number S0189

**Stemline II Hematopoietic Stem Cell Expansion Medium**
without antibiotics or cytokines

Catalog Number S0192

Storage Temperature 2-8 °C

Synonyms: Hematopoietic Progenitor Cell Expansion Medium, CD34⁺ Expansion Medium

**Product Description**
The use of human Hematopoietic Stem Cells (HSC) isolated from umbilical cord blood (CB) as a source of cellular reconstitution following high-dose chemotherapy is now a common therapeutic modality for the treatment of malignancy. However, due to the low yield of HSC from CB, HSC transfusion is most effective in children and has limited application in adults. In order to obtain sufficient numbers of cells for applications of this therapeutic approach in adults, ex vivo expansion has been utilized to ensure successful engraftment and minimize the short-term effects of neutropenia and thrombocytopenia.

We have developed two serum-free media formulations for the optimal expansion of CB HSC, Stemline Hematopoietic Stem Cell Expansion Medium, Catalog Number S0189, and Stemline II Hematopoietic Stem Cell Expansion Medium, Catalog Number S0192. These media promote the expansion of CD34⁺ hematopoietic stem cells derived from umbilical cord blood and are also effective for the ex vivo expansion of HSC isolated from bone marrow or mobilized peripheral blood. These media support high viable cell densities. The elimination of serum reduces variability in the performance of the media and eliminates safety risks associated with possible adventitious agents in serum.

**Precautions and Disclaimer**
This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

The safety and efficacy of these products have not been established for clinical applications.

**Components**
Stemline Hematopoietic Stem Cell Expansion Medium and Stemline II Hematopoietic Stem Cell Expansion Medium are both proprietary formulations without antibiotics and cytokines. The Stemline Hematopoietic Stem Cell Expansion Medium, Catalog Number S0189, does not contain glutamine.

Human serum albumin is the only animal-derived protein in both media. It has been found to be non-reactive (donor level) for anti-HIV 1 & 2, anti-HCV and HBsAg. Handle as if potentially infectious.

Stemline Hematopoietic Stem Cell Expansion Medium, Catalog Number S0189, also contains cholesterol from sheep wool. Stemline II Hematopoietic Stem Cell Expansion Medium, Catalog Number S0192, contains no other animal-derived components.

**Preparation Instructions**
Media are supplied as sterile liquids (1X) and must be supplemented with cytokines and antibiotics, if desired. Stemline Hematopoietic Stem Cell Expansion Medium, Catalog Number S0189, must also be supplemented with glutamine. Add 20 ml of 200 mM L-glutamine solution or 0.5844 g powder (irradiated) per liter of medium.
Storage/Stability
These media are stable, when stored at 2-8 °C and protected from light, until the date indicated on the label.

Procedure
Plating Cultures
1. Prepare either fresh or frozen CD34+ cells as directed by the supplier or in accordance with established protocols.
2. Count cells using a hemacytometer.
3. Transfer the proper number of cells to the desired culture vessel with medium supplemented with cytokines (and antibiotics if desired).
4. Place the culture vessel in a humidified incubator at 37 °C and 5% CO2.

Product Profile
Sigma’s Stemline Hematopoietic Stem Cell Expansion Media demonstrate excellent expansion of total nucleated cells (TNC), committed progenitors (CD34+/CD38−) and primitive progenitors (CD34+/CD38+). These products were compared with several other commercially available serum-free expansion media for their ability to expand cord blood CD34+ cells in a 24-well microplate culture system. For these small-scale experiments, triplicate 1-ml cultures of 10,000 cord blood CD34+ cells/ml were incubated for 10 days in Stemline media or other commercial products containing 100 ng/ml each of SCF (Stem Cell Factor, Catalog Number S7901), G-CSF (Granulocyte-Colony Stimulating Factor, Catalog Number G0407) and TPO (Thrombopoietin, Catalog Number T1568).

The study was then expanded to a 2-step, clinical-scale protocol using Teflon® culture bags, which were assayed for TNC expansion prior to transplantation into NOD/SCID mice. For clinical-scale studies, cord blood CD34+ cells were cultured for 7 days in 100 ml Teflon culture bags containing 50 ml of culture medium plus cytokines. Cells were harvested from these bags and a 5-ml aliquot was transferred to a second 100 ml Teflon bag containing 50 ml of selected medium plus cytokines and cultured for an additional 7-day culture period. At the end of both culture protocols, cells were harvested and assayed as previously described. The data are summarized in Figures 1, 2, and 3. Both media demonstrated superior expansion of total nucleated cells (TNC), as well as both committed and primitive progenitors. The expanded cells were able to successfully engraft (first and second recipient) NOD/SCID mice (data not shown).

Figure 1. Comparison of Stemline Media to Commercially Available Products. Cells were seeded in triplicate at 10,000 cells per well in 24-well tissue culture plates containing either Stemline Hematopoietic Stem Cell Expansion Medium, Stemline II Hematopoietic Stem Cell Expansion Medium or one of several leading competitors’ media. The wells were supplemented with 100 ng/ml each of SCF, G-CSF, and TPO. Each well was triturated and harvested after a 10-day expansion. TNC were counted using a hemacytometer and averaged for all 3 wells in each condition.

Figure 2. Comparison of Stemline Media in Clinical-Scale 14-Day Expansions in Teflon Bags. Cells were seeded in Teflon culture bags containing either Stemline Hematopoietic Stem Cell Expansion Medium or Stemline II Hematopoietic Stem Cell Expansion Medium. The bags were supplemented with 50 ng/ml of FLT3L and 100 ng/ml each of SCF, G-CSF, and TPO. The cells were expanded for 7 days prior to being harvested and seeded into a new culture bag for an additional 7 days. Cells were harvested from the second
culture bag and TNC were counted. These expanded cells were used to successfully engraft NOD/SCID mice (data not shown).

Figure 3. Flow Cytometric Analysis of CD34⁺ Cord Blood Cells from the Clinical-Scale Expansion in Teflon Bags. Cells were expanded as previously described and assayed for CD34 and CD38 using standard flow cytometry procedures.

References

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