

High quality and yield

Standardized and clinically compliant

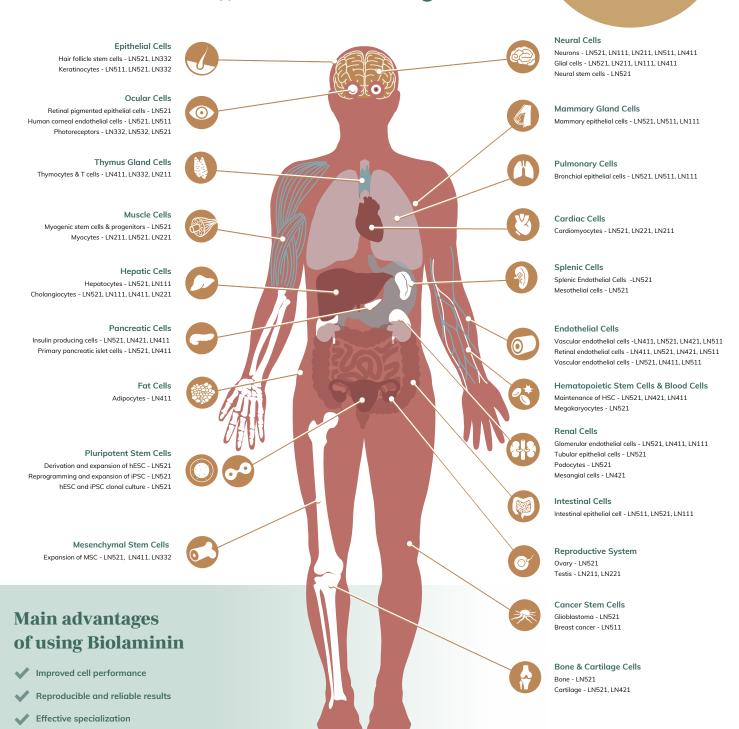
# Tissue-specific cell culture matrices

Imitate the natural cell-matrix interactions for improved cell functionality

The chemically defined and animal origin-free laminin cell culture matrices, Biolaminin®, allow you to imitate the natural cell-matrix interaction in vitro. Laminins are key components of the extracellular matrix. Through their interactions with specific receptors, laminins trigger the authentic cellular responses, pivotal for cell anchorage, survival, proliferation, migration, organization and specialization, leading to improved cell functionality.

Read more about different cell applications for our Biolaminins

We offer an expansive
portfolio of recombinant
laminin proteins for a variety
of applications, including
reliable expansion of pluripotent
cells and differentiation and
maintenance of specialized
cell types



## **Highlighted Biolaminin applications**



#### Robust self-renewal of high quality hPSCs on the Biolaminin 521 stem cell matrix

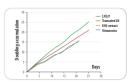
Laminin-521 is a key cell adhesion protein of the natural stem cell niche. The Biolaminin 521 (LN521) substrate supports efficient expansion at low dénsities of single-cell plated human pluripotent stem cells (hPSCs) under defined and animal origin-free  $conditions. \, LN521 \, is \, compatible \, with \, any \, medium \, and \, supports \, weekend-free feeding. \, Importantly, \, the \, cells \, behave \, predictably, \, and \, conditions \, conditions$ are homogenously pluripotent and karyotypically stable.



LN521 is naturally expressed and secreted by hPSCs in the inner cell mass of the embryo



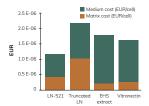
hPSCs can be seeded as single cells without ROCKI (day 0), grown as a homogenous monolayer (day 3) and can be cultured to high confluence without spontaneous differentiation.



hPSCs propagate faster on LN521 compared to other feeder-free



The hPSCs remain pluripotent (Oct4+; pink) and show no areas of differentiation (only DAPI staining;



Due to faster growth rate and higher cell yield, the total cost per cell and passage is lowest for LN521 compared to other feeder-free matrices.



#### Biolaminin 111 generates high yield of clinically compliant dopaminergic neurons

Biolaminin 111 (LN111) supports efficient, GMP compliant differentiation of a homogenous population of hPSC-derived dopaminergic (DA) progenitor cells. Compared to embryoid bodies (EB)-based protocols, the yield of DA progenitors is >40x on LN111. Starting from a single 6-well plate of hESCs, DA progenitor cells can be produced on a scale suitable for clinical production.









The cells become TH+ neurons at the site of transplantation in rats

**Yield of VM progentitors d16** Starting from 1\*10<sup>6</sup> cells on day 0 500

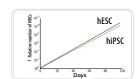


43-fold increase in yield of DA progenitors from human ES cells differentiated on LN111, compared to standard EB-based protocols.

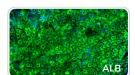


### Biolaminin 521 and 111 support hPSC-derived hepatocyte differentiation and self-organization

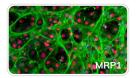
Human ES cells differentiated on Biolaminin 521 (LN521) and 111 (LN111) demonstrate efficient hepatocyte maturation and cell organization with significant improvements in cell function and stability of phenotype. The cells form candicular-like structures express multidrug resistance protein 1 (MRP1) and 2 (MRP2) and are capable of biliary efflux. The cell organization is coherent with the enhanced cellular function



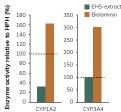
Efficient clonal expansion and maintenance of hESC- and hiPSC-derived hepatoblast-like cells (HBCs).



High ratio of hepatocyte-like cells express albumin (ALB; green).



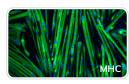
The cells are highly organized and express transporter protein MRP1 (green).



Large increase in P450 metabolic enzyme activity compared to cells on Engelbreth-Holm-Swarm (EHS) mouse hepatocytes (HPH; dotted line).



#### Biolaminin 521 maintains satellite cell-derived myoblast differentiation potential long-term



Biolaminin 521 (LN521) supports superior muscle cell performance in vitro by dramatically improving muscle cell proliferation and differentiation performance, with larger myotubes and higher amounts of nuclei per myotube. Importantly, LN521 supports more consistent and reliable differentiation over long-term culture, and without altering the traditional Pax7/MyoD paradigm.

The cells form myotubes after 8 passages on LN521. Myosin heavy chain expression (MHC; green).

#### **REFERENCES:**

Clonal culturing of human embryonic stem cells on laminin-521/E-cadherin matrix in defined and xeno-free environment. Rodin et al., Nat Commun., 2014 Monolayer culturing and cloning of human pluripotent stem cells on laminin-521 based matrices under xeno-free and chemically defined conditions. Rodin et al., Nat Prot., 2014 hPSCs

Predictive Markers Guide Differentiation to Improve Graft Outcome in Clinical Translation of hESC-Based Therapy for Parkinson's Disease. Kirkeby et al., Cell Stem Cell, 2017 DA neurons - Efficiently Specified VM DA Neurons From Human PSCs Under Xeno-Free Conditions Restore Motor Deficits in Parkinsonian Rodents, Niclis et al., Stem Cells Transl Med., 2017

- Recombinant Laminins Drive the Differentiation and Self-Organization of hESC-Derived Hepatocytes. Cameron et al., Stem Cell Reports, 2015 - Long-Term Self-Renewal of Human ES/iPS-Derived Hepatoblast-like Cells on Human Laminin 111-Coated Dishes. Takayama et al., Stem Cell Reports, 2013 Hepatocytes

- Laminin 521 maintains differentiation potential of mouse and human satellite cell-derived myoblasts during long-term culture expansion. Penton et al., Skeletal Muscle, 2016 Skeletal muscle

Scan to see all applications!





Keep in touch! Email: support@biolamina.com



BioLamina AB, Löfströms Allé 5, SE-172 66 Stockholm, Sweden





