

TISSUE SPECIFIC CELL CULTURE MATRICES

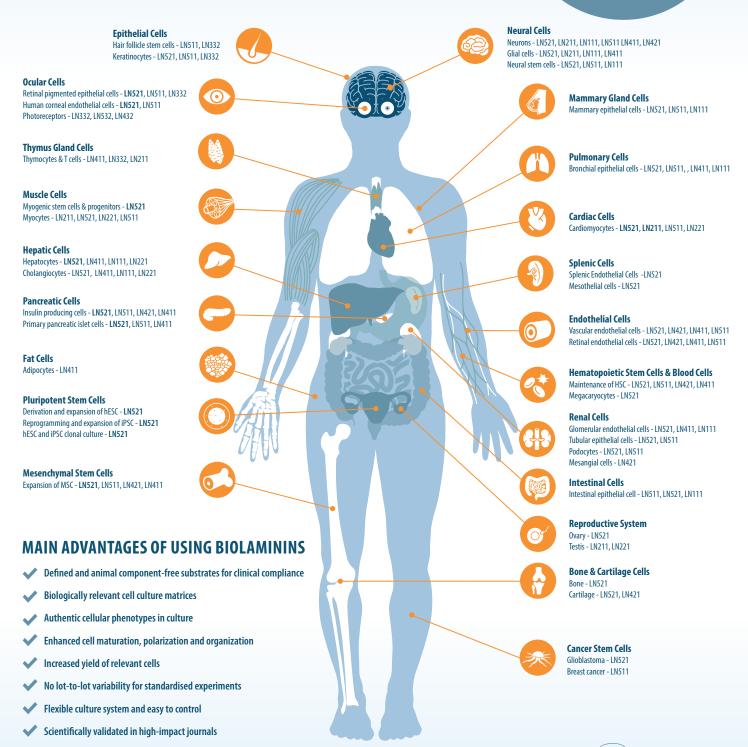
IMITATE THE NATRURAL CELL-MATRIX INTERACTIONS FOR IMPROVED CELL FUNCTIONALITY

BioLamina's chemically defined and animal component-free laminin cell culture matrices, Biolaminin™ matrices, 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



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

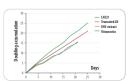
Laminin 521 is a key cell adhesion protein of the natural stem cell niche. The Biolaminin 521 (LN521) substrate supports efficient expansion at low densities of single-cell plated human pluripotent stem cells (hPSCs) under defined and animal component-free conditions. LN521 is compatible with any medium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



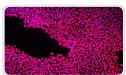
ressed and secreted by hPSCs in the inner cell mass of the embryo.



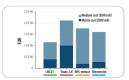
without ROCKi (day 0), grown as a homogenous monolayer (day 3) and can be cultured to high confluence without spontaneous differentiation.



hPSCs propagate faster on LN521 compared to other feeder-free matrices



(Oct4+; pink) and show no areas of differentiation (only DAPI staining;



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



LN111 generates high yield of clinically compliant dopaminergic neurons

Biolaminin 111 (LN111) supports efficient, GMP compliant differentiation of a homogenous population of hPSCderived 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 in a scale suitable for clinical production.

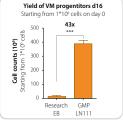








The cells become TH+ neurons at the site of



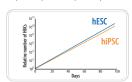
ID: BG-002-04. Valid from 2018-010-16

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

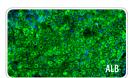


LN521 and LN111 support hPSC derived hepatocyte differentiation and self-organization

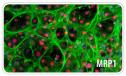
Human ES cells differentiated on Biolaminin 521 (LN521) and 111 (LN111) demonstrate efficient hepatocyte maturation and cell organization with significant improvements in cell function and stability of phenotype. The cells form canalicular-like structures, express multidrug resistance protein 1 (MRP1)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



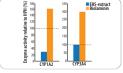
Efficient clonal expansion and maintenance of hESC and hiPSC derived hepatoblast-like cells (HBCs).



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



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

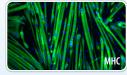


enzyme activity compared to cells on Engelbreth-Holm-Swarm (EHS) mouse sarcoma extract or human primary hepatocytes (HPH; dotted line).



LN521 maintains differentiation potential of satellite cell-derived myoblasts during long-term culture

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

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, 2016 DA neurons:

Neurons From Human Pluripotent Stem Cells Under Xeno-Free Conditions Restore Motor Deficits in Parkinsonian Rodents. Niclis et al., Stem cells transl med., 2016

Recombinant Laminins Drive the Differentiation and Self-Organization of hESC-Derived Hepatocytes. Cameron et al., Stem Cell Reports, 2015 Hepatocytes

Long-Term Self-Renewal of Human ES/iPS-Derived Hepatoblast-like Cells on Human Laminin 111-Coated Dishes, Takayama et al., Cell Stem Cell Reports, 2013

- 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:

KFFP IN TOUCH

TEL: +46-8-5888 5180 EMAIL: SALES@BIOLAMINA.COM BIOLAMINA AB LÖFSTRÖMS ALLÉ 5A 172 66 STOCKHOLM, SWEDEN

For more information and publications visit WWW.THESCIENCEROOM.COM

www.biolamina.com

BIOLAMINA - REVOLUTIONIZING CELL CULTURE



