

SFB 829 post Klenk Minisymposium

NEXT GENERATION FEMALE LEADERS PUSHING EPIDERMAL-DERMAL CROSSTALK

WHERE
CECAD LECTURE HALL
JOSEPH-STELZMANN-STRAßE 26
50931 COLOGNE

WHEN 17TH OCTOBER 2017 2:30 PM - 4:30 PM

PROGRAM

Session 1: Fibroblast function in skin homeostasis and regeneration

14.30 – 15.00 Claire Higgins (London) - *In vitro and in silico models to study skin function*

15.00 – 15.30 Beate Lichtenberger (Wien) - *Fibroblast heterogeneity in skin homeostasis, regeneration and disease*

Session 2: Epithelial function in skin homeostasis and regeneration

15.30 – 16.00 Ning Xu Landén (Stockholm) - The role of regulatory RNAs in wound healing

16.00 – 16.30 Maria Alcolea (Cambridge) - Oesophageal Stem Cell Plasticity - Relevance for Tumour Development





MARIA ALCOLEA

MARIA IS A CELL BIOLOGIST WHO HAS FOCUSED ON UN-DERSTANDING THE BEHAVIOUR OF EPITHELIAL STEM AND PROGENITOR CELLS IN HEALTH AND DISEASE. USING THE MOUSE OESOPHAGUS AS A MODEL, AND BY COMBINING GENETIC LINEAGE TRACING APPROACHES WITH METHODS FROM STATISTICAL PHYSICS, MARIA'S WORK REVEALED THE REMARKABLE CELLULAR PLASTICITY OF EPITHELIAL CELLS IN THIS TISSUE. CELLS ARE NOT ONLY ABLE TO REDEFINE THEIR PROGRAMME OF CELL BEHAVIOUR IN RESPONSE TO INJURY AND EARLY TUMOUR FORMATION, BUT THEY ALSO ADAPT TO THESE PERTURBATIONS BY REACHING A NEW STEADY STATE THAT ENSURES TISSUE MAINTENANCE. INVESTIGAT-ING THE CELLULAR AND MOLECULAR MECHANISMS GOV-ERNING THIS DYNAMIC CELL BEHAVIOUR, AND THE POTEN-TIAL IMPLICATIONS FOR EARLY CANCER DEVELOPMENT REPRESENT MARIA'S MAIN INTERESTS.



NING XU LANDÉN

AFTER OBTAINING A PHD DEGREE IN MOLECULAR VIROLO-GY AT UPPSALA UNIVERSITY SWEDEN, NING XU LANDÉN JOINED PROF. MONA STÄHLE'S LAB AT KAROLINSKA INSTI-TUTE AS A POSTDOC IN 2009, STUDYING THE ROLES OF MICRORNAS IN THE PATHOGENESIS OF PSORIASIS AND SKIN CANCERS. NING STARTED HER RESEARCH GROUP IN 2014, WITH FOCUS ON REGULATORY RNAS (E.G. MI-CRORNA AND LONG NON-CODING RNAS) IN SKIN WOUND REPARATION. HER GOAL IS TO UNDERSTAND THE FUNCTIONS OF REGULATORY RNAS IN WOUND HEALING TO DEVELOP RNA-BASED THERAPY FOR CHRONIC WOUNDS. SINCE THEN NING'S TEAM HAS CHARACTERIZED THE EXPRESSION PROFILES OF REGULATORY RNAS IN IN-JURED HUMAN SKIN DURING NORMAL WOUND HEALING PROCESS AND IN CHRONIC WOUNDS. THEY HAVE RE-VEALED BIOLOGICAL FUNCTIONS AND UNDERLYING MO-LECULAR MECHANISMS OF SEVERAL MIRNAS AND LONG NON-CODING RNAS IN WOUND HEALING. THEY ALSO EX-PLORE THE THERAPEUTIC POTENTIAL OF MIRNAS USING HUMAN EX VIVO AND MOUSE IN VIVO WOUND MODELS.



BEATE LICHTENBERGER

FOR HER POSTDOC, BEATE LICHTENBERGER JOINED THE LAB OF PROF. FIONA WATT, ONE OF THE WORLD-LEAD-SKIN AND LABS IN STEM BIOLOGY, AT THE UNIVERSITY IN CAMBRIDGE AND LATER AT KING'S COLLEGE LONDON AS A POSTDOCTORAL FELLOW. DURING HER POSTDOCTORAL STUDIES SHE DIS-COVERED THAT SKIN DERMIS COMPRISES TWO FIBRO-BLAST LINEAGES WITH DIFFERENT FUNCTIONS IN SKIN PHYSIOLOGY AND PATHOLOGY, AND INVESTIGATED WHICH PATHWAYS ARE INVOLVED IN THE NICHE SIGNAL-LING OF EPIDERMAL STEM CELLS AND FIBROBLASTS SINCE APRIL 2016 SHE IS A PRINCIPAL INVESTIGATOR AT THE SKIN & ENDOTHELIUM RESEARCH DIVISION, THE DEPARTMENT OF DERMATOLOGY AT THE MEDICAL UNI-VERSITY OF VIENNA. HER RESEARCH FOCUSES ON DIS-SECTING THE ROLE OF DISTINCT FIBROBLAST SUBPOPU-LATIONS IN NON-MELANOMA AND MELANOMA SKIN CANCER AS WELL AS IN FIBROTIC SKIN DISEASES AND ON INVESTIGATING IF MODULATING DERMAL SIGNALLING AFFECTS SKIN REGENERATION AND CANCER PROGRES-SION.

CLAIRE HIGGINS

CLAIRE JOINED THE DEPARTMENT OF BIOENGINEERING AT IMPERIAL COLLEGE LONDON IN 2014, AND HAS AS-SEMBLED A RESEARCH LAB WHICH FOCUSES ON FIBRO-BLAST FUNCTION, TISSUE REPAIR AND REGENERATION SPECIFICALLY IN THE CONTEXT OF HUMAN SKIN, LAB IN-TERESTS INCLUDE DEVELOPING IN VITRO MODELS TO ASSESS THE ROLE OF DERMAL FIBROBLAST SUB-TYPES IN WOUND REPAIR AND HAIR GROWTH. COMPARATIVELY, IN SILICO MODELS ARE BEING DEVELOPED AND USED TO INVESTIGATE THE STRUCTURAL FEATURES OF SKIN WHICH ENABLE SITES SUCH AS THE FEET TO HAVE LOAD BEARING CAPACITY. ULTIMATELY THIS RESEARCH IS BEING CONDUCTED WITH THE GOAL OF TRANSLATING THE FINDINGS TO ENABLE OTHER BODY SITES TO BECOME CAPABLE OF BEARING LOAD.