EXECUTIVE BOARD



SPOKESPERSON

Prof. Dr. Ulrich S. Schubert Laboratory of Organic and Macromolecular Chemistry (IOMC) Friedrich Schiller University Jena Humboldtstr. 10 07743 Jena



DEPUTY SPOKESPERSON

Prof. Dr. Michael Bauer Center for Sepsis Control and Care (CSCC) Jena University Hospital Am Klinikum 1 07747 Jena



SCIENTIFIC MANAGER

apl. Prof. Dr. Michael Gottschaldt Laboratory of Organic and Macromolecular Chemistry (IOMC) Friedrich Schiller University Jena Humboldtstr. 10 07743 Jena



PD Dr. Stephanie Höppener Jena Center for Soft Matter (JCSM) Friedrich Schiller University Jena Philosophenweg 7 07743 Jena



Prof. Dr. Oliver Werz Institute of Pharmacy (IP) Friedrich Schiller University Jena Philosophenweg 14 07743 Jena



FRIEDRICH-SCHILLERUNIVERSITÄT JENA Collaborative Research Center CRC 1278

CONTACT

Friedrich Schiller University Jena Jena Center for Soft Matter (JCSM) Prof. Dr. Ulrich S. Schubert apl. Prof. Dr. Michael Gottschaldt Philosophenweg 7 07743 Jena

Phone: +49 3641 9-48201 Fax: +49 3641 9-48202 E-Mail: polytarget@uni-jena.de

Published by Jena Center for Soft Matter (JCSM)
Photos: Jan-Peter Kasper, Annegret Günther,
Klinisches Medienzentrum Universitätsklinikum Jena
Lavout: Stabsstelle Kommunikation



www.jcsm.uni-jena.de



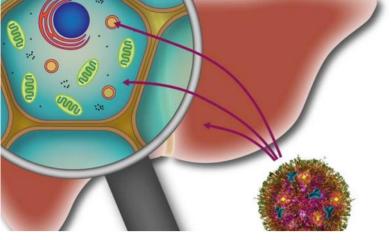


POLYTARGET

Polymer-based Nanoparticle Libraries for Targeted Anti-inflammatory Strategies



www.polytarget.uni-jena.de



THE CRC POLYTARGET

The goal of the CRC PolyTarget is the development of new strategies for the treatment of infection-triggered inflammatory states, centered on a rational design of tailor-made nanoparticulate drug carriers. Pharmacologically active nanoparticles based on functional synthetic polymers and (modified) biopolymers are utilized and characterized to address the fundamental questions of targeted nanomedicine from the bottom up. Based on the establishment of polymer libraries and a detailed molecular and morphological characterization of the nanoparticles, structure-property relationships are studied to optimize the nanoparticles with respect to their biological and pharmaceutical function.

Systematic polymer and particle libraries

Multiple, advanced characterization methods combined with detailed biological studies, GMP laboratory

Elucidation of quantitative structure-property relationships

Transition from trial & error experimentation towards knowledge-based design of multifunctional polymer-based nanoparticles

Cell and organ specific delivery systems for inflammation-related diseases

PROJECTS

Subject Area A: CORE

- A01 Tailor-made multifunctional polymers and nanoparticles with optimized compatibility between biodegradable core and encapsulated drug
- A02 Multifunctional nanoparticles based on polysaccharides for targeted drug delivery with two-step release behavior
- A03 Photoacids and bases as responsive elements in block copolymer nanostructures for uptake and transport
- A04 Spatial and temporal targeting of membranebound mPGES-1 and FLAP / 5-LO by dual inhibitors employing polymer-based nanocarriers
- A05 Targetable nanoparticles for efficient translocation across gastrointestinal barriers
- A06 Controlling stealth and barrier breaking behavior: Hybrid protein nanofibers and POxylation on polymeric nanoparticles with structurally tailored thermal properties

Subject Area B: SHELL

- B01 Targeted nanoparticle mediated delivery of nucleic acids into muscle stem cells for prevention of critical illness myopathy
- B02 Macromolecular prodrug nanoparticles for antimicrobial therapy
- B03 Bioinspired guanidinium-containing nanoparticles for gene delivery
- B04 Nanoscale monitoring of surface effects, structural changes, and encapsulation in block copolymer nanostructures using tip-enhanced Raman spectroscopy

Subject Area C: MEDIUM

C01 Biophotonic characterisation of the interaction of nanoparticles and drugs with hepatic stellate cells

- CO2 Tailored delivery of anti-inflammatory natural drugs using polymer-based nanocarriers to prevent cytokine and eicosanoid storms in infectious inflammation
- CO4 Investigation of cellular response to nanoparticle uptake by dual TEM and superresolution fluorescence imaging
- C06 Prevention of late phase liver damage by targeted modulation of the liver's immune response

Subject Area D: VIRAL

- D01 Mimicking viral entry mechanisms with polymeric nanoparticles
- D02 Delivery platforms for antiviral and antiinflammatory agents targeting infections of respiratory viruses with pandemic potential

Subject Area T: TRANSFER

T01 Targeting renal phosphoinositide 3-kinase γ (PI3Kγ) by dye-tagged nanoparticles

Subject area Ö: Educational institutions and the public

Ö01 Science education on target: Didactic reconstruction of current research at the interface between nanotechnology and medicine

Subject area Z: Central research platform and supporting projects

- Z01 Research platform for the synthesis, formulation, and advanced physicochemical characterization of polymers and nanoparticles
- Z02 Integrated Research Training Group
- Z03 Central tasks of the Collaborative Research Center