



[WWW.4GPCRNET.DE](http://WWW.4GPCRNET.DE)

# 4GPCRnet

International Symposium



# PROGRAM

**SEPTEMBER 26–29, 2022**

**Leipzig, Germany**

NEUES AUGUSTEUM  
LEIPZIG UNIVERSITY  
CAMPUS AUGUSTUSPLATZ  
04103 LEIPZIG, GERMANY

# 4GPCRnet

*International Symposium*

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Dear friends and colleagues,

It is with joyous hearts that we welcome you all to the beautiful and historic city of Leipzig for this celebratory 4GPCRnet International Symposium. This meeting takes place in the spirit of unity: our four Germany-based GPCR-focused research networks have joined forces to reunite the international community after two long pandemic years. Over the next four days, we will together experience the latest findings and approaches to GPCR signal transduction research spanning a broad range of perspectives from the molecular, cellular and physiological levels. The inclusive program highlights diverse and emerging topics in our field that will inspire us all. The Symposium aims to promote international and interdisciplinary cooperation by fostering an open synergistic environment. Among the 450 participants of the meeting are many of our field's most renowned researchers and also - most importantly - many early career investigators who we hope will be invigorated with new ideas and networking opportunities.

We wish you all an enjoyable experience in Leipzig!

Sincerely yours,



**Annette G.  
Beck-Sickinger**  
*Leipzig University,  
Chair of SFB 1423*



**Evi Kostenis**  
*University of Bonn,  
Chair of FOR2372*



**Andreas Bock**  
*Leipzig University & Max  
Delbrück Center of Molecular  
Medicine, Grant Holder  
of CA 18133 ERNEST*



**Ines Liebscher**  
*Leipzig University, Grant  
Holder of CA 18240  
Adher'n Rise*



**Simone Prömel**  
*Heinrich Heine University  
Düsseldorf, Chair of  
CA 18240 Adher'n Rise*



**Martha Sommer**  
*ISAR Bioscience, Chair  
of CA 18133 ERNEST*

## CONFERENCE VENUE

**Leipzig University Neues Augusteum**  
Campus Augustusplatz  
04103 Leipzig, Germany

## CONFERENCE OFFICE

**event lab. GmbH**  
Richard-Lehmann-Str. 12  
04275 Leipzig  
[gpcr-symposium@eventlab.org](mailto:gpcr-symposium@eventlab.org)

## REGISTRATION DESK

**Location:** Ground floor

### Opening hours:

Monday, 26 September 2022	12:00 – 20:00
Tuesday, 27 September 2022	08:00 – 17:30
Wednesday, 28 September 2022	08:00 – 20:00
Thursday, 29 September 2022	08:00 – 18:00

## NAME TAGS

Official conference name tags will be required for admission to all conference functions and lectures and to the exhibition and poster area.

## SPEAKERS PREVIEW

**Location:** Room 122, First floor

## POSTER EXHIBITION

**Location:** in the Paulinum, Ground floor

All posters must be removed by 15:00  
Thursday or will be disposed

### CLOAKROOM

**Location:** Ground floor at the main entrance

**Opening hours:**

Monday, 26 September 2022 08:30 – 21:00

Tuesday, 27 September 2022 08:30 – 17:45

Wednesday, 28 September 2022 08:30 – 21:00

Thursday, 29 September 2022 08:30 – 18:15

Storage room for your coats and baggage.

### COVID NOTICE

We would like to ask you, to please pay attention to your own health and the health of others.

The following hygiene measures apply:

- While indoors, it is strongly recommended to wear a mouth-nose protection and make sure you keep a sufficient distance.
- In case of symptoms, please do not participate in the scientific- or the social program. Please test yourself regularly before entering the venue.
- The organiser will not provide masks and tests.

### WIFI ACCESS

Wireless internet access will be available in most areas.

Network: eduinfo

event name: 4GPCRnet-Symposium

Password: d48de-2022

### TAXIS

Taxis are available around-the-clock at the taxi stand at the Goethestraße or at the Main Station.



In the Collaborative Research Centre 1423 “Structural Dynamics of GPCR Activation and Signalling”, scientists from the life sciences, medicine, pharmacy and bioinformatics of Leipzig University are working together with partners at the Charité Medical University and Max Delbrück Center in Berlin, as well as the Martin Luther University Halle-Wittenberg. The aim of this German Research Foundation-funded initiative is

to understand two understudied classes of GPCRs, the peptide receptors and the adhesion receptors, specifically the structural dynamics of ligand binding, signal transduction, and downstream control of G protein- and arrestin-signalling pathways. The goal of the CRC is to clarify the dynamic structural states of these GPCRs in order to understand their functions and thereby facilitate development of novel therapeutics.

Website: <https://research.uni-leipzig.de/sfb1423/>



The European Research Network on Signal Transduction (ERNEST) is an Action funded by COST (European Cooperation in Science and Technology, [www.cost.eu](http://www.cost.eu)). The main scientific objective of the Action is to develop a common, comprehensive and holistic understanding of signal transduction that will advance development of pathway-specific chemical modulators. ERNEST comprises

a diverse multidisciplinary network of nearly 800 researchers worldwide. Diversity is manifest not only in core expertise, but also career level, gender, country of residence, and institutional sector. Since 2019, ERNEST has served the community with bi-annual meetings, training schools and ECI-organised initiatives, as well as financial support of cross-border scientific exchanges, open access publications, and reliable communication channels that support networking and cooperation.

Website: <https://ernest-gpcr.eu/>



Based at the University of Bonn, the Research Unit 2372 “G-protein signalling cascades: with new molecular probes and agents to new pharmacological concepts” is funded by the German Research Foundation and involves partners at the BMRZ-Goethe University in Frankfurt, Otto-von-Guericke-University Magdeburg, and the Paul Scherrer Institute in Switzerland. This multi-disciplinary consortium aims at the

rational design and the generation of novel, cell-permeable signalling inhibitors with selectivity for G protein families. Molecular-mechanistic analyses of these inhibitors provide insights into the relevance of individual signalling cascades within complex signalling networks and (patho-)physiological events. Since GPCRs serve as molecular targets for a plethora of drugs, novel principles for piloting intracellular signalling might also inspire the development of therapies addressing GPCRs themselves.

Website: <https://www.for2372.uni-bonn.de/>



Adher 'N Rise (ADHEsion GPCR Network: Research and Implementation Set the path for future Exploration) is an Action funded by COST (European Cooperation in Science and Technology, [www.cost.eu](http://www.cost.eu)). Its aim is to promote, stimulate and translate research on Adhesion-G

protein-coupled receptors (aGPCRs) ‘from bench to bedside’ in Europe. Scientists as well as clinicians with divergent expertise and interests begin to recognise the biological importance of aGPCRs and their unexploited pharmacological potential. The Action supports this relatively young and growing community of aGPCR researchers by providing communication platforms and opportunities to interact. Early Career Investigators (ECIs) are of utmost importance to Adher 'N Rise as they ensure the development of novel ideas and the long-term progress of the field.

Website: <https://www.adhernrise.eu/>

## **Brian Kobilka, MD**

Professor, Department of Molecular and Cellular Physiology

Hélène Irwin Fagan Chair in Cardiology

Stanford University



Brian Kobilka received Bachelor of Science Degrees in Biology and Chemistry from the University of Minnesota, Duluth in 1977. He graduated from Yale University School of Medicine in 1981, and completed residency training in Internal Medicine at the Barnes Hospital, Washington University School of Medicine, St. Louis, Missouri in 1984. From 1984-1989 he was a postdoctoral fellow in the laboratory of Robert Lefkowitz at Duke University.

While in the Lefkowitz lab, he and his colleagues cloned the gene that encodes the receptor for the hormone adrenaline. They found that the receptor was similar to rhodopsin, the light sensing receptor. It was later discovered that there is an entire family of receptors that look and act in similar ways. These receptors are known as G-protein-coupled receptors (GPCRs); they are responsible for the body's response to the majority of hormones and neurotransmitters.

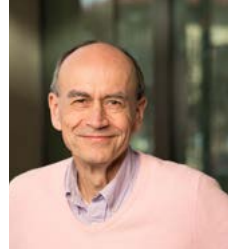
In 1989 he joined the faculty of Medicine and Molecular and Cellular Physiology at Stanford University. Research in the Kobilka lab focuses on the structure and mechanism of action of GPCRs. They apply a spectrum of biochemical, biophysical and structural approaches to understand GPCR signalling at the molecular level. He is a member of the National Academy of Sciences, the National Academy of Medicine, and the American Academy of Arts and Sciences. In 2012, Kobilka was awarded the Nobel Prize in Chemistry with Robert Lefkowitz for their work on GPCRs.



**Thomas C. Südhof, M.D.**

Avram Goldstein Chair,

Professor of Molecular and Cellular Physiology, and  
Investigator of the Howard Hughes Medical Institute,  
Stanford University School of Medicine



Thomas Christian Südhof, M.D., serves as the Avram Goldstein Professor in the School of Medicine of Stanford University, as a Professor of Molecular & Cellular Physiology, Neurosurgery, and by courtesy, of Neurology and Psychiatry, and as an Investigator of the Howard Hughes Medical Institute. Südhof's research interests focus on how synapses that transmit and process information in neural circuits are formed and specified, and how their role becomes dysfunctional in neuropsychiatric and neurodegenerative diseases. Towards this goal, Südhof uses genetic manipulations in human and mouse neurons as well as biophysical and cell-biological approaches to dissect the contributions of trans-synaptic signaling complexes to the construction of synapses and their properties and plasticity.

**M. Madan Babu | St. Jude Children's Research Hospital | USA**

Wednesday | 28 September 2022 | 16:45 - 17:15

Variation in GPCR Signaling: Implications for Physiology and Drug Discovery.

Session 5: Computational Design and GPCR Therapeutics

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**Meritxell Canals Buj | The University of Nottingham | UK**

Monday | 26 September 2022 | 14:30 - 15:00

Mu-opioid receptor induced changes in beta-arrestin2 conformations

Session 1: Modulations of GPCR Signal Transduction

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**Andy Chevigné | Luxembourg Institute of Health | Luxembourg**

Tuesday | 27 September 2022 | 11:30 - 12:00

Dual-activity GPCR: ACKR3 at the crossroads between chemokine and opioid peptide regulation

Session 2: Holistic Concepts in GPCR Signal Transduction

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**I. Sadaf Farooqi | University of Cambridge | UK**

Wednesday | 28 September 2022 | 14:30 - 15:00

GPCRs and metabolic disease

Session 4: GPCR in Physiology and Disease

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**Michael Freissmuth | Medical University Wien | Austria**

Wednesday | 28 September 2022 | 14:00 - 14:30

Targeting GPCRs to enhance haematopoietic stem cell transplantation

Session 4: GPCR in Physiology and Disease



**Joshua Frenster | Universitat Pompeu Fabra | Spain**

Tuesday | 27 September 2022 | 14:30-15:00

GPR133 (ADGRD1) signaling is increased by the dissociation of its extracellular NTF and by binding of its new positive allosteric modulator PTK7

Session 3: aGPCR: from Structure to Function

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**Heidi Hamm | Vanderbilt University | USA**

Tuesday | 27 September 2022 | 9:00 - 9:30

Role of GPCRs in regulation of hormone and neurotransmitter release.

Session 2: Holistic Concepts in GPCR Signal Transduction

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**Adriaan Ijzermann | Leiden University | Netherlands**

Wednesday | 28 September 2022 | 16:15 - 16:45

Partial agonism in the picture

Session 5: Computational Design and GPCR Therapeutics

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**Brian Kobilka | Stanford University | USA**

Thursday | 29 September 2022 | 16:15 - 17:30

The role of protein dynamics in GPCR signaling

Keynote Lecture

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**Mickey Kosloff | University of Haifa | Israel**

Tuesday | 27 September 2022 | 11:00 - 11:30

Deciphering structural design principles for interaction specificity in G protein signaling

Session 2: Holistic Concepts in GPCR Signal Transduction

**Josef Lazar | Institute of Organic Chemistry and Biochemistry  
of the Czech Academy of Sciences | Czech Republic**

Monday | 26 September 2022 | 16:30 - 17:00

A Simple and Obvious (in Retrospect) Method for  
Microscopy Imaging of G-protein Signaling

Session 1: Modulations of GPCR Signal Transduction

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**Kelly Monk | The Vollum Institute  
Oregon Health & Science University | USA**

Tuesday | 27 September 2022 | 16:15 - 16:45

In vivo investigations of adhesion GPCR function

Session 3: aGPCR: from Structure to Function

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**Katarina Nikolic | Belgrade University | Serbia**

Thursday | 29 September 2022 | 9:00 - 9:30

Application of computational methods in rational design of multi-target  
ligands as potential antipsychotics and antidepressants

Session 5: Computational Design and GPCR Therapeutics

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**Mette Rosenkilde | University of Copenhagen | Denmark**

Wednesday | 28 September 2022 | 11:15 - 11:45

How come that GIPR agonists and antagonists both result in weight loss?

Session 4: GPCR in Physiology and Disease



**Bryan Roth | University of North Carolina UNC | USA**

Thursday | 29 September 2022 | 9:30 - 10:00

Integrated computational and structural approaches for GPCR drug discovery

Session 5: Computational Design and GPCR Therapeutics

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**Gunnar Schulte | Karolinska Institute | Sweden**

Monday | 26 September 2022 | 17:00 - 17:30

Mechanisms of Class F receptor activation and signal specification

Session 1: Modulations of GPCR Signal Transduction

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**Arun K. Shukla | Indian Institute of Technology | India**

Thursday | 29 September 2022 | 14:00 - 14:30

Structural insights into GPCR-beta-arrestin interaction and signaling

Session 6: Molecular Structure and Dynamics

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**Thomas C. Südhof | Stanford University School of Medicine | USA**

Wednesday | 28 September 2022 | 9:00 - 10:15

Adhesion-GPCR signaling in synapse formation

Keynote Lecture

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**Jin-Peng Sun | Shandong University | China**

Tuesday | 27 September 2022 | 14:00 - 14:30

Ligands identification and Structural studies of several aGPCRs

Session 3: aGPCR: from Structure to Function

**Roger Sunahara | University of California San Diego | USA**

Tuesday | 27 September 2022 | 9:30 - 10:00

GPCR-mediated endosomal ERK signaling through  
G proteins: implications to disease

Session 2: Holistic Concepts in GPCR Signal Transduction

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**Christopher A. Tate | MRC Laboratory of Molecular Biology | UK**

Thursday | 29 September 2022 | 11:30 - 12:00

Structural insights into allostery in GPCRs: G proteins, arrestin and dimers

Session 6: Molecular Structure and Dynamics

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**Mark von Zastrow | University of California | USA**

Monday | 26 September 2022 | 15:00 - 15:30

Toward understanding the subcellular organization  
and reorganization of GPCR signaling

Session 1: Modulations of GPCR Signal Transduction

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**Nina Wettschureck | Max Planck Institute for Heart and Lung Research**

**Department of Pharmacology | Germany**

Wednesday | 28 September 2022 | 10:45 - 11:15

Orphan GPCRs in the vascular and immune system

Session 4: GPCR in Physiology and Disease



**Denise Wootten | Monash University**  
**Monash Institute of Pharmaceutical Sciences | Australia**

Thursday | 29 September 2022 | 14:30 - 15:00

Structural and Mechanistic insights into class B1  
GPCR signalling and allostery (Cryo-EM)

Session 6: Molecular Structure and Dynamics

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**Beili Wu | Chinese Academy of Sciences | China**

Thursday | 29 September 2022 | 11:00 - 11:30

Structural insights into signal transduction of metabotropic glutamate receptors

Session 6: Molecular Structure and Dynamics of GPCRs

09:00	<b>COST ACTION Meetings</b>	
11:30	<b>COST ACTION Board Meeting</b>	
13:30	<b>Welcome Ceremony</b>	
14:15	<b>Break</b>	
14:30	<p><b>Session 1</b></p> <hr/> <p><b>Modulation of GPCR Signal Transduction</b></p> <p><i>Chair: Evi Kostenis</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Mu-opioid receptor induced changes in beta-arrestin2 conformations</b> Meritxell Canals Buj, The University of Nottingham, UK</p> <p><b>Toward understanding the subcellular organization and reorganization of GPCR signaling</b> Mark von Zastrow, University of California, USA</p> <p><b>Chemogenetic G protein-ligand pairs for causal investigation of cellular biology in vitro and in vivo</b> Judith Ahlenfelder, University of Bonn, GER</p> <p><b>Cotranslational insertion of GPCRs into nanoparticle membranes and their structural evaluation by cryo-electron microscopy in complex with G-proteins</b> Zoe Köck, Goethe University Frankfurt, GER</p>
16:00	<b>Coffee Break</b>	
16:30	<p><b>Session 1</b></p> <hr/> <p><b>Modulation of GPCR Signal Transduction</b></p> <p><i>Chair: Torsten Schöneberg</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>A Simple and Obvious (in Retrospect) Method for Microscopy Imaging of G-protein Signaling</b> Josef Lazar, Czech Academy of Sciences, CZ</p> <p><b>Mechanisms of Class F receptor activation and signal specification</b> Gunnar Schulte, Karolinska Institute, SWE</p> <p><b>In vitro and in vivo regulation of <math>\beta</math>-Adrenoceptors signaling using synthetic light-regulated molecules</b> Xavier Rovira, The Spanish National Research Council (CSIC), ESP</p> <p><b>Cis-on photoswitchable G protein-biased ligand for melatonin receptors</b> Gloria Somalo-Barranco, Université de Paris, FRA</p> <p><b><math>\beta</math>-arrestin 1 and 2 exhibit distinct phosphorylation-dependent conformations when coupling to the same GPCR in living cells</b> Carsten Hoffmann, University Hospital Jena, GER</p>
18:15 18:45	<p><b>Poster Appetizer Session</b></p> <p><b>Room: Audimax   Chair: Ralf Jockers</b></p>	
19:00 21:00	<p><b>Poster Session with Drinks and Snacks supported by IRN</b></p> <p><b>Room: Paulinum</b></p>	



09:00	<p><b>Session 2</b></p> <hr/> <p><b>Holistic Concepts in GPCR Signal Transduction</b></p> <p><i>Chair: Martha Sommer</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Role of GPCRs in regulation of hormone and neurotransmitter release</b> Heidi Hamm, Vanderbilt University, USA</p> <p><b>GPCR-mediated endosomal ERK signaling through G proteins: implications to disease</b> Roger Sunahara, University of California San Diego, USA</p> <p><b>New nanobodies illuminate opioid receptor signaling with subcellular precision</b> Miriam Stoeber, University of Geneva, CHE</p> <p><b>Molecular mechanism for GPCR spatial organization at the cell surface</b> Gabriele Kockelkoren, University of Copenhagen, DNK</p>
10:30	<b>Coffee Break</b>	
11:00	<p><b>Session 2</b></p> <hr/> <p><b>Holistic Concepts in GPCR Signal Transduction</b></p> <p><i>Chair: Andreas Bock</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Deciphering structural design principles for interaction specificity in G protein signaling</b> Mickey Kosloff, University of Haifa, ISR</p> <p><b>Dual-activity GPCR: ACKR3 at the crossroads between chemokine and opioid peptide regulation</b> Andy Chevigné, Luxembourg Institute of Health, LUX</p> <p><b>Lipid regulation of ligand binding to atypical chemokine receptor 3</b> Martin Gustavsson, University of Copenhagen, DNK</p> <p><b>Unveiling of the elusive signaling pathways activated by the super conserved orphan receptors GPR27, GPR85 and GPR173</b> Julien Hanson, University of Liege, BEL</p>
12:30	<p><b>Poster Appetizer Session</b></p> <p>Room Audimax   Chair: Norbert Sträter</p>	
13:00	<p><b>Lunch   Room: Foyer</b></p> <p><b>and</b></p> <p><b>Poster Session   Room: Paulinum</b></p>	

14:00	<p><b>Session 3</b></p> <hr/> <p><b>aGPCR from Structure to Function</b></p> <p><i>Chair: Ines Liebscher</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Ligands identification and Structural studies of several aGPCRs</b> Jin-Peng Sun, Shandong University, CHN</p> <p><b>GPR133 (ADGRD1) signaling is increased by the dissociation of its extracellular NTF and by binding of its new positive allosteric modulator PTK7</b> Joshua Frenster, Universitat Pompeu Fabra, ESP</p> <p><b>Identifying Novel Ligands for ADGRG5/GPR114 with a new Ultra-large Library Screening Algorithm in Rosetta</b> Jens Meiler, Leipzig University, GER</p> <p><b>A transgenic system to monitor adhesion G protein-coupled receptor dissociation</b> Tobias Langenhan, Leipzig University, GER</p>
15:30	<b>Coffee Break</b>	
16:15	<p><b>Session 3</b></p> <hr/> <p><b>aGPCR from Structure to Function</b></p> <p><i>Chair: Simone Prömel</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>In vivo investigations of adhesion GPCR function</b> Kelly Monk, The Vollum Institute Oregon Health &amp; Science University, USA</p> <p><b>Adhesion GPCRs in oncology: emerging roles, functions and therapeutic directions</b> David Favara, University of Cambridge, UK</p> <p><b>Regulation of glucose homeostasis by Adhesion GPCRs</b> Doreen Thor, Leipzig University, GER</p>
18:30	<b>Transfer to Zoo</b>	
19:00 01:00	<b>Conference dinner at Zoo Leipzig</b>	

09:00	<b>Keynote Lecture</b> <i>Chair: Nicole Scholz</i> <hr/> <b>Room: Audimax</b>	<b>Adhesion-GPCR signaling in synapse formation</b> Thomas C. Südhof, Stanford University School of Medicine, USA
10:15	<b>Coffee Break</b>	
10:45	<b>Session 4</b> <hr/> <b>GPCR in Physiology and Disease</b> <i>Chair: Martin Lohse</i> <hr/> <b>Room: Audimax</b>	<b>Orphan GPCRs in the vascular and immune system</b> Nina Wettschureck, MPI, GER  <b>How come that GIPR agonists and antagonists both result in weight loss ?</b> Mette Rosenkilde, University of Copenhagen, DNK  <b>Oncomodulation explained: cancer going viral</b> Nick D. Bergkamp, Vrije Universiteit Amsterdam, NLD  <b>Structural and Functional Diversity Among Agonist-Bound States of the GLP-1 Receptor</b> Brian P. Cary, University of Wisconsin-Madison, USA  <b>The orphan G protein-coupled receptor, GPR50, and its involvement in obesity and type2 diabetes</b> Julie Dam, Inserm CNRS Paris, FRA
12:30	<b>Poster Appetizer Session</b> Room Audimax   <i>Chair: Peter Schmidt</i>	
13:00	<b>Lunch   Room: Foyer and</b> <b>Poster Session   Room: Paulinum</b>	

14:00	<p><b>Session 4</b></p> <hr/> <p><b>GPCR in Physiology and Disease</b></p> <p><i>Chair: Peter Kühnen</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Targeting GPCRs to enhance haematopoietic stem cell transplantation</b> Michael Freissmuth, Medical University of Vienna, AUT</p> <p><b>GPCRs and metabolic disease</b> I Sadaf Farooqi, University of Cambridge, UK</p> <p><b>A NanoBiT/BRET-based binding assay for CRISPR-Cas9-edited FZD7 allows real-time analysis of ligand binding in the colorectal cancer model</b> Pawel Kozielowicz, Karolinska Institutet, SWE</p> <p><b>G protein-coupled receptor 84 is an evolutionary conserved Gα15-coupled immune cell receptor</b> Claudia Stäubert, Leipzig University, GER</p>
15:30	<b>Coffee Break</b>	
16:15	<p><b>Session 5</b></p> <hr/> <p><b>Computational Design and GPCR Therapeutics</b></p> <p><i>Chair: Andrea Sinz</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Partial agonism in the picture</b> Adriaan Ijzermann, Leiden University, NLD</p> <p><b>Variation in GPCR Signaling: Implications for Physiology and Drug Discovery</b> M Madan Babu, St. Jude Children's Research Hospital, USA</p> <p><b>Structure-Based Computer Driven Discovery of GPCR Ligands with New Chemotypes and Functional Selectivity</b> Vsevolod Katrich, University of Southern California, USA</p> <p><b>Optical control of family A GPCRs using azobenzene-derived photoswitchable ligands</b> Rob Leurs, Vrije Universiteit Amsterdam, NLD</p> <p><b>Caenorhabditis elegans as model organism for GPCR research</b> Anette Kaiser, Leipzig University, GER</p>
18:00	<b>Short Break</b>	
18:15 18:45	<p><b>Greetings &amp; Poster Appetizer Session</b></p> <p><b>Room: Audimax</b>   <i>Chair: Marie Hoffmann (Project Manager Science Funding, Stiftung Charité) &amp; Peter Hildebrand</i></p>	
19:00 21:00	<p><b>Poster Session with Drinks and Snacks</b> <b>- funded by Stiftung Charité</b></p> <p><b>Room: Paulinum</b></p>	

09:00	<p><b>Session 5</b></p> <hr/> <p><b>Computational Design and GPCR Therapeutics</b></p> <p><i>Chair: Peter Stadler</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Application of computational methods in rational design of multi-target ligands as potential antipsychotics and antidepressants</b> Katarina Nikolic, Belgrade University, SRB</p> <p><b>Integrated computational and structural approaches for GPCR drug discovery</b> Bryan Roth, University of North Carolina UNC, USA</p> <p><b>In Silico Hit Discovery for GPCRs across Scale and Sector</b> Brian Bender, Sosei Heptares, UK</p> <p><b>GPCR structure analysis platform and activation mechanisms</b> David Gloriam, University of Copenhagen, DNK</p>
10:30	<b>Coffee Break</b>	
11:00	<p><b>Session 6</b></p> <hr/> <p><b>Molecular Structure and Dynamics</b></p> <p><i>Chair: Patrick Scheerer</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Structural insights into signal transduction of metabotropic glutamate receptors</b> Beili Wu, Chinese Academy of Sciences, CHN</p> <p><b>Structural insights into allostery in GPCRs: G proteins, arrestin and dimers</b> Christopher G Tate, MRC Laboratory of Molecular Biology, UK</p> <p><b>Live-cell structural insights into arrestin interactions with a secretin-like GPCR</b> Irene Coin, Leipzig University, GER</p> <p><b>A modular ligand-directed approach to label endogenous aminergic GPCRs in live cells</b> Xavier Gómez-Santacana, Université de Montpellier, CNRS and INSERM, ESP</p>
12:30	<p><b>Poster Appetizer Session</b></p> <p>Room Audimax   Chair: Paolo Annibale</p>	
13:00	<p><b>Lunch   Room: Foyer and</b></p> <p><b>Poster Session   Room: Paulinum</b></p>	

14:00	<p><b>Session 6</b></p> <hr/> <p><b>Molecular Structure and Dynamics</b> <i>Chair: Daniel Huster</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>Structural insights into GPCR-beta-arrestin interaction and signaling</b> Arun A Shukla, Indian Institute of Technology, IND</p> <p><b>Structural and Mechanistic insights into class B1 GPCR signalling and allostery (Cryo-EM)</b> Denise Wootten, Monash University, AUS</p> <p><b>Allosteric modulation of metabotropic glutamate receptor activation resolved by single molecule FRET using minimally invasive labeling strategies</b> Robert B. Quast, Centre de Biologie Structurale (CBS), FRA</p> <p><b>Structural and functional studies of CC chemokine receptor 5 phosphorylation and its interaction with arrestin2</b> Polina Isaikina, University of Basel, CHE</p>
15:45	<b>Coffee Break</b>	
16:00	<p><b>Keynote Lecture</b> <i>Chair: Peter Hildebrand</i></p> <hr/> <p><b>Room: Audimax</b></p>	<p><b>The role of protein dynamics in GPCR signaling</b> Brian Kobilka, Stanford University, USA</p>
17:30	<p><b>CRC Closing Remarks &amp; Farewell</b> Annette G. Beck-Sickinger Room: Audimax</p>	
17:45 22:00	<p><b>Closing Party</b> <b>Early Career Investigator Session and Awarding</b> at Moritzbastei</p>	

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We are very pleased with the high number of scientific contributions submitted.

The posters can be viewed in the Paulinum during regular opening hours.

During the five Poster Sessions, especially young investigators have the opportunity to present their work to an international audience. The schedule of the poster sessions is as follows:

The poster authors have been given the opportunity to present their poster at the beginning of each poster session, each with 1-2 slides in the form of a short talk – so called Poster Appetizer.

**The following posters will be presented during the Poster Appetizer Sessions:**

**Monday**

P-I-07, P-I-09, P-I-10, P-I-11, P-I-13, P-I-19, P-I-29, P-I-37, P-I-40, P-I-77, P-I-80

**Tuesday**

P-I-21, P-I-26, P-I-33, P-I-50, P-I-54, P-I-57, P-I-68, P-I-71

**Wednesday - Lunch**

P-I-55, P-II-02, P-II-05, P-II-06, P-IV-02, P-IV-06, P-IV-18, P-IV-29

**Wednesday - Evening**

P-V-01, P-V-02, P-V-04, P-V-11, P-V-14, P-V-17, P-VI-01, P-VI-02, P-VI-07

**Thursday**

P-VI-18, P-VI-31, P-VI-38, P-VI-39, P-VI-40, P-VI-46



### Poster Session Monday – supported by IRN



18:15 – 18:45 Poster Appetizer Session, Audimax

19:00 – 20:00 poster viewing of the poster topics P-I – P-III

20:00 – 21:00 poster viewing of the poster topics P-IV – P-VI

Food and Drinks will be provided

### Poster Session Wednesday – funded by Stiftung Charité



18:15 – 18:45 Poster Appetizer Session, Audimax  
Introduction and greetings: Marie Hoffmann,  
Project Manager Science Funding, Stiftung Charité

19:00 – 20:00 poster viewing of the poster topics P-IV – P-VI

20:00 – 21:00 poster viewing of the poster topics P-I – P-III

Food and Drinks will be provided

A selected poster jury will evaluate all posters. During the Closing Party at Mortizbastei on Thursday, the best posters will be honoured with a prize.

Be there and support the poster authors!

## P-I – Modulation of GPCR Signal Transduction

<b>P-I-01</b>	Voltage sensitivity of G protein coupled receptors	<i>Y. Ben-Chaim</i>
<b>P-I-02</b>	Nanodelivery: A new and efficient method for transfer and characterization of cell-free synthesized G-protein coupled receptors into living cells	<i>S. Umbach et al.</i>
<b>P-I-03</b>	Constitutive vs. agonist-induced internalization of mGlu receptors	<i>M. Cimadevila et al.</i>
<b>P-I-05</b>	Allosteric modulation of GPCR-induced $\beta$ -arrestin trafficking and signaling by a synthetic intrabody	<i>M. Chaturvedi et al.</i>
<b>P-I-06</b>	Pathway selectivity in Frizzleds is achieved by conserved microswitches defining pathway-determining, active conformations	<i>L. Grätz et al.</i>
<b>P-I-07</b>	The impact of lipid polyunsaturation in the Dopamine D2 activation and signaling	<i>I. D. Alves</i>
<b>P-I-08</b>	Initiation and regulation of glucagon receptor signaling by glucagon, stable glucagon analogs, and RAMPs	<i>J. M. Mathiesen et al.</i>
<b>P-I-09</b>	Heteromerization with the cell adhesion molecule CD44 modulates the constitutive activity and agonist-induced signaling of the serotonin receptor 7	<i>S. Borsdorf et al.</i>
<b>P-I-10</b>	Different Effects of Ageing on the Expression of Bitter Taste Receptors in Human and Rat Colon	<i>F. Jalševac et al.</i>
<b>P-I-11</b>	New insights into the non-canonical desensitization of Gq-signaling by GRK2/3 expression levels	<i>N. Jaiswal et al.</i>
<b>P-I-12</b>	Pharmacological characterization of new Angiotensin II receptors fluorescent ligands as tracer for in vitro and in vivo labeling	<i>C. Delaitre et al.</i>
<b>P-I-13</b>	Dissecting the GPCR signaling cascade one step at the time: a fluorescence spectroscopy investigation of adrenergic-mediated signaling upon cell swelling	<i>A. Sirbu et al.</i>
<b>P-I-14</b>	Location-resolved signaling profiles of opioid receptors	<i>A. Radoux-Mergault et al.</i>
<b>P-I-15</b>	Development of a $\beta$ -Arrestin2-biased Photoswitchable Cannabinoid 2 Receptor Agonist	<i>S. A.M. Steinmüller et al.</i>
<b>P-I-16</b>	Tuning GPCR expression, stability and allostery by directed evolution in mammalian cells	<i>C. Klenk et al.</i>
<b>P-I-17</b>	Shedding light on the internalisation mechanism of glucagon-like peptide 1 receptor	<i>E.V. Moo et al.</i>

<b>P-I-18</b>	Single cell studies reveal distinct pharmacological profiles of GPCRs depending on their oligomerization state	<i>K. L. Martinez et al.</i>
<b>P-I-19</b>	Functional modulation of PTH1R activation and signalling by RAMP2	<i>K. Nemec et al.</i>
<b>P-I-20</b>	The Electrostatic Switch: a new model for CCR5 desensitization based on dynamic modulation of negative charges at the receptor-ligand interface	<i>I. Pinheiro et al.</i>
<b>P-I-21</b>	The influence of succinate receptor 1 on cellular metabolism and its subcellular localization	<i>A.-D. Liebing et al.</i>
<b>P-I-22</b>	The orphan receptor GPR101 as a novel cause of growth hormone deregulation	<i>C. Abboud et al.</i>
<b>P-I-23</b>	Kinetic analysis of endogenous $\beta$ 2-adrenoceptor-mediated cAMP GloSensor <sup>TM</sup> responses in HEK293 cells.	<i>S. A. Cullum et al.</i>
<b>P-I-24</b>	WNT stimulation induce conformational dynamics of Frizzled-Dishevelled interaction	<i>C.-F. Bowin et al.</i>
<b>P-I-25</b>	Pharmacological characterization of human calcium-sensing receptor mutants	<i>W. Du et al.</i>
<b>P-I-26</b>	The role of the intracellular loop 3 of M2 and M4 muscarinic receptors in GRK-mediated $\beta$ -arrestin recruitment and internalization	<i>J. C. Filor et al.</i>
<b>P-I-27</b>	The two endogenous ACKR3 agonists exhibit distinct effects on $\beta$ -arrestin 2	<i>N. Youssef et al.</i>
<b>P-I-28</b>	Gs priming of non-Gs protein activation and signalling at the glucagon-like peptide-1 receptor	<i>D. Wootten et al.</i>
<b>P-I-29</b>	Negative Allosteric Modulation of the Human Neuropeptide Y4 Receptor by the Small Molecule (S)-VU0637120	<i>C. Schüß et al.</i>
<b>P-I-30</b>	Allosteric Modulation of Peptide GPCRs	<i>H. Lentschat et al.</i>
<b>P-I-31</b>	Pharmacological characterisation of positive allosteric modulators for the delta opioid receptors to treat gastrointestinal motility disorders	<i>S. Alvi et al.</i>
<b>P-I-32</b>	Context-dependent adaption of vascular smooth muscle cell G-protein signaling by regulator of G-protein signaling 5	<i>J. Garg et al.</i>
<b>P-I-33</b>	Pharmacological characterization of H1R and H3R small molecule photoligands – reaching beyond the UV spectrum	<i>I. Josimovic et al.</i>
<b>P-I-34</b>	Catch me if you can: Where is Gs-derived G $\beta\gamma$ ?	<i>J. Brands et al.</i>
<b>P-I-35</b>	Nanobody Toolkit to Sense and Modulate GPCR Activity	<i>Z. Valbret et al.</i>
<b>P-I-36</b>	NOMAD® biosensors for multiplexing Ca <sup>2+</sup> and $\beta$ -Arrestin functional assays of Proteinase-activated receptor 2 (PAR2)	<i>P. Villacé et al.</i>

<b>P-I-37</b>	Characterisation of the transducer coupling profiles of PAC1 receptor splice isoforms	<i>J. Lu et al.</i>
<b>P-I-38</b>	Improvement of a small molecule hit structure at chemerin receptor	<i>T. Weiß et al.</i>
<b>P-I-39</b>	Introduction and characterization of PSB-18061 - a novel tool compound targeting the orphan primate-specific Mas-related G protein-coupled receptor MRGPRX4	<i>R. Gedschold et al.</i>
<b>P-I-40</b>	Optical control of $\beta$ -adrenergic signalling via light-sensitive ligands	<i>S. Shi et al.</i>
<b>P-I-41</b>	Bivalent ligands targeting dopamine D3 and neurotensin NTS1 receptors	<i>H. Vogt et al.</i>
<b>P-I-42</b>	Exploring the Effects of Dimerization in the NPY System Utilizing a Live Cell Labeling Approach	<i>A. Mohr et al.</i>
<b>P-I-43</b>	$\beta$ -hairpins and the conserved disulfide bridge in the extracellular loop 2 of peptide GPCRs: A contributing factor to pathway specificity	<i>M. Wygas et al.</i>
<b>P-I-44</b>	Profiling small molecule inhibitors for CXCR4 using systems biology modelling	<i>K. S. Pan et al.</i>
<b>P-I-45</b>	GPCR-arrestin complex stability and dynamics probed with atomic force microscopy and molecular dynamics simulations	<i>F. M. Wilhelm et al.</i>
<b>P-I-46</b>	Combining cell-based and computational methods to unravel TAS2R14 novel bitter receptor inhibitors	<i>L. Peri et al.</i>
<b>P-I-47</b>	Peptides as modulators of S1P signaling affect immune cell migration	<i>J. Gattringer et al.</i>
<b>P-I-48</b>	Employing nanobodies to functionally modulate GPCRs	<i>M. A. Skiba et al.</i>
<b>P-I-49</b>	CXCR4 oligomerization: detection, modulation and functional consequences	<i>C. V. Perez Almeria et al.</i>
<b>P-I-50</b>	Photoswitchable small-molecule ligands to optically modulate chemokine receptors	<i>R. Leurs et al.</i>
<b>P-I-51</b>	Inactivation of the ACKR3 receptor by a small-molecule inverse agonist	<i>D. Nesheva et al.</i>
<b>P-I-52</b>	Bridging the binding sites of the cannabinoid 2 receptor with dualsteric ligands	<i>A. Tutov et al.</i>
<b>P-I-53</b>	Photoswitchable Benzimidazole Azo-arenes as $\beta$ -Arrestin2-biased Cannabinoid 2 Receptor-Selective Agonists	<i>S. A.M. Steinmüller et al.</i>
<b>P-I-54</b>	Pharmacological characterization of photocaged histamine H3 and H4 receptor agonists	<i>M. Gao et al.</i>
<b>P-I-55</b>	Identification and functional analysis of a cannabinoid 2 / $\kappa$ -opioid receptor interaction	<i>N. Tomašević et al.</i>

<b>P-I-56</b>	Comparison of human and mouse formyl peptide receptor 2 (FPR2) endocytic trafficking.	<i>C. Jack et al.</i>
<b>P-I-57</b>	Utilising CRISPR/Cas9 techniques for endogenous promotor expression of split Nanoluciferase tagged $\beta 1/2$ adrenoceptors and miniGs proteins in HEK293T cells	<i>L. J. Humphrys et al.</i>
<b>P-I-58</b>	Protein kinase A and GRKs differentially influence direct interaction of $\beta$ -arrestin2 with the b1AR, $\beta$ -arrestin2 translocation to the plasma membrane, and b1AR internalization	<i>L. E. Kletzin et al.</i>
<b>P-I-59</b>	Imaging GPCR signaling by polarization-resolved fluorescence microscopy	<i>V. Marková et al.</i>
<b>P-I-60</b>	A bead-based GPCR phosphorylation immunoassay for high-throughput ligand profiling and GRK inhibitor screening	<i>S. Schulz et al.</i>
<b>P-I-61</b>	Imaging G-protein signaling by polarization microscopy	<i>P. S. Miclea et al.</i>
<b>P-I-62</b>	POPDC proteins – potential regulators of compartmentalized cAMP signaling	<i>L. M. Martin et al.</i>
<b>P-I-63</b>	Development of BRET-based G protein biosensors to study ligand bias	<i>K. Boon et al.</i>
<b>P-I-64</b>	Deconvolution of CCR7 signalling using label-free cellular electrical impedance	<i>N. Vanalken et al.</i>
<b>P-I-65</b>	Characterization of PH sensitive ligands at the $\beta 1$ adrenergic and the $\mu$ opioid receptor	<i>T. Pröll et al.</i>
<b>P-I-66</b>	TRPV1 modulation of MOR signalling	<i>J. Sanchez et al.</i>
<b>P-I-67</b>	Ligand-induced changes in the plasma membrane organisation of the chemokine receptor CXCR4	<i>N. Karsai et al.</i>
<b>P-I-68</b>	Multivalent VHH formats as modulators of chemokine receptor CXCR4 oligomerization	<i>S. M. Anbuhl et al.</i>
<b>P-I-69</b>	Distinct roles of $\beta$ -arrestins in GPCR endocytosis	<i>J. Liu et al.</i>
<b>P-I-70</b>	HCMV-encoded viral GPCR US28 signals to the Hippo pathway via Gq/11	<i>E. M. Pfeil et al.</i>
<b>P-I-71</b>	Spatiotemporal signalling and location of fractalkine receptor CX3CR1 and its natural genetic variants associated to disease CX3CR1-V249I/T280M and CX3CR1-A55T	<i>M. Castro et al.</i>
<b>P-I-72</b>	The role of spinophilin in modulating the K-opioid receptorsignaling	<i>A. Symeonof et al.</i>
<b>P-I-73</b>	Biased signaling due to oligomerization of a G protein-coupled receptor	<i>J. Liu et al.</i>
<b>P-I-74</b>	Design of "CIS-ON" agonists as opticalmodulators of the human cannabinoid receptor type 1 (hCB1R)	<i>D. A. Rodriguez Soacha et al.</i>

<b>P-I-75</b>	Suitable use of FRET-based Biosensors for Quantitative Detection of GPCR Activation	<i>N. K. Brinkenfeldt et al.</i>
<b>P-I-76</b>	Conservative mutations of key lysines in the N-domain of arrestin inhibit C-tail release and thereby attenuate arrestin interactions with GPCRs	<i>A. K. Sztyler et al.</i>
<b>P-I-77</b>	Modulation of Dimerization and Signaling in Melanocortin-4 Receptor (MC4R) by accessory protein MRAP2	<i>I. Sohail et al.</i>
<b>P-I-78</b>	Prime time for unlocking inhibition of Gs?	<i>F. Eryilmaz et al.</i>
<b>P-I-79</b>	Towards unveiling the importance of Gα12/13 signaling using the Gq selective inhibitor FR900359	<i>N. Merten et al.</i>
<b>P-I-80</b>	Discovery of new synthetic agonists and antagonists for OR51E1, an ectopically expressed olfactory receptor linked to prostate cancer.	<i>V. Slepak et al.</i>
<b>P-I-81</b>	Dissecting CCR5 activation mechanism	<i>C. Branco et al.</i>
<b>P-I-82</b>	The Gβ subunit is the secret mediator in Gαq/11 inhibition	<i>J. Mühle et al.</i>
<b>P-I-83</b>	Development of Generic G Protein Peptidomimetics Able to Stabilize Active State Gs Protein-Coupled Receptors for Application in Drug Discovery	<i>S. Ballet et al.</i>

## P-II – Holistic Concepts in GPCR Signal Transduction

<b>P-II-01</b>	Gi/o and Gq/11 proteins cooperate in asymmetric GPCR dimers to activate ERK signaling	<i>E. Cecon et al.</i>
<b>P-II-02</b>	Molecular origins and principles governing adrenaline efficacy and potency in the human β2-adrenergic receptor	<i>F. M. Heydenreich et al.</i>
<b>P-II-03</b>	The role of preformed complexes between G protein-coupled receptors and G proteins in cell signaling	<i>A. Bondar et al.</i>
<b>P-II-04</b>	Bioluminescence Resonance Energy Transfer and NanoLuciferase-based complementation assays for the real-time monitoring of mitochondrial protein interactions	<i>A. O. Abdulrahman et al.</i>
<b>P-II-05</b>	The extended N-terminal domain confers atypical chemokine receptor properties to CXCR3-B	<i>G. D'Uonno et al.</i>
<b>P-II-06</b>	Agonist- and expression-dependent coupling of the promiscuous adenosine A2B receptor to Gα protein subunits	<i>J. H. Voss et al.</i>
<b>P-II-07</b>	Investigating the carboxy-terminal phosphorylation of the BB2 bombesin receptor using phospho-site-specific antibodies	<i>D. Freund et al.</i>

**P-II-08** A Fusion Protein Platform for Analyzing Tethered Agonism in the Adhesion Family of G Protein-Coupled Receptors *A. N. Dates et al.*

**P-II-09** The pre-existence of G protein heterotrimer is not required for G $\beta$ -mediated internalization of the C-C chemokine receptor 8 (CCR8) *L. Liu et al.*

### P-III – Adhesion GPCRs: from Structure to Function / Physiology

**P-III-01** Structural studies on the ADGRB2 (BAI2) GAIN domain and its resistance to GPS autoproteolysis *F. Pohl et al.*

**P-III-02** Agonist efficacy at the  $\beta$ 2-adrenoceptor ( $\beta$ 2AR) is driven by agonist induced conformational differences that increase the affinity andkonof the Gsprotein rescrutment *C. Harwood et al.*

**P-III-04** Potential allosteric modulators of the  $\beta$ 2-adrenergic receptor ( $\beta$ 2AR): an in silico and in vitro study *M. Sencanski et al.*

**P-III-03** Alternative splicing of Adhesion GPCR is required for neuronal mechanosensing *N. Scholz et al.*

**P-III-06** Elucidating the role of the aGPCR Latrophilin-1 in mammalian synapse formation *D. Matúš et al.*

**P-III-07** The role of GPR110 (ADGRF1) in osmotic regulatory processes in the kidney *S. Pick et al.*

**P-III-08** From sequence to function: the variant repertoire of the Latrophilin homolog LAT-1 *V. E. Groß et al.*

**P-III-09** The enigmatic trans function of the Adhesion GPCR Latrophilin cross-talks with the Notch pathway in germ cell proliferation *W.B. Post et al.*

**P-III-10** ST171, a Gi-biased 5-HT1A receptor agonist: ligand development and cryo-EM *A. Ullrich et al.*

**P-III-11** Tethered agonist activated GPR110-miniG $\alpha$ s/q structure reveals molecular preference for G $\alpha$ q signaling *D. T.D. Jones et al.*

**P-III-12** The N Terminus of Adhesion G Protein-Coupled Receptor GPR126/ADGRG6 as Allosteric Force Integrator *J. Mitgau et al.*

## P-IV – GPCR Signal Transduction in Physiology and Disease

<b>P-IV-01</b>	Activation of M1-muscarinic acetylcholine receptor reduces pathology and slows progression of neurodegenerative disease	<i>L. Dwomoh et al.</i>
<b>P-IV-02</b>	Nanobodies targeting the viral GPCR US28 for investigation of US28 signaling pathways, targeted cancer therapy and clearance of latent human cytomegalovirus reservoir	<i>T. W. De Groof et al.</i>
<b>P-IV-03</b>	Deciphering the role of the GPR101 orphan receptor in growth hormone hypersecretion	<i>D. Abboud et al.</i>
<b>P-IV-04</b>	Pharmacological Gq targeting induces strong pulmonary vasorelaxation in health and disease	<i>A. Seidinger et al.</i>
<b>P-IV-05</b>	Targeting serotonin 5-HT7 receptor to ameliorate Tau pathology and memory deficits	<i>E. Ponimaskin et al.</i>
<b>P-IV-07</b>	Pharmacology of kappa opioid receptors: novel assays and ligands	<i>D. Malfacini et al.</i>
<b>P-IV-08</b>	Interpreting the molecular mechanisms of disease variants in GPCRs and human membrane proteins	<i>J. K.S. Tiemann et al.</i>
<b>P-IV-09</b>	Noradrenergic Regulation of Astroglial Aerobic Glycolysis and Lipid Droplet Metabolism in Health and Disease	<i>N. Vardjan et al.</i>
<b>P-IV-10</b>	GPCRs in the era of biobanks	<i>A. S. Hauser et al.</i>
<b>P-IV-11</b>	Serotonin receptors - novel targets in the treatment of amyotrophic lateral sclerosis	<i>J. Labus et al.</i>
<b>P-IV-12</b>	Bivalent Ligands Targeting the D1R-H3R and D2R-H3R Heteromer	<i>S. Pockes</i>
<b>P-IV-13</b>	Chemogenomic approach to identifying chemoreceptor drug targets in parasitic nematodes	<i>A. Langeland et al.</i>
<b>P-IV-14</b>	Identification of melatonin-like activities in plant extracts	<i>N. Labani et al.</i>
<b>P-IV-15</b>	Application of BRET Method to Investigate the Functional Properties of GPCR-like Proteins of the Plasmodium falciparum	<i>M.K. Singh et al.</i>
<b>P-IV-16</b>	Duchenne muscular dystrophy: The role of eicosanoids and their GPCRs in bone health	<i>M. Hoxha et al.</i>
<b>P-IV-17</b>	Different G protein regulation profile for hallucinogenic and non-hallucinogenic 5-HT2A receptor drugs in postmortem human brain	<i>R. Diez-Alarcia et al.</i>
<b>P-IV-18</b>	Evaluation of G-protein signalling pattern and polypharmacological profile of different antipsychotic drugs in post-mortem human native brain tissue.	<i>I. Muneta-Arrate et al.</i>



<b>P-IV-19</b>	A nanoBRET-based Ligand Binding Assay for the Melanocortin Receptors to Find Novel Variants by Phage Display	<i>M. C. Troll et al.</i>
<b>P-IV-20</b>	Identification of an orphan GPCR, GPR151, as a potential actor in reproduction and metabolic function	<i>C. Jacquinet et al.</i>
<b>P-IV-21</b>	Stimulation of the free fatty acid receptor 4 (FFAR4) induces strong pulmonary vasorelaxation	<i>A. Seidinger et al.</i>
<b>P-IV-22</b>	The expression changes of the individual aGPCRs in lung adenocarcinoma cell line infected with SARS-CoV-2	<i>J. Trylcova et al.</i>
<b>P-IV-06</b>	Functional analysis of melanocortin-4 receptor variants linked to obesity	<i>A. V. Rodríguez Rondón et al.</i>
<b>P-IV-24</b>	Analysis of high-impact missense variants in G Protein-Coupled Receptors	<i>A. Peralta-García et al.</i>
<b>P-IV-25</b>	Fluorescent peptide tracers for oxytocin receptor visualization	<i>M. Perisic et al.</i>
<b>P-IV-26</b>	Dual-acting small molecules: selective CB2R/BChE hybrids with agonism on CB2R show neuroprotective activity in an Alzheimer's disease mouse model	<i>P. Spatz et al.</i>
<b>P-IV-28</b>	A small molecule agonist of the atypical chemokine receptor 3 (ACKR3) alleviates fibrosis in a preclinical liver but not lung injury model	<i>T. Van Loy et al.</i>
<b>P-IV-29</b>	GPCRs and their role in EGF-mediated small airway epithelial remodeling	<i>J. van den Bor et al.</i>
<b>P-IV-30</b>	Nanobody-based sensors reveal a high proportion of mGlu heterodimers in the brain	<i>P.-A. Lafon et al.</i>
<b>P-IV-31</b>	Hypertension causing PDE3A catalytic mutation leads to alteration of cAMP signaling	<i>C. Kayser et al.</i>
<b>P-IV-32</b>	A2B agonists as vasorelaxants in pulmonary arteries	<i>J. Lewandowski et al.</i>
<b>P-IV-33</b>	Bivalent Ligands for the D1R-H3R-Heteromer Reduce $\beta$ -Amyloid-Induced Cell Death	<i>N. Rosier et al.</i>
<b>P-IV-34</b>	Synthesis and biological evaluation of bivalent ligands targeting the D2-H3 receptor heteromer	<i>M. Nagl et al.</i>
<b>P-IV-35</b>	Regulation of the GPCRome in adipose tissue of obese individuals	<i>I. Kaczmarek et al.</i>
<b>P-IV-36</b>	Metabolite-sensing GPCRs mediate cross-talk of adipocyte signaling and metabolic state with transcriptional regulation in obesity	<i>M. Sandhu et al.</i>
<b>P-IV-37</b>	Phosphorylation state-specific antibodies as biosensors for $\beta$ 2-adrenoceptor activation in vivo	<i>E. Miess-Tanneberg et al.</i>
<b>P-IV-38</b>	Effect of mutations causing visual disease, and trace metals, on the structure and stability of the visual G protein-coupled receptor rhodopsin	<i>P. Fernandez et al.</i>

**P-IV-39** Characterization of the D1R-H3R and D2R-H3R heteromers in co-expressing systems using bivalent ligands *D. Moennich et al.*

**P-IV-40** Signatures of cannabinoids phosphorylation patterns *V. Stammer et al.*

**P-IV-41** Role of GRKs in MOP mediated effects and side effects in vivo. *B. Herböck et al.*

## P-V – Computational Design and GPCR Therapeutics

**P-V-01** Discovery of small molecule modulators for FZD7 using in silico docking screens *M. M. Scharf et al.*

**P-V-02** AZD5462 is a clinical, oral small molecule RXFP1 agonist that mimics the signaling pharmacology of relaxin-2, the cognate peptide ligand at this receptor *N. Larsson et al.*

**P-V-03** Identification of Novel Pyrazolo[3,4-c]pyridine Antagonists with Nanomolar affinity and Long Residence Time for A1 / A3 Adenosine Receptors and SAR Investigation Using Mutagenesis Experiments and TI/MD calculations *M. Stampelou et al.*

**P-V-04** Discovery and development of Ligands for Super-Conserved Receptors Expressed in the Brain (SREB) *T. Pillaiyar et al.*

**P-V-06** GPCR photopharmacology: using light operated ligands for a precise control of receptor activity *A. Llebaria*

**P-V-07** Combined docking and machine learning identifies key molecular determinants of ligand pharmacological activity on 2 adrenoceptor. *D. B. Veprintsev et al.*

**P-V-08** Structure-based comparisons of GPCR interactions with nanobodies and G proteins *Z. Shapiro Tuchman et al.*

**P-V-10** Computational and In Vitro Evaluation of Potent Thiazolo[4,5-d]pyrimidine Corticotropin Releasing Factor (CRF) Receptor Antagonists *V. Panagiotopoulos et al.*

**P-V-11** Targeting Cancer Cells with Peptide-Drug-Conjugates addressing the CMKLR1 Receptor *A. S. Czerniak et al.*

**P-V-12** Selective Addressing of Adipocytes with the F,P-NPY/Y1R -System *A. Kohler et al.*

**P-V-13** Interpretability tools in the prediction of properties of biologically active compounds *S. Podlowska et al.*

**P-V-14** Discovery of Novel Allosteric CCR9 Ligands as Anticorectal Cancer Agents by Sequential Virtual Screening *Y. M. Mandour et al.*

**P-V-15** Evolution of Neuropeptide Y/Rfamamide-like receptors in Nematodes *F. Reinhart et al.*

**P-V-16** Conformational thermostabilisation of GPCRs for structure-based drug design *F. Autore et al.*

<b>P-V-17</b>	A single point mutation blocks the entrance of ligands to the cannabinoid CB2 receptor via the lipid bilayer	<i>N. Casajuana-Martin et al.</i>
<b>P-V-18</b>	Virtual screening of known allosteric GPCR binding sites using a DOCK3.7-based workflow	<i>M. Persechino et al.</i>
<b>P-V-19</b>	Discovery of Novel Cannabinoid Receptor-2 (CB2R) Agonists and Inverse Agonists via In-Silico Docking	<i>M. Dennis et al.</i>
<b>P-V-20</b>	Bitter peptide GPCRs: identifying peptide signatures and key receptor residues using computational tools	<i>A. Steuer et al.</i>
<b>P-V-21</b>	Estimation of the absolute free energy of binding using the linear interaction energy method with continuum electrostatics for a diverse set of ligands of the 5-HT2A receptor	<i>A. Shahraki et al.</i>
<b>P-V-22</b>	Molecular determinants for species differences in Histamine H4 receptor	<i>T.D. Tran et al.</i>
<b>P-V-23</b>	Ultra-large Library Docking in Rosetta EvoLigand Tackles Challenging GPCR Targets In Drug Discovery	<i>F. Liessmann et al.</i>
<b>P-V-24</b>	Wanted urgently: Specific GPR151 ligands	<i>C. S. Tautermann</i>

## P-VI – Molecular Structure and Dynamics of GPCRs

<b>P-VI-01</b>	Structural insights into glucagon-like peptide 1 receptor activation and allosteric modulation by non-peptidic ligands	<i>X. Zhang et al.</i>
<b>P-VI-02</b>	Dynamic drug targets: Using Cryo-EM data and MD simulations to create realistic 3D animations of Class B1 GPCR activation	<i>S. Piper et al.</i>
<b>P-VI-03</b>	Can novel structures of 7TM ancestors reveal a missing link to evolved signaling pathways?	<i>M. Shalev-Benani</i>
<b>P-VI-04</b>	Cholesterol Binding to Active and Inactive States of Adenosine Receptor Subtype-1	<i>E. Tzortzini et al.</i>
<b>P-VI-05</b>	Allosteric modulations of human kinin G-protein-coupled receptors bradykinin 1 receptor-peptide agonist interactions by ACE inhibitors	<i>J. Mao et al.</i>
<b>P-VI-06</b>	Investigating nanobody interactions with Chemokine Receptors CXCR4 and ACKR3 using Hydrogen/Deuterium Exchange Mass Spectrometry	<i>O. Otun et al.</i>
<b>P-VI-07</b>	Novel single color fluorescent GPCR biosensors to study allosteric coupling in GPCR activation	<i>R. Thomas et al.</i>

<b>P-VI-08</b>	The extracellular N-terminus of natural receptor isoforms allosterically modulates GPR35-transducer coupling and mediates intracellular pathway bias	<i>H. Schihada</i>
<b>P-VI-09</b>	How to avoid hallucinogenic side effects? Comparative investigation of ibogainalog (IBG) and its non-hallucinogenic tabernanthalog (TBG)	<i>D. J. Kiss et al.</i>
<b>P-VI-10</b>	Small label ≠ no effect: how labelling of arrestin-1 influences activation and GPCR binding	<i>B. Bollmann et al.</i>
<b>P-VI-11</b>	Investigating Cell Free Expressed Neuropeptide Y4 Receptor by using Photo-Crosslinking Studies	<i>V. Behr et al.</i>
<b>P-VI-12</b>	Molecular Dynamics simulations of human $\beta 1/\beta 2$ -adrenoceptors provide a possible explanation for the effect of the I118/H93 mutation in transmembrane 2 on the $\beta 1$ selectivity of ICI89406 and xamoterol.	<i>J.Y.V. Lim et al.</i>
<b>P-VI-13</b>	Investigation of the Histidine residues of the Growth Hormone Secretagogue Receptor by Solid-State NMR	<i>E. M. Pacull et al.</i>
<b>P-VI-14</b>	Pairwise crosslinking reveals details of neuropeptide Y binding at the Y5 receptor	<i>K. Philipp et al.</i>
<b>P-VI-15</b>	Structural basis of neuropeptide Y signaling through human neuropeptide Y1 receptor	<i>J. Kim et al.</i>
<b>P-VI-16</b>	Mapping the Dynamic Interface of $\beta$ -Arrestin 1 and a Secretin-like GPCR via Genetically Encoded Crosslinkers.	<i>Y. Aydin et al.</i>
<b>P-VI-17</b>	Structural analysis of the neurokinin 1 receptor in complex with Gq and Gs proteins reveals substance P binding mode and unique features of activation and antagonism	<i>C. Thom et al.</i>
<b>P-VI-18</b>	Studies of GPCR oligomerization in a eukaryotic Cell-free System	<i>J. Ullrich et al.</i>
<b>P-VI-19</b>	The Conformational Changes in Gq Activated by the M3 Muscarinic Acetylcholine Receptor	<i>D. Ham et al.</i>
<b>P-VI-20</b>	Interaction between Dopamine D2 receptor and $\beta$ -arrestin 2	<i>K. Kim et al.</i>
<b>P-VI-21</b>	The Human Neuropeptide Y Type 4 Receptor (hY4R) and its Extracellular Coupling Partners: Insights into Structural Dynamics via Solid-State NMR Spectroscopy	<i>M. Gozzi et al.</i>
<b>P-VI-22</b>	Application of MD simulation and protein-protein docking for analysis of the structural effect of mutations in GPCR	<i>N. Kulik et al.</i>
<b>P-VI-23</b>	In silico study of GPCR signaling bias using molecular dynamics and machine learning	<i>A. Morales Pastor et al.</i>

<b>P-VI-24</b>	A Novel Non-radioactive Binding Assay for the Neuropeptide Y5 Receptor	<i>L. C. Peuker et al.</i>
<b>P-VI-25</b>	Mapping Interactions of the M2-Receptor with Arrestin-3 in the Live Cell	<i>T. Müller et al.</i>
<b>P-VI-26</b>	Structural investigations of Arrestin-3 in complex with neuropeptide Y receptors	<i>V. Gupta et al.</i>
<b>P-VI-27</b>	Molecular mechanism of activation and transducer-coupling of $\beta$ -arrestin-biased receptors	<i>J. Maharana et al.</i>
<b>P-VI-28</b>	MYCBP2, a Novel G $\alpha$ s $\alpha$ -Helical Domain-binding partner	<i>D. Ahn et al.</i>
<b>P-VI-29</b>	Novel insights into GPCR dynamics	<i>D. Aranda-Garcia et al.</i>
<b>P-VI-30</b>	Evolution of the sodium binding site in the angiotensin II receptors: Implications for the receptor functional properties	<i>M. Chabbert et al.</i>
<b>P-VI-31</b>	Interactions of Neuropeptide Y with Y2 Receptor studied by Cross-linking Mass Spectrometry	<i>J. C. Rojas Echeverri et al.</i>
<b>P-VI-32</b>	Minimal-sized FRET sensors for studying conformational changes in GPCRs	<i>C. De Faveri et al.</i>
<b>P-VI-33</b>	Common structural features in the vasopressin receptor 2 and the $\mu$ -opioid receptor for biased signaling	<i>X. Cong et al.</i>
<b>P-VI-34</b>	A small genetically encoded fluorophore for conformational change tracking	<i>A. López Sierra et al.</i>
<b>P-VI-35</b>	Distinct Binding Mode of Neuropeptide Y toward the Y2 Receptor	<i>C. Park et al.</i>
<b>P-VI-36</b>	In situ Visualization of therapeutic drug effects using phosphosite-specific GPCR antibodies	<i>S. Fritzwanker et al.</i>
<b>P-VI-37</b>	Heteromerization and kinetic screening of metabotropic glutamate receptors	<i>T. Kukaj et al.</i>
<b>P-VI-38</b>	mdciao: Accessible Analysis and Visualization of Molecular Dynamics Simulation Data	<i>G. Pérez Hernández et al.</i>
<b>P-VI-39</b>	Mechanistic study of receptor mediated G-protein activation	<i>H. Batebi et al.</i>
<b>P-VI-40</b>	The molecular mechanism of tethered peptide binding in adhesion GPCRs	<i>R. Guixà-González et al.</i>
<b>P-VI-41</b>	Analysis of G proteins and family-specific inhibitors by solution and solid-state NMR spectroscopy	<i>C. Bonifer et al.</i>
<b>P-VI-42</b>	Biophysical characterisation of cell-free synthesized free fatty acid receptor 2 via various <sup>19</sup> F probes and novel solid-state NMR approaches	<i>S. Seidl et al.</i>
<b>P-VI-43</b>	Insights into the dynamics of human neuropeptide Y Y1 receptor with solid-state NMR spectroscopy	<i>M. Voitel et al.</i>
<b>P-VI-44</b>	Investigating mGluR cooperativity with FRET	<i>S. Dadashkhan et al.</i>

<b>P-VI-45</b>	Preambly of 5-hydroxytryptamine receptor type 7 (5HT7R) with the Gs protein: snapshots from protein-protein interface models	Z. Zara <i>et al.</i>
<b>P-VI-46</b>	Structural mechanism of endogenous MC4R agonists b-MSH and a-MSH binding reveals a novel interface for drug design	N. A. Heyder <i>et al.</i>
<b>P-VI-47</b>	Investigating the Conformational Dynamics of the Y2 -Receptor by Site-directed Spin Labeling EPR Spectroscopy	J. M. Laugwitz <i>et al.</i>
<b>P-VI-48</b>	Detection of GPCR oligomerization in single cells by quantitative fluorescence microscopy	G. J. von Rosing <i>et al.</i>
<b>P-VI-49</b>	Structural and biochemical characterization of G<math>\alpha\gamma</math> interactions with a pre-fusion trans-SNARE mimetic	A. R. Eitel <i>et al.</i>
<b>P-VI-50</b>	Dynamics of -Arrestin Interaction with a G-Protein Coupled Receptor	M. Pankonin <i>et al.</i>
<b>P-VI-51</b>	Cryo-EM studies on GPCR-G protein complexes	O. Tejero <i>et al.</i>



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## **GALA DINNER AT LEIPZIG ZOO**

**Date:** Tuesday, 27  
September 2022

**Entrance:** 19:00

**Address:** Gondwanaland  
Pfaffendorfer Straße 25

In Germany's largest tropical adventure world, Gondwanaland, the beautifully Asian-designed Patakan restaurant is at your disposal. Experience the flora and fauna of the rainforest after the zoo has closed and gain an insight into the history of the earth's origins during a boat trip on the primeval forest river Gamanil.

The ticket price of 70.00 EUR per person includes the following services:

- the entrance to the tropical hall, Gondwanaland, with a boat tour through the rainforest
- Welcome drink
- varied Buffet and midnight snack
- alcoholic free drinks, beer and wine

**Tickets can be purchased directly at the registration counter.\***

*\*According to availability*





### **CLOSING PARTY AT MORITZBASTEI LEIPZIG**

#### **Early Career Investigator Session & Awarding**

**Date:** Thursday, 29 September 2022

**Entrance:** 18:00

**Address:** Moritzbastei Leipzig | Kurt-Masur-Platz 1

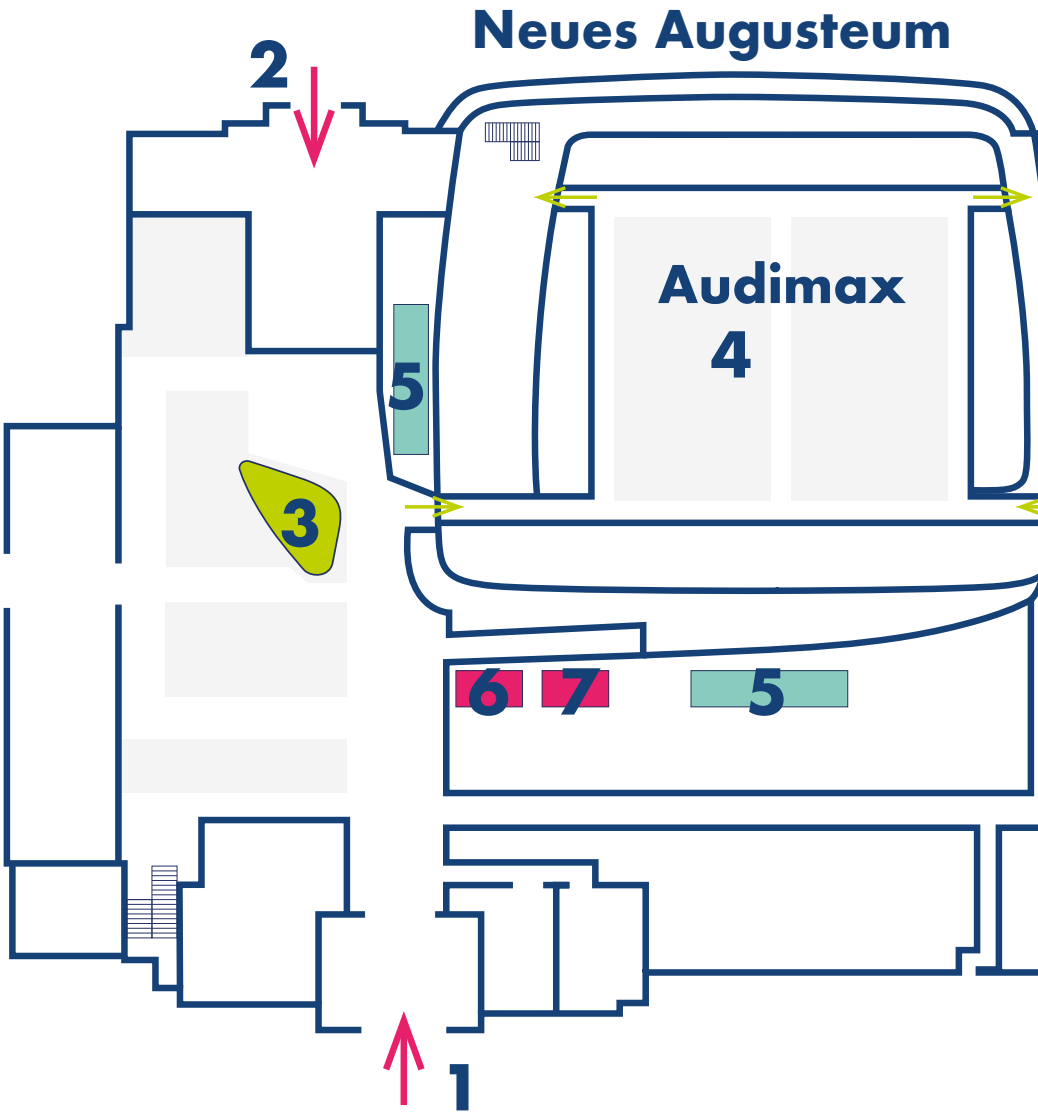
The Moritzbastei is Leipzig's best-known cultural centre. Located in the heart of the city, right next to the New Gewandhaus and the university, it combines historical architecture and modern cultural life in all its diversity.

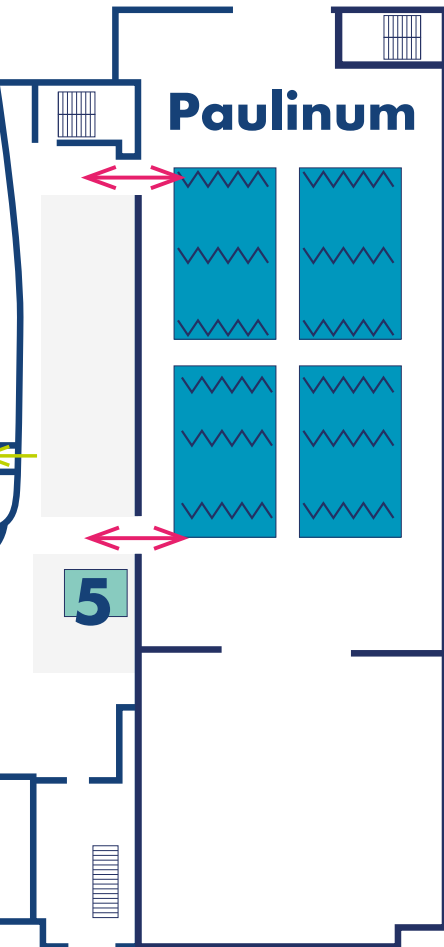
The Moritzbastei is the last remaining part of Leipzig's old city fortifications. Since it was built in 1551-1553, it can look back on an eventful history.

Enjoy the evening in convivial company incl. drinks, buffet and music.

Admission is free of charge. However, for better planning, we kindly ask you to register.







## LEIPZIG UNIVERSITY NEUES AUGUSTEUM

Campus Augustusplatz  
04103 Leipzig, Germany

- 1.** Entrance via Augustusplatz
- 2.** Entrance via Leibniz-Forum
- 3.** Registration Counter
- 4.** Audimax
- 5.** Catering
- 6.** Promega GmbH
- 7.** Keyence Deutschland GmbH

## LEIPZIG UNIVERSITY PAULINUM

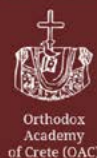
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- P-III**
- P-IV**
- P-V**
- P-VI**

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