

Yearbook 2008/2009

**MSc/PhD/MD-PhD
Neuroscience Program**
at the University of Göttingen

**International Max Planck
Research School**

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Letter from the President

The international Master's / PhD Programs Molecular Biology and Neurosciences were established by the Georg August University Göttingen, together with the Max Planck Society for the Advancement of Science, in the year 2000 to attract excellent students from all over the world and provide them with an outstanding, research-oriented graduate program. Both programs are taught in English by internationally renowned scientists and offer a high level of services and individual support.

Several hundred students from all over the world apply for the 20 study places available in each of the programs every year. Both programs have introduced and combined elements of international recruitment, competitive admission procedures, advanced curricula, research training, social integration programs, extracurricular support and evaluation procedures into successful working structures. They have achieved excellent recommendations in several external evaluations and have been awarded the 2004 prize for excellent support services for foreign students by the German Federal Foreign Office. For the newly established Georg August University School of Science (GAUSS) and other graduate schools in Göttingen, the Molecular Biology and Neuroscience Programs are considered exemplary and serve as best practice models.

In October 2006, the two programs were awarded the label „Top 10 International Master's Degree Courses made in Germany“ by the „Stifterverband für die Deutsche Wissenschaft“ and the German Academic Exchange Service (DAAD) in a national contest, in which 121 Master's programs of 77 universities participated. The Göttingen Molecular Biology and Neuroscience programs were the only Master's programs in the natural sciences and medicine which received this award. Both programs are members of the Göttingen Graduate School for Neurosciences and Molecular Biosciences (GGNB), which was successful in the recent Excellence Competition by the German Federal and State Governments to promote science and research at German universities.

Five Göttingen University faculties, three Göttingen Max Planck Institutes as well as the German Primate Center participate in the programs. International guest lecturers are also involved. The Max Planck Society contributes through its newly established International Max Planck Research Schools. Both programs keep close contacts with the relevant industries to further enhance the chances of the graduates for a successful professional career.

I would very much like to thank all scientific bodies and institutions for their committed support in establishing these international programs and, last but not least, the German Academic Exchange Service (DAAD), the Lower Saxony Ministry of Science and Culture, and the various generous donors.

The Georg August University of Göttingen is proud of its long-standing international experience the two attractive and innovative programs have already become an integral part of. The university will continue to support these programs within the setting of Göttingen's lively urban, cultural, and social life, in itself a prerequisite for creative teaching and research.

Prof. Dr. Kurt von Figura
(President of the Georg August University Göttingen)



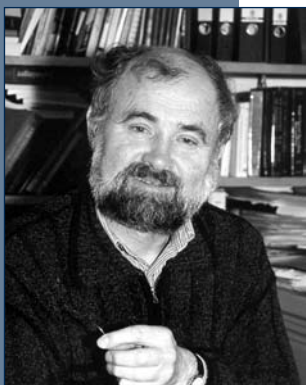
Letter from the Max Planck Society

The mission of the Max Planck Society is to conduct basic research in science and humanities at the highest level. More than 80 Max Planck Institutes are located on scientific campuses across Germany, most of them close to universities.

Scientific ties between Max Planck Institutes and universities are traditionally strong. In 1998, during the 50th year celebration of the Max Planck Society in Göttingen, the Max Planck Society, together with the Hochschulrektorenkonferenz, launched the International Max Planck Research Schools as a new joint program to further intensify cooperation.

The goals of the International Max Planck Research Schools are

- to attract excellent students from all around the world to intensive Ph.D. training programs in Germany, preparing them for careers in science,
- to integrate Max Planck scientists in top-level scientific training of junior scientists,
- to intensify the ties to the universities owing to the participation of internationally renowned Max Planck scientists in joint teaching activities, and
- to strengthen international relationships by providing individual support to each student and by exposing foreign students to German culture and the German language.



By now, 51 International Max Planck Research Schools have been established involving 65 Max Planck Institutes, 48 German universities with 70 participating faculties and more than 15 universities abroad. More than 1900 (mostly PhD) students from 87 countries are presently enrolled. Approximately 850 PhD students have graduated to date from an International Max Planck Research School.

Since their foundation in the year 2000, the Göttingen International Max Planck Research Schools in Molecular Biology and Neurosciences have met with extraordinary success. Every year, the programs receive hundreds of applications, with the quality of the students consistently being very high. Most students graduated so far have moved on to postdoctoral positions, many at prestigious international institutions. In the past years, the Göttingen Schools received unanimous acclaim during external evaluations and won national awards. For instance they are the only Life Science Programs within Germany that were selected for the “Top Ten International Master’s Degree Courses”. The Schools have also reshaped the local scientific community, strengthening the ties between the participating institutions, and initiated new scientific collaborations that augment the international reputation of Göttingen as a center of scientific excellence. Furthermore, the Schools served as role models and founding members of the Göttingen Graduate School for Neurosciences and Molecular Biosciences, thus were being instrumental for the success of the University in the German Excellence Initiative. We hope that in the years to come the students of the International Max Planck Research Schools will be successful in their professional careers. We also hope that they will remember their training period in Göttingen as an exciting and stimulating phase in their lives.

Peter Gruss
President
Max Planck Society

Erwin Neher
Dean of the IMPRS
Neurosciences

Overview

This yearbook is intended to provide information on the International MSc/PhD/MD-PhD Neuroscience Program in Göttingen, Germany, which was established in 2000. In addition to general information on the program, the yearbook introduces the current year's students, the faculty members, the program committee, and the coordination team.

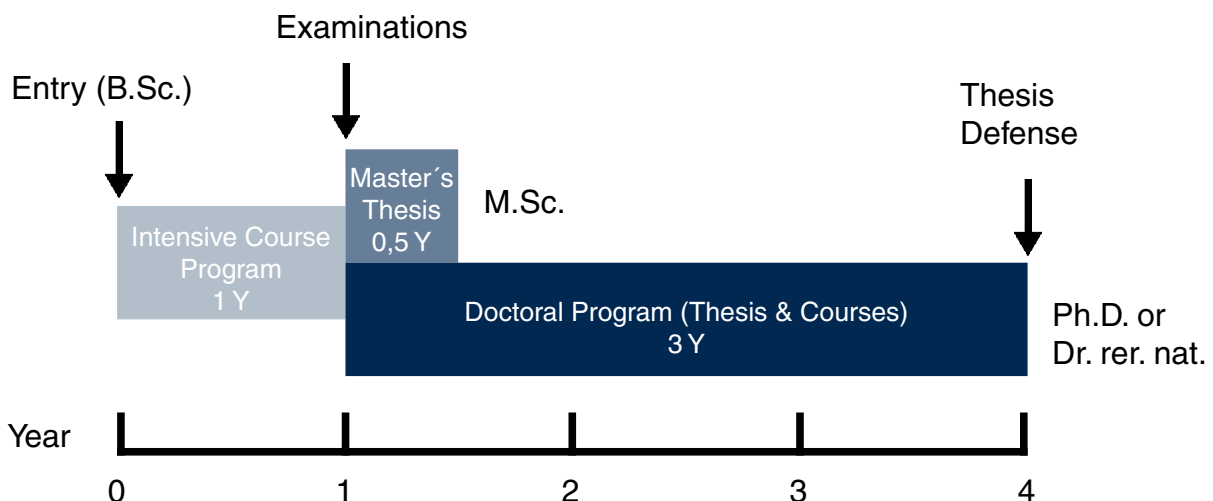
The program is a member of the Göttingen Graduate School for Neurosciences and Molecular Biosciences (GGNB), which is funded by the Excellence Initiative of the German Federal and State Governments. It is offered by the University of Göttingen, the Max Planck Institute for Biophysical Chemistry (MPIbpc), the Max Planck Institute for Experimental Medicine (MPIem), the Max Planck Institute for Dynamics and Self-Organization (MPIs), the German Primate Center (DPZ), and the European Neuroscience Institute (ENI). Further to their active participation in the Neuroscience Program, the above-mentioned partners closely cooperate in the DFG Research Center for Molecular Physiology of the Brain (CMPB), the Göttingen Center for Molecular Biosciences (GZMB), the Center for Systems Neuroscience (ZNV), in several collaborative research centers (Sonderforschungsbereiche, SFB) and in interdisciplinary doctoral programs (Graduiertenkollegs, GK).

The International MSc/PhD/MD-PhD Neuroscience Program qualifies students for professional work in the neurosciences. The program is open to students from Germany and from abroad, who hold a Bachelor's degree (or equivalent) in the biosciences, medicine, psychology, physics, or related fields. All courses are held in English. Scholarships are available. The academic year starts in October and is preceded by a three week orientation program. Applications may be submitted until January 15 of the year of enrollment. To ensure a high standard of individual training, the number of participants is limited to 20 students per year.

All students initially participate in one year of intensive course work. This first segment of the program comprises lectures, tutorials, seminars, methods courses, and independent, individually supervised research projects (laboratory rotations). The traditional German structure of academic semesters is not followed. The condensed schedule allows students to accumulate 90 credits (ECTS) within one year, which would normally require three semesters.

Subsequently, two separate segments are offered:

- **PhD Program:** Good to excellent results after the first year qualify for direct admission to a three-year doctoral project in one of the participating research groups. The Master's thesis requirement is waived in this case. After successful defense of a doctoral thesis, the degree Doctor of Philosophy (Ph.D.) or the equivalent title Doctor rerum naturalium (Dr. rer. nat.) is conferred. Students who finished medical school can apply for an MD-PhD title.
- **MSc Program:** Alternatively, students may conclude the program with a Master's thesis, based on six months of experimental scientific research. The degree Master of Science (MSc) is awarded upon successful completion of the Master's thesis.



Funding of the Program

The Neuroscience Program thanks the following institutions and funding initiatives, who contributed to the success of the Neuroscience Program:

DAAD

German Academic Exchange Service (DAAD),
Bonn, Germany, <http://www.daad.de>

*International Degree Programs -
Auslandsorientierte Studiengänge (AS)*


IPP made in Germany 

*International Postgraduate Programs –
Internationale Promotionsprogramme (IPP)*



Max Planck Society for the Advancement of Science,
Munich, Germany, <http://www.mpg.de>

International Max Planck Research Schools

 Niedersächsisches Ministerium
für Wissenschaft und Kultur

Ministry of Lower Saxony for Science and Culture,
Hannover, Germany, <http://www.mwk.niedersachsen.de/home/>

Innovationsoffensive

Doctoral Programs - Promotionsprogramme

Stifterverband
für die Deutsche Wissenschaft

Stifterverband für die Deutsche Wissenschaft,
Essen, Germany, <http://www.stifterverband.org>



Exzellenzstiftung zur Förderung der Max-Planck-Gesellschaft,
<http://www.exzellenzstiftung.de>

Gemeinnützige
Hertie-Stiftung 

Gemeinnützige Hertie-Stiftung, Frankfurt am Main,
<http://www.ghst.de>

Donors

The Neuroscience Program thanks the following companies for their donations, which were used to financially support students during the first year of studies:



Bayer AG, Leverkusen, Germany



Carl Zeiss Lichtmikroskopie, Göttingen, Germany



Degussa AG, Düsseldorf, Germany



DeveloGen AG, Göttingen, Germany



Heka Elektronik GmbH, Lambrecht / Pfalz, Germany



Hellma GmbH & Co. KG, Müllheim / Baden, Germany



KWS Saat AG, Einbeck, Germany



Leica Microsystems GmbH, Bensheim, Germany



Luigs & Neumann, Ratingen, Germany



Olympus Deutschland GmbH, Hamburg, Germany



Roche Diagnostics GmbH, Penzberg, Germany



Sartorius AG, Göttingen, Germany



Solvay Pharmaceuticals, Hannover, Germany



Springer Verlag, Heidelberg, Germany



Vossius & Partner, München, Germany

Intensive Course Program (First Year)

Throughout the first year, current topics in the neurosciences are covered by

- lectures
- tutorials
- methods courses
- laboratory rotations
- seminars

Lectures and Tutorials

A comprehensive lecture series is organized into a sequence of 4-6 week units. The following topics are taught on an advanced level throughout the first year (36 weeks, 4 hours per week):

- A. Neuroanatomy**
- B. Physiology and Basic Statistics**
- C. Modelling, Autonomous Nervous System, Pharmacology**
- D. Molecular Biology, Development, and Neurogenetics**
- E. Sensory and Motor Systems**
- F. Clinical Neurosciences and Higher Brain Functions**
- G. Specialization Seminars and Tutorials**

Each lecture is accompanied by a tutorial session, where students meet with a tutor in small groups. Tutorials involve exercises, review of lecture material, and discussion of related topics.

Methods Courses

During the first months of the Neuroscience Program, students participate in a series of methods courses to introduce them to principles and practical aspects of basic scientific techniques and the handling of model organisms. The practical courses and tutorials comprise the following topics:

I Neuroanatomy

- comparative development of the vertebrate brain
- cytology and ultrastructure of the human brain
- functional neuroanatomy of sensory and motor systems
- immunocytochemical techniques
- single neuron staining and recording
- invertebrate model systems

II Physiology and Basic Statistics

- introduction to medical statistics
- electrophysiological techniques
- membrane physiology / synaptic transmission
- FLIM / Ca-imaging / FCS techniques
- sensory and behavioral physiology

III Modelling, Autonomous Nervous System, Pharmacology

- neuronal modelling
- behavioral analysis
- neuroendocrinology / neuropharmacology
- protein separation techniques

IV Molecular Biology, Development, and Neurogenetics

- cell culture methods
- methods in molecular biology

Laboratory Rotations

Starting in January, every student carries out three independent research projects (laboratory rotations) in participating laboratories. Each project is individually supervised and involves seven weeks of experimental work, followed by one week for data analysis and presentation. For each project, a report must be completed in the format of a scientific publication. The laboratory rotations must cover at least two different subjects.

Seminars

Seminars start in March. The class meets weekly for two hours to discuss two student presentations. The presentations are research reports based on work from the laboratory rotations.

Examinations

After the first year of intensive training, all students take one written and two oral Master's examinations. The Master's examinations explore the students' theoretical background in topics covered by lectures and tutorials. All candidates are examined both in the field of anatomy and physiology in two separate oral exams.

PhD Program

Students who have passed the Master's examinations with good or excellent results qualify for direct admission to a three-year doctoral project in one of the participating research groups without being required to complete a Master's thesis first.

The PhD program emphasizes independent research on the part of the students. Doctoral students select three faculty members as their doctoral thesis committee which closely monitors progress and advises students in their research project. Laboratory work is accompanied by seminars and lecture series, a wide variety of advanced methods courses, training in scientific writing and oral presentation skills, courses in intercultural communication, bioethics and research ethics, elective courses, and participation in international conferences or workshops.

At the end of the PhD training program, a doctoral thesis is submitted either in the traditional format, or as a collection of scientific publications in internationally recognized journals along with a general introduction and a discussion of the results. The degree PhD or, alternatively, Dr. rer. nat. will be awarded after the successful defense of the doctoral thesis. Having fulfilled all PhD degree requirements, medical students may apply for the degree of an MD-PhD at the Medical Faculty.

Master's Program

After the first year of intensive training, students may conclude the program with a six-month thesis project, leading to a Master of Science degree. The thesis project involves experimental work under the supervision of faculty members of the Neuroscience Program. Students have the opportunity to conduct their Master's thesis project at a research institution abroad.

Orientation, Language Courses, Social Activities

A three-week orientation prior to the program provides assistance and advice for managing day-to-day life, including arrangements for bank account, health insurance, residence permit, housing, and enrollment. Students have the opportunity to meet faculty members and visit laboratories of the participating institutions. In addition, the orientation program informs students about computing and library facilities, the city and university of Göttingen, sports facilities, and cultural events.

An intensive basic language course in German is offered in cooperation with the *Lektorat Deutsch als Fremdsprache* to facilitate the start in Göttingen. Additional language courses and social activities accompany the program.

Application, Selection and Admission 2008

Applicants must hold a Bachelor's degree or equivalent in biology, medicine, psychology, physics, or related fields. Applicants who are not native speakers of English should demonstrate adequate competence of the English language by acceptable results in an internationally recognized test.

In the year 2008, the coordination office received 177 applications from 36 countries

Continent	Applications	Admissions
Europe (total)	41	9
Germany	28	7
other West Europe	6	2
East Europe	7	0
America (total)	10	3
North America	2	0
Central/South America	8	3
Africa (total)	8	0
North Africa	3	0
Central/South Africa	5	0
Asia (total)	118	8
Near East	33	0
Central Asia/ Far East	85	8
Australia	0	0

Students 2008/2009

Name		Home Country
Malte	Alf	Germany
Jonas	Barth	Germany
I-Wen	Chen	Taiwan
Carolina	Cunha	Brazil
Juan Daniel	Flórez Weidinger	Colombia
Gorur Shandilya	Srinivas	India
Xiaojie	Huang	P.R. China
Cordelia	Imig	Germany
Zhizi	Jing	P.R. China
Maria-Eleni	Kastriti	Greece
Alejandro	Mendoza Schulz	Germany
Jatin	Nagpal	India
Srinivas	Parthasarathy	India
Natalia	Revelo Nuncira	Colombia
Meike Annika	Schweisfurth	Germany
Nicolas	Snaidero	France
Swathi	Srivatsa	India
Roman	Stilling	Germany
Benjamin	Wilhelm	Germany
Aaron Benson	Wong	Hong Kong, China



Germany

Malte Alf

EDUCATION

College / University

2004 – 2007: Freie Universität Berlin

2007 – 2008: Yale University Grad School

Highest Degree

Vordiplom in Physics, Vordiplom in Psychology

Major Subjects

Physics, Psychology, Cognitive Neuroscience

Lab Experience

Basic physics and inorganic chemistry labs, eye tracking and EEG at FU Berlin, fMRI at MPI for Human Development and Yale University

Projects / Research

2006 – 2007: fMRI study at MPI for Human Development (Research assistant for U. Lindenberger and Shu-Chen Li)

2007 – 2008: fMRI/DTI study at Yale U (independent work, advisers Marcia Johnson and Todd Constable)

Scholarships / Awards

2008 – 2009: International Max Planck Research School support

2007 – 2008: Fulbright Scholarship and Yale University Scholarship



Germany

Jonas Barth

EDUCATION

College / University:

University of Applied Sciences Giessen-Friedberg, Germany

Highest Degree:

Diploma

Major Subjects:

Bioinformatics

Lab Experience:

Programming and other bioinformatic techniques, techniques in molecular biology

Projects / Research:

Wet Lab:

Mar 08 – Aug 08: Dept. of Molecular Neurogenetics at the European Neuroscience Institute Göttingen: analysis of *C.elegans* Rab family

Computational Labs:

Mar 07 – Sep 07: Centre for Molecular and Biomolecular Informatics Radboud Uni. Nijmegen (NL) "Comparison of mutation and SNP impacts on GPCRs"

Sep 06 – Feb 07: NeuroPI-group of the Department of Cognitive Neurosciences Radboud Uni. Nijmegen (NL) "Establishing an associative search for the database COCOMAC (Collations of Connectivity data on the Macaque Brain)"

Scholarships:

2008 – 2009: International Max Planck Research School support



Taiwan

I-Wen Chen

EDUCATION

College / University

National Taiwan University

Highest Degree:

B.Sc.

Major Subjects:

Life Science

Lab Experience:

Basic methods in molecular biology, developmental biology, and physiology

Projects / Research:

2006 – 2007: Cold acclimation in zebrafish gill

2005 – 2006: Ionocytes differentiation in zebrafish(embryos and adult fish gills)

Publications:

Chou MY, Hsiao CD, Chen SC, Chen IW, Liu ST, Hwang PP. Effects of hypothermia on gene expression in zebrafish gills: upregulation in differentiation and function of ionocytes as compensatory responses. *J Exp Biol.* 2008 Oct;211 (Pt 19):3077-84

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

2006 and 2007: Scholarships for undergraduate research, NSC, Taiwan

2005: Presidential Award in National Taiwan University, Taiwan



Brazil

Carolina Cunha

EDUCATION

College / University

2002 – 2007: College of Pharmacy, Federal University of Pernambuco, Brazil

Highest Degree:

B.Sc.

Major Subjects:

Pharmaceutical Sciences

Lab Experience:

Organic synthesis, analytical chemistry, pharmacological and toxicological analysis, drug formulation

Projects / Research:

Sep 07 – Jan 08: "Schizophrenia: Current Pharmacological Therapy and Future Perspectives", internship Ulysses Pernambucano Psychiatric Institute

Sep 06 – Nov 06: Internship at the College of Pharmacy, Federal University of Pernambuco, in Pharmaceutical Compounding and Drug Dispensation

Sep 04 – Aug 05: Pre-clinical toxicological research and evaluation of *Carapa guianensis* extract

Mar 03 – Apr 04: Research towards new pathways in organic synthesis with the usage of organometallic compounds

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society



Colombia

Juan Daniel Flórez Weidinger

EDUCATION

College / University

2004 – 2007: Universidad de los Andes, Bogotá, Colombia

Highest Degree:

B.Sc.

Major Subjects:

Physics

Lab Experience:

Techniques in Biophysics, GUV's, SUV's and supported lipid membrane formation, basic fluorescence imaging. *In silico* experiments with Monte Carlo simulations

Projects / Research:

Mar – Aug 2008: determining the factors involved in the variability in the rhodopsin activated signal transduction cascade in photoreceptors

Aug – Dec 2007: Thesis Project "Measurement of the diffusion coefficient in a lipid monolayer with and without domains using FRAP and Monte Carlo

Apr – May 2007: measuring the growth of lipid domains in a supported monolayer through an increase of lateral pressure using a homemade Langmuir Trough

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

Aug 04 – Dec 07: scholarship for finishing the program in 7 semesters with a GPA above 4.0/5.0



India

Srinivas Gorur Shandilya

EDUCATION

College / University

St. Stephen's College, University of Delhi, India

Highest Degree:

B.Sc. in Physical Sciences

Major Subjects:

Physics, Chemistry, and Mathematics

Lab Experience:

Elementary techniques in biochemistry, and *Drosophila* and *C. elegans* genetics

Projects / Research:

2008: Worked on pilot experiments involving behavioral assays to study if Soldier flies pay attention to visual cues while exhibiting landing behavior

2008: Using MATLAB language to find regions of similar anisotropic spectral distribution on a cell surface, indicating the location of 'rafts' on the cell surface

2007: Built an experimental protocol to screen *C. elegans* mutants for deviations in distributions of mitochondrial volume along the axon

2006: Project on the genetic basis of photophobic response in *Drosophila*

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

2005 – 2008: KVPY Fellowship, Indian Institute of Science/Government of India



P.R. China

Xiaojie Huang

EDUCATION

College / University

Tsinghua University

Highest Degree:

B.Sc.

Major Subjects:

Biological Science

Lab Experience:

Oct 06 – Jul 08: Centre for learning and memory, School of Medicine, Tsinghua University, project “The effect of a new compound on rodent to improve learning and memory”

2005 – 2006: Institute of Biomechanics and Medical Engineering, School of Aerospace, Tsinghua University, project “Mechanical mechanism of the avascular necrosis of femoral head (ANFH)”

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

2005 – 2008: Tang’s scholarship



Germany

Cordelia Imig

EDUCATION

College / University

Philipps-Universität Marburg

Highest Degree:

B.Sc. in Biology

Major Subjects:

Cell Biology and Genetics, Animal Physiology, Neurobiology and Behavioral Physiology

Lab Experience:

Molecular biology techniques, DNA cloning techniques, *in situ* hybridization, MALDI-TOF mass spectrometry, immunofluorescence stainings, confocal microscopy

Projects / Research:

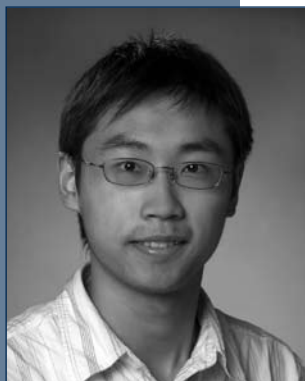
2008: “Anatomical and histological processes during brain development and about the role of signalling molecules in patterning processes during the development of the cerebral cortex”, University of Edinburgh

2007/08: “The role of *amontillado* (PC2) in neuropeptide processing in *Drosophila*”, Philipps-Universität Marburg

Scholarships:

2008 – 2009: International Max Planck Research School support

May – Aug 2008: Erasmus Lifelong Learning Programme (LLP)



P.R. China

Zhizi Jing

EDUCATION

College / University

Sep 04 – Jul 08: College of Life Sciences, Nankai University

Highest Degree:

B.Sc.

Major Subjects:

Molecular Biology, Cell Biology, Biochemistry

Lab Experience:

Jul 07 – Jul 08: Lab of Tumor Immunology and Molecular Biology, Dept. of Immunology, School of Medicine, Nankai University

Mar – Sep 2007: “100 Projects” of Creative Research for the undergraduates of Nankai University

Projects / Research:

Project 1: Study the effect of Folic Acid on the growth and development of primary cortex neurons in culture (advisor: Dr. Yanqiang Liu)

Project 2: High-throughput screen of functional gene involved in NF- κ B signaling by using a limited siRNA library. Investigate the biological function of a human unknown functional gene induced in dendrite cells by stimulation of specific receptors involved in immune response (advisor: Dr. Rongcun Yang)

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

2006 – 2007: Third Prize Excellent UG Scholarship of Nankai University



Greece

Maria-Eleni Kastriti

EDUCATION

College / University

School of Natural Sciences, University of Patras

Highest Degree:

B.Sc. in Biology

Major Subjects:

Genetics, Cellular-Molecular Biology, Physiology

Lab Experience:

SDS-PAGE electrophoresis, flow cytometry, isolation of white cells

Projects / Research:

Identification of p-Elk-1 in hemocytes of *C. capitata*, estimation of the effect of the plasma factors of people who suffer from chronic kidney diseases on their white cells phagocytic activity

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society



Germany

Alejandro Mendoza Schulz

EDUCATION

College / University

Universität Duisburg Essen

Highest Degree:

B.Sc. in "Water Science"

Major Subjects:

Analytics, Microbiology

Lab Experience:

Chemical analytics, organic synthesis, cell culture, western blot, other basic techniques in molecular biology and biochemistry

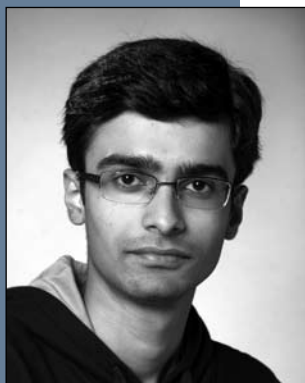
Projects / Research:

2007: The effect of fluoride on cell viability, cell proliferation and cell migration of the anterior pituitary cells: GH4C1 and Follicle Stellate (CINVESTAV, Mexico D.F.)

Scholarships:

2008 – 2009: International Max Planck Research School support

Apr – Aug 2007: "Stipend for a temporary stay abroad with the purpose of writing a final paper" granted by the German Academic Exchange Service



India

Jatin Nagpal

EDUCATION

College / University

Sri Venkateswara College, University of Delhi

Highest Degree:

B.Sc. (Honors) in Biochemistry

Major Subjects:

Biochemistry, Molecular and Cell Biology, Immunology, Membrane Biology, Bioenergetics, Genetics

Lab Experience:

Molecular biology, protein purification and gene cloning techniques, behavioral and molecular study of circadian rhythms in *Drosophila*

Projects / Research:

May – Jun 2008: To investigate the role of glial cells in regulating the circadian behavioral phenotype in *Drosophila melanogaster*

Sep – Oct 2007: To clone the gene Rv0903c which encodes for a two component response regulator involved in signal transduction in *Mycobacterium tuberculosis*

May – Jun 2007: To ablate LNvs using Gal4-UAS lines and studying its effect on the eclosion activity in *Drosophila melanogaster*

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

2006 – 2008: 'Project Oriented Biological Education' science outreach program of JNCASR

2005 – 2008: CBSE merit scholarship for undergraduates in basic sciences



India

Srinivas Parthasarathy

EDUCATION

College / University

Sri Venkateswaran College, University of Delhi

Highest Degree:

B.Sc. in Biochemistry

Major Subjects:

Biochemistry, Molecular and Cell Biology, Immunology, Membrane Biology and Bioenergetics, Genetics

Lab Experience:

Molecular Biology and Biochemistry related techniques, animal handling and behavior paradigms including anxiety tests

Projects / Research:

Sep – Dec 2007: Cloning and characterization of the signal transduction genes of the *M.tuberculosis* two component system

Jun – Jul 2007: Role of aminoguanidine in ameliorating the temporary memory impairment caused by acute Hypobaric Hypoxia in mice

May – Jun 2007: Over expression of *M.tuberculosis* protein Rv2027 in *E.coli* and purification using Ni-NTA column

Scholarships:

2008 – 2009: Stipend International Max Planck Research School, Germany

2005 – 2008: CBSE merit scholarship for outstanding performance in the All India Senior Secondary Certificate Examination (Class XII)



Colombia

Natalia Revelo Nuncira

EDUCATION

College / University

2003 – 2007: Universidad Nacional de Colombia, Bogotá

Highest Degree:

B.Sc. in Biology (Honor degree)

Major Subjects:

Animal physiology, Cell Biology, Neurobiology

Lab Experience:

Primary and tridimensional cultures of sciatic nerve and dorsal root ganglion cells. Immunohistochemistry. Stereotaxic surgery.

Projects / Research:

2008 Content based recovery system for histological images

2008 Thyroperoxidase location in mouse thyroid follicles

2007 Mouse sciatic nerve Schwann cells and endoneural fibroblast tridimensional cultures in a collagen matrix

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

2008: Young and innovative researchers program, COLCIENCIAS, Colombia

2007: Study Visits / Seminars and Practicals in Germany (DAAD)

2003 – 2007: Honor matriculation for best grade point average



Germany

Meike Annika Schweisfurth

EDUCATION

College / University

2007 – 2008: Darwin College , University of Cambridge

2004 – 2007: University of Göttingen

Highest Degree:

Certificate of Advanced Studies of Mathematics (CASM)

Major Subjects:

Mathematics and Theoretical Physics

Projects / Research:

2008: Cambridge Essay about Rayleigh-Bénard convection, a problem in hydrodynamic systems

Scholarships:

2008 – 2009: International Max Planck Research School support

2007 – 2008: Cambridge European Trust, Hölderlin-Stipendium der Allianz, DAAD Europäisches Exzellenzprogramm

since 2005: Studienstiftung des deutschen Volkes

since 2004: e-fellows.net-online scholarship



France

Nicolas Snaidero

EDUCATION

College / University

University Joseph Fourier, Grenoble, France and University of Glasgow, Scotland

Highest Degree:

B.Sc. Neurosciences, Maitrise science and technology mention biology

Major Subjects:

Neurosciences

Lab Experience:

Animal experimentation, identification and extraction of CNS regions in mice, DNA and ARN extraction and quantification, protein assay, RT PCR, Western Blot, immunohistochemistry fluorescence and confocal, bacterial culture and genetical analysis of *E. coli*

Projects/Research:

Jun – Sep 2008: Validation of gene expression patterns in genetic models of spastic paraplegia, Max Planck Institute for Experimental Medicine, Göttingen

Oct 07 – May 08: Biochemical investigation of axonal transport regulators in myelin mutant mice, Applied Neurobiology Group in the Faculty of Veterinary Medicine, University of Glasgow, UK

Jun – Sep 2006: Analysed networks of the genetic expression in *E. coli* bacteria, Laboratoire Adaptation et Pathogénie des Microorganismes CNRS, La Tronche, France

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society



India

Swathi Srivatsa

EDUCATION

College / University

Mount Carmel College, Bangalore University

Highest Degree:

B.Sc.

Major Subjects:

Chemistry, Zoology, Microbiology

Lab Experience:

Molecular biological techniques (cloning, DNA sequencing, PCR) protein chemistry, Western Blotting, working with animal cell lines for transfection/gene expression, flow cytometry, immune- techniques (Latex agglutination test) Counter Current Immunoelectrophoresis, ELISA, Radial and Double Diffusion assay, chromatography, spectroscopy

Projects/Research:

Developing novel HIV-1 reporter vectors with transactivation control, construction of a dual reporter expression vector under modified C-LTR promoter, construction of reporter gene expression inversely correlated to Tat protein expression

Scholarships:

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

2006 – 2008: Project Oriented Biological Education Fellowship, Jawaharlal Nehru Centre for Advanced Scientific Research, and Kishore Vaigyanik Prothshahan Yojna Fellowship, Indian Institute of Science, Bangalore

Apr 2006: National Initiative on Undergraduate Science Fellowship, Homi Babha Centre of Science Education, Mumbai



Germany

Roman Stilling

EDUCATION

College / University

Westfälische Wilhelms-Universität Münster

Highest Degree:

B.Sc.

Major Subjects:

Biology

Lab Experience:

Basic techniques in molecular biology and genetics

Projects / Research:

2008: The role of *Psc* in olfactory neuron differentiation of *Drosophila melanogaster*

Scholarships:

2008 – 2009: International Max Planck Research School support



Germany

Benjamin Wilhelm

EDUCATION

College / University

Oct 05 – Sep 08 Georg-August University Göttingen

Highest Degree:

Vordiplom

Major Subjects:

Biology

Lab Experience:

Basic techniques in molecular biology, microbiology, biochemistry, cell culture, and fluorescence microscopy

Projects / Research:

Aug 06 – Jul 08: Research assistant at the Max Planck Institute for Biophysical Chemistry (Dept. of Membrane Biophysics) investigating in the processing of APP

Scholarships:

2008 – 2009: International Max Planck Research School support

Apr 2008 – today: online stipend from e-fellows.net

Aug 2007: Travel stipend from the German Academic Exchange Service



China

Aaron Benson Wong

EDUCATION

College / University

2004 – 2008: The Chinese University of Hong Kong

2006 – 2007: Exchange Program to University of California, Riverside

Highest Degree:

B.Sc. with Honors, First Class

Major Subjects:

Molecular Biotechnology

Lab Experience:

Molecular cloning, bacterial protein expression, RT-PCR, 2D-GE, mammalian cell culture, drug screening through bioassays

Projects / Research

Jul 2007 – May 2008: Screening of herbal materials for promoting neural cell growth and differentiation

Jun 2006 – Aug 2006: Cloning, expression and purification of Bee Venom Protein

Scholarships

2008 – 2009: Stipend of the Excellence Foundation for the Promotion of the Max Planck Society

2007 – 2008: Chung Chi College Departmental Prize – Molecular Biotechnology (Chung Chi College, CUHK) and Kong E Suen Memorial Scholarship (Faculty of Science, CUHK)

2004 – 2008 Fr. Barrett Memorial Scholarship for Undergraduate Studies

Faculty

Name		Institute	
Mathias	Bähr	Neurology	U Göttingen
Nils	Brose	Molecular Neurobiology	MPI em
Wolfgang	Brück	Neuropathology	U Göttingen
Edgar	Brunner	Medical Statistics	U Göttingen
Hannelore	Ehrenreich	Clinical Neurosciences	MPI em
Stefan	Eimer	Molecular Neurogenetics	ENI
Wolfgang	Engel	Human Genetics	U Göttingen
André	Fiala	Molecular Neurobiology of Behavior	U Göttingen
André	Fischer	Laboratory for Aging and Cognitive Diseases	ENI
Gabriele	Flügge	Neurobiology	DPZ
Jens	Frahm	Biomedical NMR Research / Physical Chemistry	MPI bpc
Eberhard	Fuchs	Animal Physiology / Neurobiology	DPZ
Theo	Geisel	Nonlinear Dynamics	MPI ds
Martin	Göpfert	Cellular Neurobiology	U Göttingen
Uwe-Karsten	Hanisch	Neuropathology	U Göttingen
Ralf	Heinrich	Neurobiology	U Göttingen
Michael	Hörner	Cell Biology/Neurobiology	U Göttingen
Swen	Hülsmann	Neuro- and Sensory Physiology	U Göttingen
Reinhard	Jahn	Neurobiology	MPI bpc
Hubertus	Jarry	Clinical and Experimental Endocrinology	U Göttingen
Jürgen	Klingauf	Membrane Biophysics	MPI bpc
Till	Marquardt	Developmental Neurobiology	ENI
Tobias	Moser	Otolaryngology	U Göttingen
Klaus-Armin	Nave	Neurogenetics	MPI em
Erwin	Neher	Membrane Biophysics	MPI bpc
Walter	Paulus	Clinical Neurophysiology	U Göttingen
Diethelm W.	Richter	Neuro- and Sensory Physiology	U Göttingen
Michael	Rickmann	Neuroanatomy	U Göttingen
Silvio O.	Rizzoli	STED Microscopy of Synaptic Function	ENI
Detlev	Schild	Molecular Neurophysiology	U Göttingen
Oliver	Schlüter	Molecular Neurobiology	ENI
Jörg B.	Schulz	Neurodegeneration	U Göttingen
Mikael	Simons	Biochemistry and Molecular Cell Biology	MPI em
Judith	Stegmüller	Cellular and Molecular Neurobiology	MPI em
Nicole	von Steinbüchel	Medical Psychology and Medical Sociology	U Göttingen
Anastassia	Stoykova	Molecular Cell Biology	MPI bpc
Walter	Stühmer	Molecular Biology of Neuronal Signals	MPI em
Andreas	Stumpner	Neurobiology	U Göttingen
Victor	Tarabykin	Molecular Biology of Neuronal Signals	MPI em
Stefan	Treue	Cognitive Neuroscience and Biological Psychology	DPZ
Andreas	Wodarz	Stem Cell Biology	U Göttingen
Fred	Wolf	Nonlinear Dynamics	MPI ds
Fred	Wouters	Cellular Biophysics	U Göttingen
Weiqi	Zhang	Neuro- and Sensory Physiology	U Göttingen

U Göttingen = Georg August University, MPI bpc = Max Planck Institute for Biophysical Chemistry, MPI em = Max Planck Institute for Experimental Medicine, MPI ds= Max Planck Institute for Dynamics and Self-Organization, DPZ = German Primate Center, ENI = European Neuroscience Institute



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Mathias Bähr

Professor of Neurology

- 1985 MD, University of Tübingen Medical School, Training in Neurology at University Hospitals in Tübingen and Düsseldorf
- DFG and Max Planck Fellow at the Max Planck Institute for Developmental Biology Tübingen and at the Department of Anatomy and Cell Biology, Washington University St.Louis
- Schilling-Foundation Professor for Clinical and Experimental Neurology, University of Tübingen
- Director at the Department of Neurology, University of Göttingen since 2001

Major Research Interests

We are interested to understand 2 basic questions in cellular and molecular neurobiology:

1. Which factors support survival of adult CNS neurons?
2. What kills these cells under pathological conditions?

Up to now, only little is known about the mechanisms that support survival of a postmitotic cell like a human neuron for eventually more than 100 years under physiological conditions. However, by examining the molecular regulation of cell survival and cell death during development and in the lesioned adult CNS, one may get some clues to answer this question.

In our group, several *in vitro* and *in vivo* model systems are used which allow examination of neuronal de- and regeneration. Our basic model is the rodent retino-tectal projection. Here, we can study development, de- and regeneration of the respective projection neurons, the retinal ganglion cells (RGCs) in single cell cultures, explants or *in vivo*. Transection or crush-axotomy of the optic nerve induces retrograde death more than 80% of RGCs within two weeks. This secondary cell loss is mainly apoptotic and involves specific changes in gene expression pattern of transcription factors (e.g. c-jun or ATF-2), pro- and anti-apoptotic genes (e.g. bcl-2 or bax) and growth-associated genes (like GAP-43). Thus, long term survival and initiation of regeneration programmes of RGCs critically depends on inhibition of apoptotic cell death. To that end, we have used a variety of techniques to interfere with the cell death cascades that follow lesions of the optic nerve in adult rats. Inhibition of neuronal apoptosis can be afforded by pharmacological administration of trophic factors or by gene therapy approaches using adeno- or adeno-associated virus vectors that can deliver neurotrophic or anti-apoptotic factors directly into neurons or into surrounding glial cells. These, and other new strategies like using peptide-transduction-domains to deliver anti-apoptotic proteins across the blood-brain-barrier are now used to develop new experimental therapy strategies in animal models of human neurological disorders like stroke, trauma, multiple sclerosis or neurodegenerative diseases (e.g. Alzheimer's or Parkinson's disease).

Selected Recent Publications

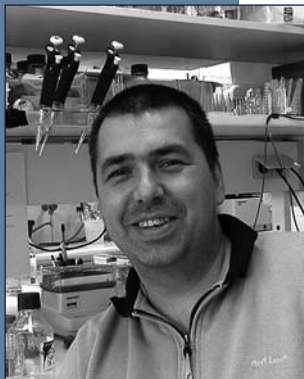
Lingor P, Tönges L, Pieper N, Bermel C, Barski E, Planchamp V, Bähr M (2008) ROCK inhibition and CNTF interact on intrinsic signalling pathways and differentially regulate survival and regeneration in retinal ganglion cells. *Brain* 131 (Pt 1): 250-63

Ganesan S, Rohde G, Eckermann K, Sroka K, Schaefer MKE, Dohm CP, Kermer P, Haase G, Wouters F, Bähr M, Weishaupt JH (2008) Mutant SOD1 detoxification mechanisms in intact single cells. *Cell Death & Diff.* 15(2): 312-21

Koeberle PD, Bähr M (2007) The Upregulation of GLAST-1 is an indirect anti-apoptotic mechanism of GDNF and Neurturin in the adult CNS. *Cell Death & Diff.* 15(3): 471-83

Meuer CK, Suppanz IE, Lingor P, Planchamp V, Göricke B, Fichtner L, Braus GH, Dietz GPH, Jakobs S, Bähr M, Weishaupt JH (2007) Cyclin-dependent kinase 5 is an upstream regulator of mitochondrial fission during neuronal apoptosis. *Cell Death & Diff.* 14(4): 651-61

Liman J, Ganesan S, Dohm CP, Krajewski S, Reed JC, Bähr M, Wouters F, Kermer P (2005) Interaction of BAG1 and Hsp70 mediates neuroprotectivity and increases chaperone activity. *Mol Cell Biol.* 25(9): 3715-25



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Nils Brose

Professor, Director at the Max Planck Institute for Experimental Medicine

- Dr. rer. nat. (Ph.D.) 1990, Ludwig Maximilians University Munich
- Appointed as Director at the Max Planck Institute for Experimental Medicine 2001

Major Research Interests

Research in the Department of Molecular Neurobiology focuses on the molecular mechanisms of synapse formation and function in the vertebrate central nervous system. Typically, synapses are formed between cellular processes of a sending and a receiving nerve cell. They are the central information processing units in the vertebrate brain where some 10¹² nerve cells are connected by 10¹⁵ synapses to form an elaborate and highly structured neuronal network that is the basis for all forms of behaviour. Signal transmission at synapses is mediated by the regulated release of signal molecules (neurotransmitters) which then diffuse to the receiving nerve cell and change its physiological state. In the Department of Molecular Neurobiology, we combine biochemical, morphological, mouse genetic, behavioural, and physiological methods to elucidate the molecular basis of synapse formation and transmitter release processes. Our synaptogenesis research concentrates on synaptic cell adhesion proteins and their role in synapse formation. Studies on the molecular mechanisms of neurotransmitter release focus on components of the presynaptic active zone and their regulatory function in synaptic vesicle fusion.

Selected Recent Publications

Jamain S, Radyushkin K, Hammerschmidt K, Granon S, Boretius S, Varoqueaux F, Ramanantsoa N, Gallego J, Ronnenberg A, Winter D, Frahm J, Fischer J, Bourgeron T, Ehrenreich H, Brose N (2008) Reduced social interaction and ultrasonic communication in a mouse model of monogenic heritable autism. *Proc Natl Acad Sci USA* 105: 1710-1715

Jockusch W, Speidel D, Sigler A, Sørensen J, Varoqueaux F, Rhee J-S, Brose N (2007) CAPS-1 and CAPS-2 are essential synaptic vesicle priming proteins. *Cell* 131: 796-808

Varoqueaux F, Aramuni G, Rawson RL, Mohrmann R, Missler M, Gottmann K, Zhang W, Südhof TC, Brose N (2006) Neuroligins determine synapse maturation and function. *Neuron* 51: 741-754

Reim K, Wegmeyer H, Brandstätter JH, Xue M, Rosenmund C, Dresbach T, Hofmann K, Brose N (2005) Structurally and functionally unique Complexins at retinal ribbon synapses. *J Cell Biol* 169: 669-680

Junge H, Rhee J-S, Jahn O, Varoqueaux F, Spiess J, Waxham MN, Rosenmund C, Brose N (2004) Calmodulin and Munc13 form a Ca²⁺-sensor/effector complex that controls short-term synaptic plasticity. *Cell* 118: 389-401

Rhee J-S, Betz A, Pyott S, Reim K, Varoqueaux F, Augustin I, Hesse D, Südhof TC, Takahashi M, Rosenmund C, Brose N (2002) b Phorbol ester- and diacylglycerol-induced augmentation of transmitter release is mediated by Munc13s and not by PKCs. *Cell* 108: 121-133



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Wolfgang Brück

Professor of Neuropathology

- 1986 MD Johannes Gutenberg University in Mainz, 1994 national boards in neuropathology
- 1996-2002 Associate professorships for neuropathology at the University of Göttingen and the Charité in Berlin
- Since 2002 full professor and director of the Department of Neuropathology, University of Göttingen

Major Research Interests

- Immunopathology of multiple sclerosis
- Brain-specific mechanisms of immune response in multiple sclerosis
- Axonal damage in inflammatory demyelination and mechanisms of remyelination
- Mechanisms and consequences of microglial activation

Selected Recent Publications

Kuhlmann T, Remington L, Maruschak B, Owens T, Brück W (2007) Nogo-A is a reliable oligodendroglial marker in human and mouse adult CNS as well as in demyelinated lesions. *J Neuropathol Exp Neurol* 66: 238-246

Albert M, Antel J, Brück W, Stadelmann C (2007) Extensive cortical remyelination in patients with chronic multiple sclerosis. *Brain Pathol* 17: 129-138

Metz I, Lucchinetti CF, Openshaw H, Garcia-Merino A, Lassmann H, Freedman MS, Azzarelli B, Kolar OJ, Atkins HL, Brück W (2007) Autologous hematopoietic stem cell transplantation fails to stop demyelination and neurodegeneration in multiple sclerosis. *Brain* 130: 1254-1262

Jack C, Antel J, Brück W, Kuhlmann T (2007) Contrasting potential of nitric oxide and peroxynitrite to mediate oligodendrocyte injury in multiple sclerosis. *Glia* 55: 926-934

Schwartz M, Butovsky O, Brück W, Hanisch UK (2006) Microglial phenotype: Is the commitment reversible? *Trends Neurosci* 29: 68-74

Merkler D, Ernsting T, Kerschensteiner M, Brück W*, Stadelmann C* (2006) A new focal EAE model of cortical demyelination: MS-like lesions with rapid resolution of inflammation and extensive remyelination. *Brain* 129: 1972-1983

Patrikios P, Stadelmann C, Kutzelnigg A, Rauschka H, Schmidbauer M, Laursen H, Sorensen P, Brück W, Lucchinetti C, Lassmann H (2006) Remyelination is extensive in a subset of Multiple Sclerosis patients. *Brain* 129: 3165-3172

Zhou D, Srivastava R, Nessler S, Grummel V, Sommer N, Brück W, Hartung HP, Stadelmann C, Hemmer B (2006) Identification of a Pathogenic Antibody Response to Native Myelin Oligodendrocyte Glycoprotein in Multiple Sclerosis. *PNAS* 103: 19057-19062

Gutenberg A, Buslei R, Fahlbusch R, Buchfelder M, Brück W (2005) Immunopathology of primary hypophysitis: implications for pathogenesis. *Am J Surg Pathol* 29: 329-38

Keegan M, König F, McClelland R, Brück W, Morales Y, Bitsch A, Panitch H, Lassmann H, Weinshenker B, Rodriguez M, Parisi J, Lucchinetti CF (2005) Humoral Multiple Sclerosis Pathology Correlates With Response To Therapeutic Plasma Exchange. *The Lancet* 366: 579-582

Merkler D, Boretius S, Stadelmann C, Ernsting T, Michaelis T, Frahm J, Brück W (2005) Multicontrast MRI of remyelination in the central nervous system. *NMR Biomed* 18: 395-403



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Edgar Brunner

Professor of Medical Statistics

- Student: WS 64/65 - SS 69, Technical University of Aachen
- Diploma: April 1969, Mathematics
- Promotion: 12. May 1971, (Dr. rer. nat.), Technical University of Aachen
Title: Eine Beziehung zwischen dem Holm-Test und dem Kolmogorov-Smirnov-Test (A Relation between Holm's Test and the Kolmogorov-Smirnov-Test)
- Habilitation: 11.11.1973, Medical Statistics
- Professor: 01.01.1976 University of Göttingen, Dept. of Medical Statistics, 01.03.1976 Head of the Department
- Editor: Biometrical Journal
- Associate Editor: Journal of Statistical Planning and Inference

Major Research Interests

Nonparametric Statistics

- Asymptotic distribution of rank statistics
- Multi-factor designs
- Adjustment for covariates

Longitudinal data

Ordered categorical data

Design and analysis of diagnostic trials

Statistical methods for the analysis of microarray data

Analysis of high-dimensional data

Selected Recent Publications

Thangavelu K, Brunner E (2006) Wilcoxon Mann-Whitney Test for Stratified Samples and Efron's Paradox Dice. *Journal of Statistical Planning and Inference* (in press)

Werner C, Brunner E (2006) Rank methods for the analysis of clustered data in diagnostic trials. *Computational Statistics and Data Analysis* (in press)

Bretz F, Landgrebe J, Brunner E, (2006) Efficient Design and Analysis of Two Color Factorial Microarray Experiments. *Computational Statistics and Data Analysis* 50: 499-517

Chen T-W, Lin B-J, Brunner E, Schild D, (2006) *In Situ* Background Estimation in Quantitative Fluorescence Imaging. *Biophysical Journal* 90: 2534-2547

Kaufmann J, Werner C, Brunner E, (2005) Nonparametric methods for analyzing the accuracy of diagnostic tests with multiple readers. *Statistical Methods in Medical Research* 14: 129-146

Brunner E, Domhof S, Langer F (2002) *Nonparametric Analysis of Longitudinal Data in Factorial Designs*. Wiley: New York

Brunner E, Munzel U (2002) *Nichtparametrische Datenanalyse*. Springer. Heidelberg

Brunner E, Munzel U, Puri ML (2001) The multivariate nonparametric Behrens-Fisher-Problem. *J. Statist Plann and Inf* 108: 37-53

Brunner E, Munzel U, Puri ML (1999) Rank-Score Tests in Factorial Designs with Repeated Measures. *Journal of Multivariate Analysis* 70: 286-317

Akritas MG, Arnold SF, Brunner E (1997) Nonparametric hypotheses and rank statistics for unbalanced designs. *Journal of the American Statistical Association* 92: 258-265



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Hannelore Ehrenreich

Professor of Neurology and Psychiatry

- 1981 Doctor of veterinary medicine, University of Munich
- 1983 Elective Period, University of Newcastle-upon-Tyne, England
- 1985 Guest Lecturer, University of the Philippines, Manila
- 1985 - 1986 Assistant, Department of Internal Medicine, University of Munich
- 1987 Graduation (Medicine), University of Munich
- 1987 - 1988 Assistant, Department of Neurology, University of Munich
- 1989 Doctor of Medicine, University of Munich
- 1989 - 1991 Guest Scientist (BMBF grant) NIAID, NIH, Bethesda, MD, USA
- 1992 - 1994 Assistant, Departments of Neurology and Psychiatry, University of Göttingen
- 1994 Habilitation (Neurology and Psychiatry)
- 1994 - present Head, Division of Clinical Neuroscience, MPIEM
- 1995 - present Consultant & Professor (1998) of Neurology & Psychiatry, University of Göttingen
- 2000 - 2002 Vice President, University of Göttingen

Major Research Interests

Translational Neuroscience

- (1) Molecular-cellular basis of neuropsychiatric disease with focus on endogenous mechanisms of neuroprotection
- (2) Clinical research on neuroprotection and neuroregeneration in acute (ischemia/hypoxia, trauma) and chronic brain disease (schizophrenia, autism, ALS, MS)
- (3) Clinical addiction research

Novel concepts for treatment of alcoholism, psychotherapeutic process-outcome research including kinetics and mechanisms of regeneration

Selected Recent Publications

Ehrenreich H, Hinze-Selch D, Stawicki S, Knolle-Veentjer S, Aust C, Wilms S, Heinz G, Erdag S, Jahn H, Degner D, Ritzen M, Mohr A, Knauth M, Wagner M, Schneider U, Bohn M, Huber M, Czernik A, Pollmächer M, Maier W, Sirén A-L, Klosterkötter J, Falkai P, Rütther E, Aldenhoff JB, Krampe H: Improvement of cognitive functions in chronic schizophrenic patients by recombinant human erythropoietin. *Mol Psychiatry* in press

Sirén A-L, Radyushkin K, Boretius S, Kämmer D, Riechers C-C, Natt O, Sargin D, Watanabe T, Sperling S, Michaelis T, Price J, Meyer B, Frahm J Ehrenreich H (2006) Global brain atrophy after unilateral parietal lesion and its prevention by erythropoietin. *Brain* 129: 480-489

Ehrenreich H, Hasselblatt M, Knerlich F, von Ahsen N, Jacob S, Sperling S, Woldt H, Nave KA, Sirén AL (2005) A hematopoietic growth factor, thrombopoietin, has a pro-apoptotic role in the brain. *Proc Natl Acad Sci USA* 102: 862-7

Ehrenreich, H, Degner, D, Meller, J, Brines, M, Béhé, M, Hasselblatt, M, Woldt, H, Falkai, M, Knerlich, F, Jacob, S., Maier, W, Brück, W, Rütther, E, Cerami, A, Becker, W, Sirén, A-L (2004) Erythropoietin: A candidate for neuroprotection in schizophrenia. *Mol Psychiatry* 9: 42-54

Sirén A-L, Fratelli M, Brines M, Goemans C, Casagrande S, Lewczuk P, Keenan S, Gleiter C, Pasquali C, Capobianco A, Mennini T, Heumann R, Cerami A, Ehrenreich H, Ghezzi P (2001): Erythropoietin prevents neuronal apoptosis after cerebral ischemia and in metabolic stress. *Proc Natl Acad Sci USA* 98: 4044-4049



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Stefan Eimer

Group Leader Molecular Neurogenetics / Neurodegeneration

- Ph.D. 2003 at the Gene Center of the Ludwig-Maximilian University (LMU in Munich)
- 2003 Postdoc at the Ecole Normale Supérieure in Paris, France
- since Oct 2005 independent group leader of the Center for Molecular Physiology of the Brain (CMPB) at the European Neuroscience Institute (ENI) in Göttingen

Major Research Interests

Neurotransmitter gated ion channels are involved in a large subset of neuronal events ranging from fast synaptic transmission to the modulation of neuronal circuits that lead to memory formation and cognition. En route to the cell surface these multimeric receptors have to undergo multiple assembly, quality control, and sorting steps to eventually reach the synapse.

Our group aims to understand the mechanisms and rules that control the trafficking and sorting of ligand gated ion channels within the secretory apparatus. In particular, we are focusing on the nicotinic acetylcholine receptor family of ligand gated ion channels, which have been implicated in numerous neurological and neurodegenerative diseases.

To find new molecules involved in these processes, we take advantage of the nematode *Caenorhabditis elegans* as a main model system, and use a combination of genetic, cell biological, and biochemical approaches as well as electro-physiology and electron-microscopy. As our main model system we are studying cholinergic neurotransmission at the neuro-muscular junction (NMJ) of *C. elegans*. Through genetic screens we have identified novel evolutionary conserved integral membrane proteins that regulate nAChR sorting at the Golgi-Endosomal interface. Further studies have implicated these molecules in the regulation and activation of small GTPases at Golgi complex. Based on these findings we have also started to study systematically how these GTPases are required for structure and function of the Golgi apparatus and how their activity affects the trafficking and neurotransmission at the NMJ of *C. elegans*.

Selected Recent Publications

Marza E, Long T, Saiardi A, Sumakovic M, Eimer S, Hall DH, Lesa GM (2007) Polyunsaturated fatty acids influence synaptojanin localization to regulate synaptic vesicle recycling. *Mol Biol Cell*, in press

Eimer S, Gottschalk A, Richmond JE, Hengartner M, Schafer W, Bessereau J-L (2007) Regulation of nicotinic receptor trafficking by the transmembrane Golgi protein UNC-50. *EMBO J* 26: 4313-23

Yamasaki A, Eimer S, Okochi M, Smialowska A, Kaether C, Baumeister R, Haass C, Steiner H (2006) The GxGD motif of presenilin contributes to catalytic function and substrate identification of gamma-secretase. *J Neurosci* 26: 3821-8

Gally C, Eimer S, Richmond JE, Bessereau J-L (2004) A transmembrane protein required for acetylcholine receptor clustering in *C. elegans*. *Nature* 431: 578-582

Eimer S, Lakowski B, Donhauser R, Baumeister R (2002) Loss of spr-5 bypasses the requirement for the presenilin sel-12 by stage-specific derepression of hop-1. *EMBO J* 21: 5787-5796



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Wolfgang Engel

Professor of Human Genetics

- Dr. med., Universität Freiburg, 1967
- Physician, Hospital Schorndorf, 1966 - 1968
- Postdoc, Institute of Human Genetics and Anthropology, Universität Freiburg, 1968 - 1977
- Habilitation (Human Genetics), Universität Freiburg, 1974
- Professor of Human Genetics and Director of the Institute, Universität Göttingen, 1977

Major Research Interests

Our research is focussed on the molecular analysis of normal human variability and genetic disturbances of development and differentiation.

Isolated genes are being analysed in detail with respect to their functional properties by animal models (transgenic and knock-out-mice). For suitable genetic diseases therapeutic strategies (substitution; gene therapy) are being developed and initial evaluation of such strategies is done in the mouse. - We are working on the genotype – phenotype correlations in neurological and cardiovascular diseases (e. g. Spastic paraplegia, Rett syndrome, mental retardation by subtelomeric microdeletions, molybdenum cofactor deficiency; cardiomyopathies, Noonan syndrome) and several genetically determined malformation syndromes (e.g. Townes-Brocks syndrome, Okihiro syndrome, Morbus Osler). We are also engaged in the molecular and cellular basis of initiation events of cancer, specifically in prostate cancer, medulloblastoma and rhabdomyosarcoma. - One main interest in our institute is the analysis of structure, expression and function of genes involved in differentiation of male gametes. The knowledge of the function of those genes can help us to clarify the genetic causes of male infertility.

We have isolated spermatogonial stem cells (SSCs) from adult mouse testis and demonstrated that these cells are as pluripotent as embryonic stem cells (ESCs). Our main interest is now to isolate and proliferate SSCs from adult human testis. These cells would be of great interest for regenerative medicine.

Selected Recent Publications

Nayerniaa K, Lee JH, Drusenheimer N, Nolte J, Wulf G, Dressel R, Gromoll J, Engel W (2006) Derivation of male germ cells from bone marrow stem cells. *Laboratory Investigation* 86: 654-663

Nayernia K, Nolte J, Michelmann HW, Lee JH, Rathsack K, Drusenheimer N, Dev A, Wulf G, Ehrmann IE, Elliott DJ, Okpanyi V, Zechner, Haaf T, Meinhardt A, Engel W (2006) *In vitro*-differentiated embryonic stem cells give rise to male gametes that can generate offspring mice. *Developmental Cell* 11: 125-132

Lee JH, Engel W, Nayernia K (2006) Stem cell protein Piwil2 modulates expression of murine spermatogonial stem cell expressed genes. *Molecular Reproduction and Development* 73: 173-179

Guan K, Nayernia K, Maier LS, Wagner S, Dressel R, Lee JH, Nolte J, Wolf, F, Li M, Engel W, Hasenfuß G (2006) Pluripotency of spermatogonial stem cells from adult mouse testis. *Nature* 440, 1199-1203

Lee HJ, Göring W, Ochs M, Mühlfeld C, Steding G, Paprotta I, Engel W, Adham IM (2004) Sox 15 is required for skeletal muscle regeneration. *Molecular and Cellular Biology* 19: 8428-8436

Nayernia K, Li, M, Jaroszynski L, Khusainow R, Wulf G, Schwandt I, Korbiowska M, Michelmann HW, Meinhardt A, Engel W (2004) Stem cells based therapeutic approach of male infertility by teratocarcinoma derived germ cells. *Human Molecular Genetics* 13: 1451-1460



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André Fiala

Professor of Molecular Neurobiology of Behavior

- 2008 Professor of Molecular Neurobiology of Behavior, University of Göttingen
- 2008 Habilitation in Neurobiology and Genetics, University of Würzburg
- 2001-2008 Research Assistant, University of Würzburg
- 2000-2001 Research Fellow, Memorial Sloan-Kettering Cancer Center, New York
- 1996-1999 PhD student, Free University of Berlin
- 1996 Degree (Diploma) in Biology, Free University of Berlin

Major Research Interests

We study neuronal mechanisms underlying olfaction, learning and memory, and goal-directed behavior using the model organism *Drosophila melanogaster*. The fruit fly *Drosophila* offers the advantage of expressing transgenes in almost any population of its about 100.000 neurons. Transgenes used by us are, for example, fluorescent sensor proteins that allow us to monitor the spatio-temporal activity of neurons, or light-sensitive proteins by which neuronal activity can be stimulated through illumination. Using these optogenetic techniques in combination with behavioral analyses we aim at unraveling the functioning of dedicated neuronal circuits, and how these circuits contribute to organizing behavior. In addition, molecular mechanisms underlying learning and memory processes are investigated.

Selected Recent Publications

Fiala A (2007) Olfaction and olfactory learning in *Drosophila*: recent progress. *Curr Opin Neurobiol* 17: 720-6

Suh GS, Ben-Tabou de Leon S, Tanimoto H, Fiala A, Benzer S, Anderson DJ (2007) Light activation of an innate olfactory avoidance response in *Drosophila*. *Curr Biol* 17: 905-8

Schroll C, Riemensperger T, Bucher D, Ehmer J, Völler T, Erbguth K, Gerber B, Hendel T, Nagel G, Buchner E, Fiala A (2006) Light-induced activation of distinct modulatory neurons triggers appetitive or aversive learning in *Drosophila* larvae. *Curr Biol* 16: 1741-7

Riemensperger T, Völler T, Stock P, Buchner E, Fiala A (2005) Punishment prediction by dopaminergic neurons in *Drosophila*. *Curr Biol* 15: 1953-60

Fiala A, Spall T, Diegelmann S, Eisermann B, Sachse S, Devaud JM, Buchner E, Galizia CG (2002) Genetically expressed cameleon in *Drosophila melanogaster* is used to visualize olfactory information in projection neurons. *Curr Biol* 12:1877-84



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André Fischer

Group Leader Laboratory for Aging and Cognitive diseases

- 2002: Dr. rer. nat.(PhD). University Goettingen/Max Planck Institute for Experimental Medicine, Germany
- 2003 - 2006: Postdoctoral Associate in the lab of Li-Huei Tsai; Harvard Medical School, Department of Pathology, Boston, USA; Picower Center for Learning and Memory, M.I.T, Cambridge, USA
- since 2006 independent group leader at the European Neuroscience Institute (ENI) in Goettingen

Major Research Interests

Our group aims to understand the molecular mechanisms underlying learning and memory processes under physiological and pathological conditions. To this end we combine molecular, biochemical, pharmacological and behavioral approaches using mice as model organisms.

We are particularly interested to understand cognitive impairment associated with normal aging as well as the pathogenesis of mental and neurodegenerative diseases, such as anxiety disorders and Alzheimer's disease.

Using animal models we deeply aim to identify therapeutic strategies that would help to reinstate neuroplasticity, learning behavior and the retrieval of lost long-term memories in patients suffering from such devastating diseases.

Selected Recent Publications

Fischer A, Sananbenesi F, Wang XY, Dobbin M, Tsai LH Recovery of learning and memory is associated with chromatin remodeling. *Nature*, doi:10.1038/nature05772

Fischer A, Radulovic M, Schrick C, Sananbenesi F, Godovac-Zimmermann J, Radulovic J (2006) Hippocampal Mek/Erk signaling mediates extinction of contextual freezing behavior. *Neurobiology of Learning and Memory* 87: 149-58

Shu T, Tseng HC, Zhou Y, Fischer A, Stern P, Coquelle F, Reiner O, Tsai LH (2006) Doublecortin-like Kinase Controls Neurogenesis by Regulating the Mitotic Spindle. *Neuron*, 49: 25-39

Fischer A, Sananbenesi F, Pang PT, Lu B, Tsai LH (2005) Opposing roles of transient and prolonged expression of p25 in synaptic plasticity and hippocampus dependent memory. *Neuron*, 48: 825-83

Park SK, Nguyen MD, Fischer A [shared co-authorship], Affar EB, Luke M, Diefenbach B, Shi Y, Tsai LH (2005) Modulation of Dopamine Signaling by Prostate Apoptosis Response 4 via Direct Interaction with Dopamine D2 Receptor. *Cell* 122: 275-287

Fischer A, Sananbenesi F, Schrick C, Spiess J, Radulovic J (2004) Distinct roles of hippocampal protein synthesis and actin rearrangement in extinction of conditioned fear. *J Neurosci* 24: 1962-1966

Sananbenesi F, Fischer A, Schrick C, Spiess J, Radulovic J (2003) Corticotropin-releasing factor receptor 2 induces mitogen-activated protein kinase signaling in the hippocampus: A possible link between stress and fear memory. *J Neurosci* 36: 11436-11443

Fischer A, Sananbenesi F, Spiess J, Radulovic J (2003) Cdk5 in the adult non-demented brain. *Current drug targets CNS* 2: 61-72

Fischer A, Sananbenesi F, Spiess J, Radulovic J (2003) Cdk5: a novel role in learning and memory. *NeuroSignals* 12: 200-208

Fischer A, Sananbenesi F, Schrick C, Spiess J, Radulovic J (2003) Regulation of contextual fear conditioning by baseline and inducible septo-hippocampal cyclin-dependent kinase 5. *Neuropharmacology* 44: 1089-1099

Sananbenesi F, Fischer A [shared first-authorship], Schrick C, Spiess J, Radulovic J (2002) Phosphorylation of hippocampal Erk-1/2, Elk-1, and p90-Rsk-1 during contextual fear conditioning: interactions between Erk-1/2 and Elk-1. *Mol Cell Neurosci* 3: 463-476



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Gabriele Flügge

Apl. Professor, Experimental Neuroscience

- Dr. rer. nat., University of Munich, 1979
- Senior Scientist, Clinical Neurobiology Laboratory at the German Primate Center

Major Research Interests

In humans, stressful or traumatic life events such as death of a close relative often represent a strong psychological load that may induce psychopathologies such as depression. The central nervous mechanisms that lead to such diseases are still not clear. We therefore investigate processes that occur in the course of chronic psychosocial stress in the brains of animals that show similar symptoms as depressed patients. Using molecular techniques, we identify central nervous genes that are regulated by stress; quantitative real time PCR, in situ hybridization and immunocytochemistry serve to localize changes in neurotransmitter systems, receptors, transporters and other molecules in distinct neurons of the brain. Similar tools are used to clarify the mechanisms that underlie the beneficial effects of antidepressant drugs. In conjunction with behavioral studies we are able to find molecular factors that play a role in central nervous processes underlying depression.

Selected Recent Publications

Abumaria N, Rygula R, Hiemke C, Fuchs E, Havemann-Reinecke U, Rütther E, Flügge G (2007). Effect of chronic citalopram on serotonin-related and stress-upregulated genes in the dorsal raphe nucleus of the rat. *Eur Neuropsychopharm* 17: 417-429

Perez-Cruz C, Muller-Keuker JIH, Heilbronner U, Fuchs E, Flügge G (2007) Morphology of pyramidal neurons in the rat prefrontal cortex: lateralized dendritic remodeling by chronic stress. *Neural Plasticity*, Vol. 2007, article ID 46276; 14 pages

Rygula R, Abumaria N, Flügge G, Hiemke C, Fuchs E, Rütther E, Havemann-Reinecke U (2006) Citalopram counteracts depressive symptoms evoked by chronic social stress in rats. *Behav Pharm* 17: 19-29

Alfonso J, Fernandez M, Cooper B, Flügge G, Frasch AC (2005) The stress-regulated protein M6a is a key modulator for neurite outgrowth and filopodium/spine formation. *Proc Natl Acad Sci USA* 102: 17196-17201

Palchadhuri M, Flügge G (2005) 5HT_{1A}-receptor expression in pyramidal neurons of cortical and limbic brain regions. *Cell & Tiss Res* 321: 159-172

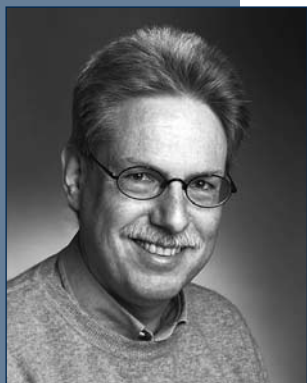
Heilbronner U, van Kampen M, Flügge G (2004) The alpha-2B adrenoceptor in the paraventricular thalamic nucleus is persistently upregulated by chronic psychosocial stress. *Cell Mol Neurobiol* 24: 815-831

Flügge G, van Kampen M, Mijster MJ (2004) Perturbations in brain monoamine systems during stress. *Cell & Tiss Res* 315: 1-14

Fuchs E, Czeh B, Flügge G (2004) Examining novel concepts of the pathophysiology of depression in the chronic psychosocial stress paradigm in tree shrews. *Behav Pharmacol* 15: 315-325

Alfonso J, Pollevick GD, Van Der Hart MG, Flügge G, Fuchs E, Frasch AC (2004) Identification of genes regulated by chronic psychosocial stress and antidepressant treatment in the hippocampus. *Eur J Neurosci* 19: 659-666

Flügge G, van Kampen M, Meyer H, Fuchs E (2003) Alpha2A and alpha2C-adrenoceptor regulation in the brain: alpha2A changes persist after chronic stress. *Eur J Neurosci* 17: 917-28



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Jens Frahm

Professor of Physical Chemistry

- Director of 'Biomedizinische NMR Forschungs GmbH'
- Biomedical Nuclear Magnetic Resonance -

Major Research Interests

General

- development and application of magnetic resonance imaging (MRI) techniques for noninvasive studies of the central nervous system of humans and animals

Methodology

- functional neuroimaging
- localized neurospectroscopy
- diffusion tensor imaging

Brain Research

- non-invasive neurobiology, human neuroscience
- structural, metabolic, and functional studies of the central nervous system
- functional mapping of neuronal activation, cognitive information processing in humans
- MRI of animal models (nonhuman primates, rats, transgenic mice, insects)

Selected Recent Publications

Merboldt KD, Baudewig J, Treue S, Frahm J (2002) Functional MRI of Self-Controlled Stereoscopic Depth Perception. *Neuroreport* 13: 1721-1725

Dechent P, Frahm J (2003) Functional Somatotopy of Finger Representations in Human Primary Motor Cortex. *Hum Brain Mapp* 18: 272-283

Frahm J, Baudewig J, Dechent P, Merboldt KD (2004) Advances in Functional MRI of the Human Brain. *Progr NMR Spectr* 44: 1-32

Watanabe T, Frahm J, Michaelis T (2004) Functional Mapping of Neural Pathways in Rodent Brain *In Vivo* Using Manganese-Enhanced Three-Dimensional Magnetic Resonance Imaging. *NMR Biomed* 17: 554-568

Hofer S, Frahm J (2006) Topography of the Human Corpus Callosum Revisited - Comprehensive Fiber Tractography Using Magnetic Resonance Diffusion Tensor Imaging. *NeuroImage* 32: 989-994



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Eberhard Fuchs

Professor of Neurobiology

- 1977: Dr. rer. nat., University of München
- 1996 - 2000: Professor (Animal Physiology), University of Karlsruhe
- 2000 - 2003: Professor for Animal Physiology, University of Göttingen
- since 2003: Professor for Neurobiology, Department of Neurology, Medical School, University of Götting

Major Research Interests

The Clinical Neurobiology Laboratory (CNL) at the German Primate Center is an interdisciplinary research laboratory using neuroanatomical, neuropharmacological, behavioral and molecular techniques to investigate functioning of the brain in animal models of psychiatric and neurodegenerative diseases. The aim of our work is to elucidate brain structures, circuits, pathways and mechanisms that underlie normal and pathological behavior. This work integrates inputs from other research fields with the ultimate aim of developing new therapeutic strategies for psychiatric and neurodegenerative diseases.

The laboratory specializes in the development, validation and investigation of animal models to detect abnormal cognitive, motor and emotional expressions of brain pathology. Currently, we are engaged in the investigation of central nervous and behavioral phenomena associated with stress and depression. In addition, we provide service platforms to study Parkinson's disease and multiple sclerosis.

Selected Recent Publications

Czéh B, Müller-Keuker JIH, Rygula R, Abumaria N, Hiemke C, Domenici E, Fuchs E (2007) Chronic social stress inhibits cell proliferation in the adult medial prefrontal cortex: hemispheric asymmetry and reversal by fluoxetine treatment. *Neuropsychopharmacology* 32: 1490-1503

Czéh B, Simon M, Schmelting B, Hiemke C, Fuchs E (2006) Astroglial plasticity in the hippocampus after chronic psychosocial stress and concomitant fluoxetine treatment. *Neuropsychopharmacology* 31:1616-26

Fuchs E, Flügge G, Czéh B (2006) Remodeling of neuronal networks by stress. *Front Biosci* 11: 2746-2758

Fuchs E, Czéh B, Kole MHP, Michaelis T, Lucassen PJ (2004) Alterations of neuroplasticity in depression: The hippocampus and beyond. *Europ Neuropharmacol* 14: 481-490

Lucassen PJ, Fuchs E, Czéh B (2004) Antidepressant treatment with tianeptine prevents apoptosis in the hippocampal dentate gyrus and temporal cortex. *Biol Psychiatry* 55: 789-796

Coe CL, Kramer M, Czéh B, Gould E, Reeves AJ, Kirschbaum C, Fuchs E (2003) Prenatal stress diminishes neurogenesis in the dentate gyrus of juvenile rhesus monkeys. *Biol Psychiat* 54: 1025-1034

Czéh B, Michaelis T, Watanabe T, Frahm J, de Biurrun G, van Kampen M, Bartolomucci A, Fuchs E (2001) Stress-induced changes in cerebral metabolites, hippocampal volume and cell proliferation are prevented by antidepressant treatment with tianeptine. *Proc Natl Acad Sci USA* 98: 12796-12801



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Theo Geisel

Professor of Theoretical Physics
Director, Max Planck Institute for Dynamics and Self-Organization
Coordinator, Bernstein Center for Computational Neuroscience

- Dr. rer.nat., University of Regensburg (1975)
- Heisenberg fellow (1983 - 1987)
- Professor of Theoretical Physics, Universities of Würzburg (1988 - 1989), Frankfurt (1989 - 1996), and Göttingen (since 1996)
- Director, Max Planck Institute for Dynamics and Self-Organization, Göttingen (since 1996)

Major Research Interests

How do the myriads of neurons in our cortex cooperate when we perceive an object or perform another task? How do they self-organize in the preceding learning process? Questions like these address the complex dynamics of spatially extended and multicomponent nonlinear systems, which still reserve many surprises. In networks of sufficiently many spiking neurons e.g. we find unstable attractors, a phenomenon which would neither have been guessed nor understood without mathematical modelling and which many physicists consider an oxymoron. They can provide a neuronal network with a high degree of flexibility to adapt to permanently changing tasks. The tools and mathematical methods developed in studies of chaotic behaviour in the past can now help us clarify the dynamics and function of complex networks and spatially extended systems and reveal the biological role of dynamical phenomena like unstable attractors. These methods lend themselves to applications in neuroscience from the level of single cells to the level of cell assemblies and large cortical networks, from the time scales of action potentials (milliseconds) to the time scales of learning and long-term memory (up to years). My work in the past has dealt among others with studies of stochastic resonance of single neurons under periodic and endogenous stimulation, detailed investigations of the properties, functions, and conditions of neuronal synchronization, and the development of neuronal maps in the visual cortex. We have elucidated the influence of the network topology on synchronization and other dynamical properties and demonstrated the existence of speed limits to network synchronization due to disordered connectivity. Besides, I am also focusing on other applications of nonlinear dynamics, e.g. for quantum chaos in semiconductor nanostructures and in mathematical models for the description and forecast of the spread of epidemics.

Selected Recent Publications

Levina A, Herrmann JM, Geisel T (2007) Dynamical Synapses Causing Self-Organized Criticality in Neural Networks. *Nature Physics*, in press

Brockmann D, Hufnagel L, Geisel T (2006) The Scaling Laws of Human Travel. *Nature* 439: 462-465

Wolf F, Timme M, Geisel T (2004) Topological speed limits to network synchronization. *Phys Rev Lett* 92: 074101

Hufnagel L, Brockmann D, Geisel T (2004) Forecast and Control of Epidemics in a Globalized World. *PNAS* 101: 15124

Denker M, Timme M, Diesmann M, Wolf F, Geisel T (2004) Breaking Synchrony by Heterogeneity in Complex Networks. *Phys Rev Lett* 92: 974193

Wolf F, Geisel T (2003) Universality in visual cortical pattern formation. *Journal of Physiology - Paris* 97: 253-264

Timme M, Wolf F, Geisel T (2002) Prevalence of unstable attractors in networks of pulse-coupled oscillators. *Phys Rev Lett* 89(15): 154105



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Martin Göpfert

Professor for Cellular Neurobiology

- 2008 Full Professor for Cellular Neurobiology, University of Göttingen
- 2008 Associate Professor for Molecular Biology and Biophysics of Sensory Systems, University of Cologne
- 2003-2008 Independent group leader, Volkswagen Foundation Group 'Active auditory mechanics in insects', Dept. Animal Physiology, University of Cologne
- 2002-2003 Royal Society University Research Fellow, School of Biological Sciences, University of Bristol
- 1998-2002 DAAD and Leopoldina Research Fellow, Dept. Neurobiology, University of Zürich and School of Biological Sciences, University of Bristol
- 1998 Degree in Biology, University of Erlangen-Nürnberg

Major Research Interests

Our group studies fundamental processes in hearing. By combining mechanical measurements with genetics, molecular biology, immunohistochemistry, electrophysiology, calcium imaging, and biophysical modelling, we are trying to decipher how molecular processes shape the performance of an ear. Our preferred model system is the hearing organ of the fruit fly *Drosophila melanogaster*, the auditory sensory cells of which share conserved molecular modules with the hair cells in our ears.

Our work has uncovered striking parallels between fly and vertebrate hearing, including the functional equivalence of the auditory transduction and adaptation machineries, the motility of auditory sensory cells, transducer-based force generation, and the expression of homologous genes. Our work also provided first insights into the diverse roles of – and interactions between – transient receptor potential (TRP) ion channels in hearing, and a model of TRP-function in the fly's auditory system has been devised. Using a novel electrostatic actuation method, we were able to identify hair cell-like signatures of transducer gating and adaptation in the fly's auditory mechanics and could show that a simple transduction model as proposed to describe hair cell mechanics comprehensively explains the macroscopic behaviour of an ear. Based on these findings, we are currently devising a computational model that allows for the high-throughput characterization of genetic hearing defects. Candidate genes for hearing, in turn, are narrowed down by expression profiling using whole-genome microarrays. By testing how these genes contribute to auditory function and performance, we aim for a comprehensive molecules-to-system description of the functional workings of an ear.

Selected Recent Publications

Nadrowski B, Albert JT, Göpfert MC (2008) Transducer-based force generation explains active process in *Drosophila* hearing. *Curr Biol* 18: 1365-72

Albert JT, Nadrowski B, Göpfert MC (2007) *Drosophila* mechanotransduction: linking functions and proteins. *Fly* 1: 238-241

Albert JT, Nadrowski B, Göpfert MC (2007) Mechanical signatures of transducer gating in the *Drosophila* ear. *Curr Biol* 17: 1000-1006

Albert JT, Winter H, Schaechtinger TJ, Weber T, Wang X, He DZZ, Hendrich O, Geisler, H-S, Zimmermann U, Oelmann K, Knipper M., Göpfert MC, Oliver D (2007) Voltage-sensitive prestin orthologue expressed in zebrafish hair cells. *J Physiol* 580: 451-461

Göpfert MC, Albert JT, Nadrowski B, Kamikouchi A (2006) Specification of auditory sensitivity by *Drosophila* TRP channels. *Nat Neurosci* 8:999-1000

Göpfert MC, Humphris AD, Albert JT, Robert D, Hendrich O (2005) Power gain exhibited by motile mechanosensory neurons in *Drosophila* ears. *Proc Natl Acad Sci USA* 102: 325-330



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Uwe-Karsten Hanisch

Professor for Experimental Neurobiology

- 1986 Diploma Degree Biochemistry University of Leipzig, Germany
- 1990 Ph.D. (Dr. rer. nat.) University of Leipzig, Germany
- 1991-1993 Douglas Hospital Research Centre, McGill University, Montreal, Canada
- 1993-2002 Department of Cellular Neurosciences, Max Delbrück Center for Molecular Medicine (MDC) Berlin, Germany
- 1999 Habilitation (Biochemistry/Neurobiology) University of Leipzig, Germany
- 2002-2004 Professor for Biochemistry University of Applied Sciences Lausitz, Germany
- 2002-2004 Guest scientist and Project leader Molecular Medicine (MDC) Berlin, Germany
- since 2004 Professor for Experimental Neurobiology Institute for Neuropathology, University of Göttingen, Germany
- since 2007 Guest Professor Medical Physiology, University of Groningen, The Netherlands

Major Research Interests

Expression and functions of cytokines in the CNS
Mechanisms of microglial activation and consequences of microglial activities
Role of plasma factors as endogenous signals for microglial cells

Selected Recent Publications

Ribes S, Ebert S, Czesnik D, Regen T, Zeug A, Bukowski S, Mildner A, Eiffert H, Hanisch UK, Hammerschmidt S, Nau R (2009) Toll-like receptor prestimulation increases phagocytosis of *Escherichia coli* DH5 and *Escherichia coli* K1 strains by murine microglial cells. *Infect Immun* 77: 557-564

Prinz M, Schmidt H, Mildner A, Knobloch KP, Hanisch UK, Detje C, Gutcher I, Mages J, Lang R, Martin R, Merkler D, Raasch J, Gold R, Becher B, Brück W, Kalinke U (2008) Distinct and nonredundant *in vivo* functions of IFNAR on myeloid cells and autoimmunity in the central nervous system. *Immunity* 28: 675-86

Hanisch UK, van Rossum (2009) Microglia properties. In Squire LR (ed), *Encyclopedia of Neuroscience*, volume 5: 853-859, Academic Press, Oxford

Hanisch UK, Johnson TV, Kipnis J (2008) Toll-like receptors: roles in neuroprotection? *Trends Neurosci* 31: 176-182

van Rossum D, Hilbert S, Straßenburg S, Hanisch UK, Brück W (2008) Myelin-phagocytosing macrophages in isolated sciatic and optic nerves reveal a unique reactive phenotype. *Glia* 56: 271-283

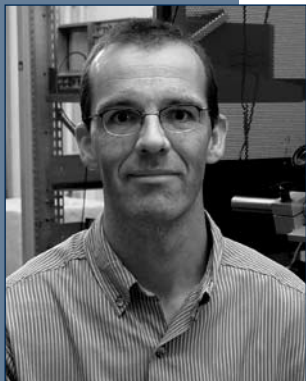
Weinstein JR, Swarts S, Bishop C, Hanisch UK, Möller T (2008) Lipopolysaccharide is a frequent and significant contaminant in commercial-grade preparations of putative microglia-activating factors. *Glia* 56: 16-26

Mildner A, Schmidt H, Nitsche M, Merkler D, Hanisch UK, Mack M, Heikenwälder M, Brück W, Priller J, Prinz M (2007) Microglia in the adult brain arise from Ly-6Chi monocytes only under defined host conditions. *Nat Neurosci* 10: 1544-1553

Hanisch UK, Kettenmann H (2007) Microglia: active sensor and versatile effector cells in the normal and pathologic brain. *Nat Neurosci* 10: 1387-1393

Schwartz M, Butovsky O, Brück W, Hanisch UK (2006) Microglial phenotype: Is the commitment reversible? *Trends Neurosci* 29: 68-74

Weinstein JR, Hong S, Kulman JD, Bishop C, Kuniyosh J, Andersen H, Ransom BR, Hanisch UK, Möller T (2005) Unraveling thrombin's true microglia-activating potential: Markedly disparate profiles of pharmaceutical-grade and commercial-grade thrombin preparations. *J Neurochem* 95: 1177-1187



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Ralf Heinrich

Juniorprofessor of Molecular Neuropharmacology of Behavior

- Dr. rer. nat., University of Göttingen, 1995
- Postdoctoral fellow, Harvard Medical School, Boston, USA, 1997 - 1999

Major Research Interests

Behavior results from integration of sensory information with internal physiological states involving complex interactions between various types of neurons. In order to study cellular and molecular mechanisms that contribute to the selection and control of situation-specific behavior, invertebrate preparations can offer unique advantages over more complex nervous systems of vertebrates, especially mammals. The nervous systems of invertebrates contain smaller numbers of neurons, many of which can be individually identified, and their behavioral repertoires are rather limited to combinations of genetically determined stereotyped components.

Studies are conducted with intact or partially dissected behaving animals (insects, crustaceans, annelids) and with isolated nervous systems or cultured organs and cells. Projects for experimental theses usually combine two or more of the following methods: neuroethology, pharmacology, electrophysiology, histology and immunocytochemistry, cell culture and molecular biology. Examples of current research projects are

- Acoustic communication in grasshoppers: control of sound production by converging signaling pathways (transmitters and second messengers) in the central complex neuropil of the brain.
- Physiological characterization of neurosecretory neurons that mediate general physiological states e.g. serotonin-releasing neurons of leeches and crustaceans.
- Control of agonistic behavior and the formation of hierarchies in crustaceans, crickets and fruitflies.
- Presence and function of erythropoietin in invertebrate nervous systems: development, regeneration and hypoxia-related functions

Selected Recent Publications

Gocht D, Heinrich R (2007) Postactivation inhibition of spontaneously active neurosecretory neurons in the medicinal leech. *J Comp Physiol A* 193: 347-361

Heinrich R, Ganter GK (2007) The role of NO in insect behavior. *Advances in Experimental Biology* 1: 107-127

Wenzel B, Kunst M, Günther C, Ganter GK, Lakes-Harlan R, Elsner N, Heinrich R (2005) Nitric oxide/cyclic GMP-signaling in the central complex of the grasshopper brain inhibits singing behavior. *J Comp Neurol*, 488: 129-139

Wenzel B, Elsner N, Heinrich R (2002) mAChRs in the grasshopper brain mediate excitation by activation of the AC/PKA and the PLC second-messenger pathways. *J Neurophysiol*, 87: 876-888

Heinrich R, Wenzel B, Elsner N (2001) A role for muscarinic excitation: Control of specific singing behavior by activation of the adenylate cyclase pathway in the brain of grasshoppers. *Proc Nat Acad Sci USA* 98: 9919-9923



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Michael Hörner

Professor of Cellular Neurobiology

- Dr. rer. nat., University of Göttingen, 1989
- Postdoctoral Fellow, Medical University of Kiel, Dept. Physiology, 1989 - 1990
- Assistant Professor, Institute for Zoology and Anthropology, Göttingen, 1990 - 1997
- Habilitation (Zoology), 1997
- Associate Professor, Institute for Zoology and Anthropology, Göttingen, 1997 - 2002
- Guest Professor, University of Science & Technology, Hongkong, 2002 - 2004
- Apl. Professor, Inst. for Zoology, Anthropol. and Develop. Biol., Göttingen, since 2004
- Research Assistant, MPI for Ethology, Seewiesen, 1985/1986
- Research Fellow, Arizona Research Labs, Tucson, USA, 1993/1996
- Feodor-Lynen/Humboldt Fellow, Harvard Medical School, Boston, USA, 1994 - 1995
- Research Fellow Marine Biological Labs, Woods Hole, USA, 1992/1997

Major Research Interests

Molecular Mechanisms Of Synaptic And Non-Synaptic Modulation

Biogenic amines such as serotonin, dopamine, histamine or octopamine (OA), the pendant of norepinephrine in invertebrates, are widely distributed within the animal kingdom. These evolutionary conserved neuroactive substances are involved in the control of vital functions in both vertebrates and invertebrates. Biogenic amines often initiate long-lasting neuro-modulatory effects in their targets, which is due to diffusion following non-synaptic release activating G-protein coupled to intracellular pathways. My work is focussed on the investigation of cellular and molecular mechanisms underlying the modulation of neuronal signaling in identified networks in invertebrate model systems. Using electrophysiological, pharmacological and immunocytochemical techniques in combination with behavioral measurements, I am investigating mechanisms of aminergic modulation in identified neurons of defined networks in insects and crustacea. To address both mechanistic and functional questions, a parallel approach has been developed, which allows to investigate single identified neurons both *in-vivo* with intact synaptic connections and *in-vitro* in primary "identified" cell culture, where neurons are separated from connections to other neurons. The functional meaning of aminergic modulation on the cellular level in behaviorally-relevant circuits is assessed by quantitative behavioral measurements. The investigations show that OA enhances the responsiveness of a neuronal network in insects ("giant fiber pathway") which triggers a fast escape reaction. The reaction to sensory stimuli in the postsynaptic giant interneurons, which are monosynaptically coupled to sensory neurons via excitatory cholinergic synapses, is significantly enhanced by OA application. Characteristic changes of the action potentials *in-vivo* ("spike broadening") and patch-clamp recordings *in-vitro* suggest, that OA selectively affects slow K⁺-conductances in postsynaptic giant interneurons

Selected Recent Publications

- Kloppenburg P, Hörner M (1998) Voltage-activated currents in identified giant interneurons isolated from adult crickets, *Gryllus bimaculatus*. J Exp Biol 201(17): 2529-2541
- Heinrich R, Cromarty SI, Hörner M, Edwards DH, Kravitz EA (1999) Autoinhibition of serotonin cells: An intrinsic regulatory mechanism sensitive to the pattern of usage of the cells. Proc Natl Acad Sci USA 96: 2473-2478
- Ferber M, Hörner M, Cepok S, Gnatzy W (2001) Digger wasp versus cricket: Mechanisms underlying the total paralysis caused by the predators venom. J Neurobiol 47: 207-2222
- Hörner M, Heinrich R, Cromarty SI, Kravitz EA (2002) Synaptic connectivity of amine-containing neurosecretory cells of lobsters: inputs to 5HT- and OCT- containing neurons. in: The Crustacean Nervous System. (ed. K. Wiese) Springer Verlag, Berlin, Heidelberg, New York, pp156-172
- Rose T, Gras H, Hörner M (2006) Activity-dependent suppression of spontaneous spike generation in the Retzius neurons of the leech, *Hirudo medicinalis* L. Invertebrate Neuro-science 6: 169-176 (DOI 10.1007/s10158-006-0030-2)



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Sven Hülsmann

Privatdozent, Department of Neurophysiology

- Dr. med., University of Münster, 1995
- Postdoctoral fellow, University of Münster Dept. of Neurosurgery, 1995 - 1996
- Postdoctoral fellow, University of Göttingen, Dept. of Neurophysiology, 1996 - 2001
- Group leader (Wissenschaftlicher Assistent) Neurophysiology, since 2001
- Principle Investigator at the DFG Research Center for Molecular Physiology of the Brain (CMPB) since 2002
- Habilitation, University of Göttingen, 2005

Major Research Interests

The majority of cells in the human brain are glial cells, outranging the number of neurons by a factor of 10. However, most behavioral aspects of life are attributed to neurons, leaving a rather white spot of knowledge about the function of the different types of glial cells. Our group aims to identify and clarify the mechanisms that allow glial cells, e.g. astrocytes to modulate and stabilize the most vital behavior of breathing.

Selected Recent Publications

Holpert M, Groß U, Bohne W (2006) Disruption of the bradyzoite-specific P-type (H⁺)-ATPase /PMA1/ in *Toxoplasma gondii* leads to decreased bradyzoite differentiation after stress stimuli but does not interfere with mature tissue cyst formation. *Mol Biochem Parasitol* 146:129-33

Fasshauer V, Groß U, Bohne W (2005) The parasitophorous vacuole membrane of *Encephalitozoon cuniculi* lacks host cell membrane proteins immediately after invasion. *Eukaryot Cell* 4: 221-224

Lüder CGK, Groß U (2005) Apoptosis and its modulation during infection with *Toxoplasma gondii*: molecular mechanisms and role in pathogenesis. *Curr Topics Microbiol Immunol* 289: 219-238

Weig M, Jäntsich L, Groß U, de Koster CG, Klis FM, de Groot PWJ (2004) Systematic identification in silico of covalently bound cell wall proteins and analysis of protein-polysaccharide linkages of the human pathogen *Candida glabrata*. *Microbiology* 150: 3129-3144

Lüder CGK, Lang C, Giraldo-Velasquez M, Algner M, Gerdes J, Groß U (2003) *Toxoplasma gondii* inhibits MHC class II expression in neural antigen-presenting cells by down-regulating the class II transactivator CIITA. *J Neuroimmunol* 134:12-24



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Professor, Director at the Max Planck Institute for Biophysical Chemistry

- Dr. rer. nat. 1981, University of Göttingen
- Assistant Professor, The Rockefeller University, New York (USA) 1985
- Junior Group leader, Max Planck Institute for Psychiatry, Martinsried, 1986
- Associate Professor of Pharmacology and Cell Biology, Yale University, and Investigator, Howard Hughes Medical Institute, New Haven (USA) 1991
- Professor of Pharmacology and Cell Biology, Yale University, New Haven, 1995
- Director, Max Planck Institute for Biophysical Chemistry, Göttingen, 1997

Major Research Interests

Our group is interested in the mechanisms of membrane fusion, with the main emphasis on regulated exocytosis in neurons. Since recent years it is known that intracellular membrane fusion events are mediated by a set of conserved membrane proteins, termed SNAREs. For fusion to occur, complementary sets of SNAREs need to be present on both of the fusing membranes. The neuronal SNAREs are among the best characterized. They are the targets of the toxins responsible for botulism and tetanus. To understand how these proteins make membranes fuse, we studied their properties in detail using biochemical and biophysical approaches. We found that they assemble into a tight complex which ties the membrane closely together and thus probably initiates bilayer mixing.

In our current approaches, we study membrane fusion at the level of isolated proteins as well as in semi-intact and intact cells. Thus, we are investigating conformational changes of the SNARE proteins before and during fusion. Furthermore, we use reconstitution of membrane fusion in cell-free assays and in proteoliposomes. Other projects of the group include the study of neurotransmitter uptake by synaptic vesicles and the function of Rab-GTPases in neuronal exocytosis

Selected Recent Publications

Zwilling D, Cypionka A, Pohl W, Fasshauer D, Walla PJ, Wahl MC, Jahn R (2007) Early endosomal SNAREs form a structurally conserved SNARE complex and fuse liposomes with multiple topologies. *EMBO J* 26: 9-18

Takamori S, Holt M, Stenius K, Lemke EA, Grønborg M, Riedel D, Urlaub H, Schenck S, Brügger B, Ringler P, Müller SA, Rammner B, Gräter F, Hub JS, De Groot BL, Mieskes G, Moriyama Y, Klingauf J, Grubmüller H, Heuser J, Wieland F, Jahn R (2006) Molecular anatomy of a trafficking organelle. *Cell* 127: 831-846

Jahn R, Scheller RH (2006) SNAREs – engines for membrane fusion. *Nature Reviews Mol Cell Biol* 7: 631-643

Willig KI, Rizzoli SO, Westphal V, Jahn R, Hell S (2006) STED-microscopy reveals that the synaptic vesicle protein synaptotagmin remains clustered after exocytosis. *Nature* 440: 935-939

Graf C, Riedel D, Schmitt HD, Jahn R (2005) Identification of functionally interacting SNAREs using complementary substitutions in the conserved '0' layer. *Mol Biol Cell* 16: 2263-2274

Schuetz CG, Hatsuzawa K, Margittai M, Stein A, Riedel D, Küster P, König M, Seidel CAM, Jahn R (2004) Determinants of liposome fusion mediated by synaptic SNARE proteins. *Proc Natl Acad Sci* 101: 2858-2863

Jahn R, Lang T, Südhof TC (2003) Membrane fusion. *Cell* 112: 519-533



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Hubertus Jarry

Professor of Clinical and Experimental Endocrinology

- 1976 - 1980 University of Göttingen, study of biology, diploma degree in biochemistry, microbiology, organic chemistry
- 1980 - 1983 PhD thesis, Department of Biochemistry, University of Göttingen,
- PhD degree in biochemistry, microbiology, organic chemistry (summa cum laude)
- Until February 1985 German Primate Center Göttingen, Dept. Reproductive Biology
- March 1985 until March 1986 Michigan State University, Dept. Pharmacology and Toxicology
- Since April 1986 Research Associate Dept. Clinical and Experimental Endocrinology University of Göttingen
- Januar 1991 Habilitation
- Dezember 1995 Promotion to Professor

Major Research Interests

The proper function of the GnRH pulse generator is essential for reproduction of all mammals studied so far. GnRH pulses are a prerequisite for proper pituitary gonadotropin release. The neurochemical mechanisms leading to pulsatile GnRH release involve norepinephrine and gamma amino butyric acid (GABA) as most important neurotransmitters. In addition, other catecholamines, amino acid neurotransmitters and neuropeptides play a modulatory role in the function of the GnRH pulse generator. Many of the GABAergic neurons in the hypothalamus are estrogen-receptive. The mechanisms by which the estrogen receptors of the alpha and beta subtype regulate gene and protein expression of neurotransmitter-producing enzymes are at present a prime focus of interest. Induction of puberty is not a gonadal but a hypothalamic maturational process. The initiation of proper GnRH pulse generator function is the ultimate trigger signal for puberty which is currently investigated. Ageing involves also neuroendocrine mechanisms. The GnRH pulse generator function deteriorates in aged rats, mechanisms which involve a variety of catecholamines and amino acid neurotransmitters which are currently investigated. Steroidal feedback signals (of estradiol, progesterone, and glucocorticoids) are crucial for the development and proper function of the adult hypothalamus of which the molecular and neurochemical mechanisms are studied with cell biological and animal experimental tools. Proper function of the GnRH pulse generator is also of crucial importance for initiation of puberty and maintenance of normal menstrual cycles in women. Many of hitherto unexplained infertilities can be explained of malfunctioning GnRH pulse generators which are studied in a series of clinical experiments.

Selected Recent Publications

- Bottner M, Leonhardt S, Wuttke W, Jarry H (2007) Changes of expression of genes related to the activity of the gonadotrophin-releasing hormone pulse generator in young versus middle-aged male rats. *J Neuroendocrinol* 19: 779-87
- Zhou L, Lehan N, Wehrenberg U, Disteldorf E, von Lossow R, Mares U, Jarry H, Rune GM (2007) Neuroprotection by estradiol: a role of aromatase against spine synapse loss after blockade of GABA(A) receptors. *Exp Neurol* 203: 72-81
- Breit A, Wolff K, Kalwa H, Jarry H, Buch T, Gudermann T (2006) The natural inverse agonist agouti-related protein induces arrestin-mediated endocytosis of melanocortin-3 and -4 receptors. *J Biol Chem* 281: 37447-56
- Fester L, Ribeiro-Gouveia V, Prange-Kiel J, von Schassen C, Bottner M, Jarry H, Rune GM (2006) Proliferation and apoptosis of hippocampal granule cells require local oestrogen synthesis. *J Neurochem* 97: 1136-44



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Jürgen Klingauf

Research Group Leader at the Max Planck Institute for Biophysical Chemistry

- Research fellow, Dept. of Molecular & Cellular Physiology, Stanford University, Ca, 1996 - 1998
- Dr. rer. nat. (Ph.D.) 1999, University of Göttingen
- Since 2000 junior group leader at the Max Planck Institute for Biophysical Chemistry

Major Research Interests

The focus of our research is the study of synaptic transmission, with the emphasis on presynaptic mechanisms. At the synapse, neurotransmitter is rapidly released from small vesicles which are triggered to fuse with the plasma membrane by the entry of Ca^{2+} ions. The maintenance of synaptic transmission requires that these vesicles be retrieved by a reverse process, i.e. endocytosis. How is this endocytic activity and subsequent formation of fusion-competent vesicles coupled to exocytosis? To delineate the mechanisms by which synaptic vesicles can be retrieved we employ high-resolution imaging techniques, like two-photon laser scanning and total internal reflection microscopy, electrophysiology, as well as biochemical approaches. By transfection of neurons in primary cell culture or the usage of knock-out models we can target or modulate specific proteins thought to be pivotal in synaptic vesicle endocytosis. Currently, we are mainly studying synapses of rodent hippocampus, down to the level of single fluorescently labeled vesicles in cultured or freshly isolated synaptic boutons. By making use of fluorescent styryl dyes with different kinetic properties we found that in central nervous synapses at least two kinetically distinct modes of endocytosis co-exist. We are now trying to characterize the respective molecular events underlying those different mechanisms using genetically encoded fluorescent probes.

Selected Recent Publications

Mueller VJ, Wienisch M, Nehring RB, Klingauf J (2004) Monitoring clathrin-mediated endocytosis during synaptic activity. *J Neurosci* 24(8): 2004-12

Jordan R, Lemke EL, Klingauf J (2005) Visualization of synaptic vesicle movement in intact synaptic boutons using fluorescence fluctuation spectroscopy. *Biophys J* 89(3): 2091-102

Lemke EL, Klingauf J (2005) Single synaptic vesicle tracking in individual hippocampal boutons at rest and during synaptic activity. *J Neurosci* 25(47): 11034-44

Vanden Berghe P, Klingauf J (2006). Synaptic vesicles in hippocampal boutons recycle to different pools in a use-dependent fashion. *J Physiol London* 572(Pt 3): 707-20

Diril MK, Wienisch M, Jung N, Klingauf J, Haucke V (2006) Stonin 2 is an AP-2-dependent endocytic sorting adaptor for synaptotagmin internalization and Recycling. *Dev Cell* 10(2): 233-44

Wienisch M, Klingauf J (2006) Vesicular proteins exocytosed and subsequently retrieved by compensatory endocytosis are non-identical. *Nature Neurosci* 9(8): 1019-27

Toonen RF, Kochubey O, de Wit H, Gulyas-Kovacs A, Konijnenburg B, Sørensen JB, Klingauf J, Verhage M (2006) Dissecting docking and tethering of secretory vesicles at the target membrane. *EMBO J* 25(16): 3725-37

Kochubey O, Majumdar A, Klingauf J (2006) Imaging clathrin dynamics in *D. melanogaster* hemocytes reveals a role for actin in vesicle fission. *Traffic* Oct 2, Epub ahead of print



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Till Marquardt

Group Leader Developmental Neurobiology Laboratory

- Since 2007: independent research group leader, DFG Emmy Noether group leader at the European Neuroscience Institute, Göttingen
- 2001 - 2006: postdoctoral research associate and staff scientist with Samuel L. Pfaff at the Salk Institute for Biological Studies in La Jolla, California, USA
- 2001: Ph.D. with Peter Gruss at the Max-Planck Institute of Biophysical Chemistry, University of Göttingen

Major Research Interests

Adequate control of body motion and posture depends on elaborate circuitries that connect both motor and sensory neurons with the musculature. The central importance of these connections is illustrated by the debilitating consequences of diseases affecting motor neurons, such as Amyotrophic Lateral Sclerosis (ALS) and diabetic neuropathy. Our research aims at understanding the molecular mechanisms driving the assembly of functional neuromuscular circuitries during embryonic and postnatal development. This includes the study of cell surface-based signaling molecules that control motor and sensory axon connectivity in mice. Another research focus of the lab aims at identifying and characterizing novel mechanisms driving the functional specification of motor neurons within the context of operative neuromuscular circuitry. We extensively take advantage of mouse genetics in order to selectively trace and manipulate specific neuron populations. We combine this genetic approach with live 3D fluorescence (*spinning disk*) microscopy, as well as electrophysiological methods to elucidate the role of cell surface and nuclear receptor proteins in sensory-motor connectivity and functional neuron specification.

Selected Recent Publications

Marquardt T, Shirasaki R, Ghosh S, Carter N, Andrews SE, Hunter T, Pfaff SL (2005) Co-expressed EphA receptors and ephrin-A ligands mediate opposing actions on growth cone navigation from distinct membrane sub-domains. *Cell* 121: 127-139

Marquardt T, Pfaff SL (2001) Cracking the transcriptional code for cell specification in the neural tube. *Cell* 106: 651-654

Marquardt T, Ashery-Padan RA, Andrejewski N, Scardigli R, Guillemot F, Gruss P (2001) Pax6 is required for the multipotent state of retinal progenitor cells. *Cell* 105: 43-55



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Tobias Moser

Professor of Experimental and Clinical Audiology

- Dr. med. (M.D.) 1995, University of Jena
- Postdoctoral fellow with E. Neher at the MPI for Biophysical Chemistry, 1994 - 1997
- Group leader at the Department of Otolaryngology, University of Göttingen since 1997

Major Research Interests

Our group focuses on the physiology and pathology of sound coding at the hair cell ribbon synapse. Molecular dissection and detailed physiological characterization of ribbon synapse function employ a spectrum of molecular and biophysical techniques such as single cell RT-PCR, immunohistochemistry of hair cells, auditory systems physiology (recordings of otoacoustic emissions, compound action potentials and auditory brainstem responses, single unit recordings), pre- or postsynaptic patch-clamp, optical methods (epifluorescence, evanescent wave and confocal imaging as well as flash photolysis of caged compounds). The group has contributed to understanding normal hair cell ribbon synapse function (reviews in Nouvian et al., 2006 and Moser et al., 2006). In our previous work we have physiologically and in part morphologically characterized mutant mice with defects in hair cell synaptic coding (Brandt et al., 2003; Khimich et al., 2005, Roux et al., 2006) and auditory nerve function (Lacas-Gervais et al., 2004). The results demonstrated that defects of hair cell synaptic sound coding cause sensorineural hearing loss in animal models – auditory synaptopathy and confirmed impaired hearing in case of nerve disorders - auditory neuropathy.

Selected Recent Publications

Nouvian R, Beutner D, Parsons TD, Moser T (2006) Structure and function of the hair cell ribbon synapse. *J Membr Biol* 209: 153-65

Roux I, Safieddine S, Nouvian R, Grati M, Simmler MC, Perfettini I, Le Gall M, Rostaing P, Hamard G, Triller A, Avan P, Moser T, Petit C (2006) Otoferlin, defective in DFNB9 deafness, is essential for the Ca²⁺-triggered synaptic exocytosis at the auditory hair cell ribbon synapse. *Cell* 127: 277-89

Moser T, Brandt A, Lysakowski A (2006) Hair cell ribbon synapses. *Cell Tissue Res* 326: 347-359

Khimich D, Nouvian R, Pujol R, tom Dieck S, Egner A, Gundelfinger ED, Moser T (2005) Hair Cell Synaptic Ribbons are Essential for Synchronous Auditory Signaling. *Nature* 434: 889-94

Brandt A, Khimich D, Moser T (2005) Few Ca_v 1.3 channels regulate a synaptic vesicle's exocytosis at the hair cell ribbon synapse. *J Neurosci* 25: 11577-11585

Beutner D, Voets T, Neher E, Moser T (2001) Calcium dependence of exocytosis and endocytosis at the cochlear inner hair cell afferent synapse. *Neuron* 29: 681-90

Moser T, Beutner D (2000) Kinetics of exocytosis and endocytosis at the cochlear inner hair cell afferent synapse of the mouse. *Proc Natl Acad Sci USA* 97: 883-888



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Klaus-Armin Nave

Professor of Molecular Biology, Director at the Max Planck Institute of Experimental Medicine

- 1987 PhD, University of California, San Diego
- 1987-1991 Postdoc, The Salk Institute, La Jolla, California
- 1991 Junior Group Leader, ZMBH, University of Heidelberg
- 1998 Professor of Molecular Biology (C4), ZMBH
- 2000 Director, Department of Neurogenetics Max Planck Institute for Experimental Medicine, Göttingen, and Professor of Biology, University of Heidelberg

Major Research Interests

We are interested in the mechanisms of neuron-glia interactions in the higher nervous system, and in the genes that are required for normal glial cell function. Here, transgenic and mutant mice have become important to study developmental processes as well as genetic diseases. For example, oligodendrocytes are glial cells highly specialized for enwrapping CNS axons with multiple layers of membranes, known to provide electrical insulation for rapid impulse propagation. We found that oligodendrocytes are also essential for maintaining the long-term integrity of myelinated axons, independent of the myelin function itself. The mechanisms by which oligodendrocytes support long-term axonal survival are still under investigation. The importance of glial cells as the “first line of neuroprotection”, however, is illustrated by several myelin-associated diseases in which axonal neurodegeneration contribute to progressive disability. These range in humans from peripheral neuropathies (CMT1) to spastic paraplegia (SPG2), and presumably multiple sclerosis (MS) and certain forms of psychiatric disorders. We are developing transgenic animal models for some of these diseases, in order to dissect the underlying disease mechanisms and, in the case of CMT1A, have used these models to design novel therapeutic strategies.

The glial “decision” to myelinate an axonal segment is partly controlled by the axon itself, but the signaling mechanism is not understood. We have found that axonal neuregulin-1 (NRG1) is the major determinant of myelination in the peripheral nervous system. We are now investigating NRG1 dysregulation also in CNS myelination, using quantifiable behavioural functions in mice. By combining genetics with environmental risk factors for schizophrenia (in collaboration with H. Ehrenreich) we will explore the hypothesis that NRG1, a known human schizophrenia susceptibility gene, points to an important role of myelinating glia in some psychiatric disorders.

Future Projects and Goals

Mechanisms of neuron-glia signalling; function of myelin proteins and lipids; transcriptional profiling of single cells *in vivo*; novel mouse models of neuropsychiatric disorders.

Selected Recent Publications

Kassmann CM, Lappe-Siefke C, Baes M, Brügger B, Mildner A, Werner HB, Natt O, Michaelis Th, Prinz M, Frahm J, Nave K-A (2007) Axonal loss and neuroinflammation caused by peroxisome-deficient oligodendrocytes. *Nature Genetics* 8: 969-976

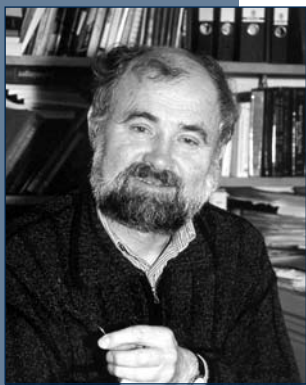
Dhaunchak A, Nave K-A (2007) A common mechanism of proteolipid protein misfolding leading to cysteine-mediated ER retention in oligodendrocytes and Pelizaeus-Merzbacher disease. *Proc Natl Acad Sci USA* (in press)

Kassmann CM, Lappe-Siefke C, Baes M, Brügger B, Mildner A, Werner HB, Natt O, Michaelis Th, Prinz M, Frahm J, Nave K-A (2007) Axonal loss and neuroinflammation caused by peroxisome-deficient oligodendrocytes. *Nature Genetics* 8: 969-976

Kramer-Albers EM, Gehrig-Burger K, Thiele C, Trotter J, Nave K-A (2006) Perturbed interactions of mutant proteolipid protein/DM20 with cholesterol and lipid rafts in oligodendroglia: implications for dysmyelination in spastic paraplegia. *J Neurosci* 26: 11743-11752

Saher G, Brügger B, Lappe-Siefke C, Möbius W, Tozawa R, Wehr M, Wieland F, Ishibashi S, Nave K-A (2005) Cholesterol is essential and rate-limiting for myelin membrane growth. *Nature Neurosci* 8: 468-475

Michailov GV, Sereda MW, Brinkmann BG, Fischer TM, Haug B, Birchmeier C, Role L, Lai C, Schwab MH, Nave K-A (2004) Axonal neuregulin-1 regulates myelin sheath thickness. *Science* 304: 700-703



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Erwin Neher

Professor, Director at the Max Planck Institute for Biophysical Chemistry

- M.Sc. (Physics), University of Wisconsin, (1967)
- Ph.D. (Physics), Institute of Technology, Munich (1970)
- Research associate at the Max Planck Institute for Biophysical Chemistry in Göttingen, Germany (1972 - 1975 and 1976 - 1982) and as a guest in the laboratory of Dr. Ch.F. Stevens at Yale University, Dept. of Physiology, New Haven, Conn. (1975 - 1976)
- Fairchild Scholar, California Institute of Technology; Pasadena, USA (1989)
- Director of the Membrane Biophysics Department at the Max Planck Institute for Biophysical Chemistry, Göttingen, Germany, since 1983

Major Research Interests

Molecular Mechanisms of Exocytosis, Neurotransmitter Release, and Short Term Synaptic Plasticity

In order to understand how the brain handles its information flow and adjusts synaptic connections on the second and subsecond timescale, one has to understand all aspects of synaptic transmission ranging from availability of vesicles for exocytosis, presynaptic electrophysiology, Ca^{++} signalling, the process of exocytosis, and postsynaptic neurotransmitter action. Our work concentrates on presynaptic aspects. We study the basic mechanisms of exocytosis, using adrenal chromaffin cells as a model system and the patch-clamp method. This work, in which intracellular Ca^{++} is manipulated (caged Ca^{++}) and measured on the single cell level aims at understanding the role of specific synaptic proteins in the maturation and exocytosis of secretory vesicles. We use neuronal cell cultures and brain slices for studying mechanisms of short term plasticity, such as depression and paired pulse facilitation. The Calyx of Held, a specialized synapse in the auditory pathway, offers unique possibilities for simultaneous pre- and postsynaptic voltage clamping. This allows a quantitative analysis of the relationship between $[Ca^{++}]$ and transmitter release.

Selected Recent Publications

Sakaba T, Stein A, Jahn R, Neher E (2005) Distinct kinetic changes in neurotransmitter release after SNARE protein cleavage. *Science* 309: 491-494

Soerensen J, Nagy G, Varoqueaux F, Nehring RB, Brose N, Wilson MC, Neher E (2003). Differential control of the releasable vesicle pools by SNAP-25 splice variants and SNAP-23. *Cell* 114: 75-86

Sakaba T, Neher E (2003) Direct modulation of synaptic vesicle priming by GABAB receptor activation at a glutamatergic synapse. *Nature* 424: 775-778

Rettig J, Neher E (2002) Emerging roles of presynaptic proteins in Ca^{++} -triggered exocytosis. *Science* 298: 781-785

Schneggenburger R, Neher E (2000) Intracellular calcium dependence of transmitter release rates at a fast central synapse. *Nature* 406: 889-893

Neher E (1998) Vesicle pools and Ca^{2+} microdomains: new tools for understanding their roles in neurotransmitter release. *Neuron* 20: 389-399

Klingauf J, Neher E (1997) Modeling buffered Ca^{2+} diffusion near the membrane: Implications for secretion in neuroendocrine cells. *Biophys J* 72: 674-690



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- Dr. med., University of Düsseldorf, 1978
- Training in Neurology at the Universities of Düsseldorf, UCL London and Munich
- Habilitation (Neurology and Clinical Neurophysiology) in Munich
- Prof. and Head of the Department of Clinical Neurophysiology 1992

Major Research Interests

Our main research goal is to development new neurophysiologically based therapies for neurological diseases incorporating excitability changes of the brain. For this we use repetitive transcranial magnetic stimulation (rTMS) and transcranial direct current stimulation (TDCS). TMS induces a short electric current in the human brain. Both rTMS and TDCS offer the prospect of inducing LTD and LTP like effects in the human brain. Diseases in our focus are Parkinson's disease, epilepsy, migraine, stroke and dystonia.

Both methods may also be used to measure excitability changes in the motor cortex or alterations in visual perception thresholds. We also evaluate rTMS and TDCS induced changes in motor cortex excitability by functional MR imaging.

Selected Recent Publications

Kuo MF, Paulus W, Nitsche MA (2007) Boosting Focally-Induced Brain Plasticity by Dopamine. *Cereb Cortex*

Nitsche MA, Roth A, Kuo MF, Fischer AK, Liebetanz D, Lang N, Tergau F, Paulus W (2007) Timing-dependent modulation of associative plasticity by general network excitability in the human motor cortex. *J Neurosci* 27(14): 3807-12

Nitsche MA, Doemkes S, Karakose T, Antal A, Liebetanz D, Lang N, Tergau F, Paulus W (2007) Shaping the effects of transcranial direct current stimulation of the human motor cortex. *J Neurophysiol* 97(4): 3109-17. Epub 2007 Jan 24



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Diethelm W. Richter

Professor of Physiology

**Chairman of the II. Department of Physiology,
University of Göttingen**

Speaker of the European Neuroscience Institute Göttingen

- 1969 - 1970 Wiss. Angestellter, I. Physiol. Inst., University of Saarland
- 1970 - 1972 Wiss. Assistent, I. Physiol. Inst., University of Saarland
- 1972 - 1974 Wiss. Assistent, I. Physiol. Inst., University of Munich
- 1974 Universitätsdozent, I. Physiol. Inst., University of Munich
- 1975 - 1976 Universitätsdozent, I. Physiol. Inst., University of Heidelberg
- 1976 - 1988 C-3 Professor, I. Physiol. Inst., University of Heidelberg
- 1988 C-4 Professor, II. Physiol. Inst., University of Göttingen

Major Research Interests

Neurotransmitters, neuromodulators, and peptide hormones are known to activate metabotropic receptor proteins that control ion channels or second messenger cascades. These receptors regulate an intracellular network of interacting signal transduction pathways by means of G-proteins. Thus, receptors transmit extracellular signals to intracellular proteins and other chemical factors. These signals are normally not transduced in a stereotypic manner, but they are integrated in a space- and time-dependent manner, resulting in highly dynamic and variable cellular responses. The specific nature of the cellular response depends on individual cell types that may differ in the expression pattern of receptor subtypes or of intracellular signaling factors. Our research group concentrates on the spatial organization of various subtypes of serotonin receptors and targets an understanding of the highly localized regulation of molecular interactions occurring simultaneously at many sites of a neuron. The goal is to achieve a refined understanding of the parallel signal processing within networks of chemical signal pathways and to clarify their effects on the properties of the neuron as a whole.

Another task addressing complex brain functions is to transfer this knowledge about molecular signaling within cells to the integrated function of neuronal networks. The problem is that modulation of network systems cannot be predicted simply on the basis of cellular reactions, because subgroups of diversely wired neurons mostly express heterogeneous receptor profiles.

Selected Recent Publications

Renner U, Glebov K, Lang T, Papusheva E, Balakrishnan S, Keller B, Richter DW, Jahn R, Ponimaskin E (2007) Localization of the 5-HT_{1A} receptor in lipid microdomains depends on its palmitoylation and is involved in receptor-mediated signaling. *Mol Pharmacol* 72(3): 502-13

Stettner GM, Huppke P, Brendel C, Richter DW, Gartner J, Dutschmann M. (2007) Breathing dysfunctions associated with impaired control of postinspiratory activity in *Mecp2*^{-/-} knockout mice. *J Physiol* 579(3): 863-76

Neusch C, Papadopoulos N, Müller M, Maletzki I, Winter SM, Hirrlinger J, Handschuh M, Bahr M, Richter DW, Kirchhoff F, Hülsmann S (2006) Lack of the Kir4.1 channel subunit abolishes K⁺ buffering properties of astrocytes in the ventral respiratory group: impact on extracellular K⁺ regulation. *J Neurophysiol* 95(3): 1843-52

Kvachnina E, Liu G, Dityatev A, Renner U, Dumuis A, Richter DW, Dityateva G, Schachner M, Voyno-Yasenetskaya TA, Ponimaskin EG (2005) 5-HT₇ receptor is coupled to G_α subunits of heterotrimeric G₁₂-protein to regulate gene transcription and neuronal morphology. *J Neurosci* 25(34): 7821-30

Büsselberg D, Bischoff AM, Richter DW (2003) A combined blockade of glycine and calcium-dependent potassium channels abolishes the respiratory rhythm. *Neuroscience* 122(3): 831-41

Gomez J, Ohno K, Hülsmann S, Armsen W, Eulenburg V, Richter DW, Laube B, Betz H (2003) Deletion of the mouse glycine transporter 2 results in a hyperekplexia phenotype and postnatal lethality. *Neuron* 40(4): 797-806



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Silvio O. Rizzoli

Group Leader STED Microscopy of Synaptic Function

- 2000 - 2004 Research assistant with William Betz at the Dep. of Physiology and Biophysics, University of Colorado Health Sciences Center (USA)
- 08/2004 PhD degree (Physiology) awarded by the University of Colorado
- 2004 - 2007 Post doctoral fellow with Reinhard Jahn at the Neurobiology Department of the Max Planck Institute for Biophysical Chemistry in Göttingen (Germany)
- since 2007 Group Leader (STED Microscopy) at the European Neuroscience Institute Göttingen (ENI-G)

Major Research Interests

Conventional fluorescence microscopy is limited by the diffraction of light: fluorescent objects that are close together cannot be discerned. Stimulated emission depletion (STED) is a recent advancement in optical physics that breaks the diffraction barrier, allowing microscopes to obtain much clearer images.

The diffraction barrier has been particularly problematic for imaging synaptic vesicles, which are among the smallest known organelles (30-50 nm in diameter). They are located in small areas in the synapses (about 1 micron in diameter). The group takes advantage of the increased imaging resolution provided by STED to investigate synaptic vesicle function, with an emphasis on synaptic vesicle recycling. Since STED microscopy also allows imaging of protein domains, the group aims at studying the patterning of protein domains in the synapse, in order to understand its molecular architecture.

Selected Recent Publications

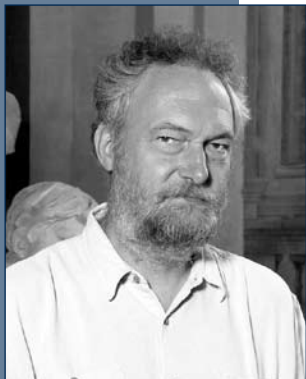
Bethani I, Lang T, Geuman U, Sieber JJ, Jahn R, Rizzoli SO (2007) The specificity of SNARE pairing in biological membranes is mediated by both proof-reading and spatial segregation. *EMBO J*, In press

Rizzoli SO, Bethani I, Zwilling D, Wenzel D, Siddiqui TJ, Brandhorst D, Jahn R (2006) Evidence for early endosome-like fusion of recently endocytosed synaptic vesicles. *Traffic* 7(9):1163-76

Gaffield MA, Rizzoli SO, Betz WJ (2006) Mobility of synaptic vesicles in different pools in resting and stimulated frog motor nerve terminals. *Neuron* 51(3): 317-25

Willig KI, Rizzoli SO, Westphal V, Jahn R, Hell SW (2006) STED microscopy reveals that synaptotagmin remains clustered after synaptic vesicle exocytosis. *Nature* 440(7086): 935-9

Rizzoli SO, Betz WJ (2005) Synaptic vesicle pools. *Nat Rev Neurosci* 6(1): 57-69. Review



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Detlev Schild

Professor of Physiology

- 1979 Diplom in Physics, University of Göttingen
- 1982 M.D., University of Göttingen
- 1985 Dr. rer.nat., University of Göttingen
- 1987 Dr. med., University of Göttingen
- 1997 Appointed head of the Department of Molecular Neurophysiology in the Center of Physiology and Pathophysiology, Medical School, University of Göttingen

Major Research Interests

We are trying to understand how the sense of smell works. Olfactory systems are able to detect and distinguish thousands of molecules in our environment. Receptor neurons are endowed with hundreds of different receptor molecules to bind odorants and transduce the chemical signals into electrical ones. Chemosensory information is thus represented in a rather high-dimensional space. The receptor neurons, which code the hitting probability of odor molecules binding to their molecular receptors, eventually generate trains of action potentials, a one-dimensional vector of stochastic processes. They convey their information onto the brain, in particular the olfactory bulb, where the receptor neuron signals are transformed into a two-dimensional neuronal image of firing activities. Glomerula, small skeins of receptor nerve fibers and synapses in the olfactory bulb, appear to be the heart of olfactory coding.

Using a combination of electrophysiological techniques, single molecule detection, photochemical and high resolution imaging techniques as well as computational and modeling methods, we are studying the biophysical and physicochemical details of

- the primary coding processes,
- the synaptic transmission in glomerula
- the generation of the neuronal chemotopic map as well as
- the processes and mechanism of odor learning and memory.

Selected Recent Publications

Czesnik D, Schild D, Kuduz J, Manzini I (2007) Endocannabinoid actions in the olfactory epithelium. *Proc Natl Acad Sci USA* 104: 2967-2972

Chen T-W, Lin B-J, Brunner E, Schild D (2006) (CMPB, BCCN) *In-situ* background estimation in quantitative fluorescence imaging. *Biophys J* 90: 2534 - 2547

Nezlin LP, Schild D (2005) Individual olfactory sensory neurons project into more than one glomerulus in *Xenopus laevis* tadpole olfactory bulb. *J Comp Neurol* 481: 233-9

Manzini I, Schild D (2004) Classes and narrowing selectivity of olfactory receptor neurons of *Xenopus laevis* tadpoles. *J. Gen Physiol* 123: 99 - 107

Manzini I, Schild D (2003) cAMP-independent olfactory transduction of amino acids in *Xenopus laevis* tadpoles. *J Physiol* 551: 115-123

Czesnik D, Rössler W, Kirchner F, Gennerich A, Schild D (2003) Neuronal representation of odorants in the olfactory bulb of *Xenopus laevis* tadpoles. *Eur J Neurosci* 17: 113-118

Gennerich A, Schild D (2002) Anisotropic diffusion in mitral cell dendrites of *Xenopus laevis* tadpoles *Biophys J* 83: 510-522



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Oliver Schlüter

Group Leader Molecular Neurobiology

- 1995 - 2001 M.D. Ph.D. with Thomas C. Südhof at the Max-Planck-Institute for Experimental Medicine in Göttingen (Germany)
- Dr. rer. nat. (PhD) 2000, University of Hannover
- Dr. med. (Medical thesis), University of Göttingen
- 2002 - 2006 Postdoc with Robert C. Malenka at Stanford University Medical Center (USA)
- Independent group leader (Emmy-Noether/DFG) at the European Neuroscience Institute Göttingen (ENI-G), since 2006

Major Research Interests

Activity-dependent modulations of synaptic transmission are important mechanisms of information processing and storage in neuronal circuits. A variety of related but mechanistically distinct forms of synaptic plasticity have been described in *in vitro* preparations of brain slices.

A major goal of my laboratory is to elucidate the underlying molecular events, leading to and regulating changes in synaptic efficacy. Newly developed techniques of molecular replacement, using mouse genetics and/or viral-mediated gene transfer allow us to manipulate the molecular composition of single neurons in a spatial and temporal controlled manner.

In particular, we are able to investigate the effects of heterologously expressed proteins on the background of wild-type neurons, or neurons, in which the endogenous protein expression is diminished. We combine this technique with simultaneous dual whole cell patch clamp recordings from rodent brain slices to monitor changes in synaptic efficacy in the manipulated cell in comparison to the neighboring control cell.

Knowledge gained from the understanding of molecular mechanisms of synaptic transmission and plasticity will ultimately provide important clues for the function of neuronal circuits and potentially the functioning of the brain

Selected Recent Publications

Schlüter* OM, Xu* W, Malenka RC (2006) Alternative N-terminal domains of PSD-95 and SAP97 govern activity-dependent regulation of synaptic AMPA receptor function. *Neuron* 51(1): 99-111

Schlüter OM, Basu J, Südhof TC, Rosenmund C (2006) Rab3 superprimes synaptic vesicles for release: implications for short-term synaptic plasticity. *J Neurosci* 26(4): 1239-46

Chandra S, Gallardo G, Fernandez-Chacon R, Schlüter OM, Südhof TC (2005) Alpha-synuclein cooperates with CSPalpha in preventing neurodegeneration. *Cell* 123(3): 383-96

Fornai F, Schlüter OM, Lenzi P, Gesi M, Ruffoli R, Ferrucci M, Lazzeri G, Busceti CL, Pontarelli F, Battaglia G, Pellegrini A, Nicoletti F, Ruggieri S, Paparelli A, Südhof TC (2005) Parkinson-like syndrome induced by continuous MPTP infusion: convergent roles of the ubiquitin-proteasome system and alpha-synuclein. *PNAS* 102(9): 3413-8

Schlüter OM, Schmitz F, Jahn R, Rosenmund C, Südhof TC (2004) A complete genetic analysis of neuronal Rab3 function. *J Neurosci* 24(29): 6629-37

Schlüter OM, Fornai F, Alessandri MG, Takamori S, Geppert M, Jahn R, Südhof TC (2003) Role of alpha-synuclein in 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine-induced parkinsonism in mice. *Neuroscience* 118(4): 985-1002

Schlüter* OM, Khvotchev* M, Jahn R, Südhof TC (2002) Localization versus function of Rab3 proteins. Evidence for a common regulatory role in controlling fusion. *J Biol Chem* 277(43): 40919-29

Schlüter OM, Schnell E, Verhage M, Tzonopoulos T, Nicoll RA, Janz R, Malenka RC, Geppert M, Südhof TC. Rabphilin knock-out mice reveal that rabphilin is not required for rab3 function in regulating neurotransmitter release. *J Neurosci*. 1999; 19(14):5834-46



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Jörg B. Schulz

Professor of Restorative Neurobiology, Director of the Department of Neurodegeneration and Restorative Research

- MD, University of Cologne Medical School, 1991
- Training in Neurology and Neuroscience at the Department of Neurology in Tübingen
- DFG Research Fellow at the Massachusetts General Hospital and Harvard Medical School, Boston
- Head of Neurodegeneration Laboratory, Hertie Institute for Clinical Brain Research and University of Tübingen, 1998 - 2004
- Habilitation, University of Tübingen, 1999
- Director of the Department of Neurodegeneration and Restorative Research, CMPB, University of Göttingen, since 2004

Major Research Interests

Our Department studies the mechanisms of degeneration in neurodegenerative disorders, including Parkinson's disease, Alzheimer's disease and cerebral ataxias. Because increased age is the major risk factor for developing a neurodegenerative disorder, we are highly interested in the mechanisms of neuronal aging. To study these mechanisms we use immortalized cell line models, primary neuronal culture models, Drosophila models, toxin-induced and transgenic mammalian (mice, rats, primates) models of Parkinson's disease. Once important pathogenetic steps have been identified we investigate their functional significance by using pharmacological or molecular tools including different transfection methods and viral gene transfer. The ultimate goal is to translate these findings into treatments that are applicable to patients. Therefore, the Department is enrolled in the outpatient clinics for Movement Disorders and Dementias and has a leading role in the German Network for Hereditary Movement Disorders (GeNeMove).

Selected Recent Publications

Strauss K, Martins LM, Plun-Favreau H, Marx F, Kautzmann S, Berg D, Gasser T, Wszolek Z, Müller T, Bornemann A, Wolburg H, Downward J, Riess O, Schulz JB, Krüger R (2005) Loss of function mutations in the gene encoding Omi/HtrA2 in Parkinson's disease. *Hum Mol Genet* 14: 2099-2111

Beier CP, Wischhusen J, Gleichmann M, Gerhardt E, Pekanovic A, Krueger A, Taylor V, Suter U, Krammer PH, Endres M, Weller M, Schulz JB (2005) FasL (CD95L/APO-1L) resistance of neurons mediated by phosphatidylinositol 3-kinase-Akt/protein kinase B-dependent expression of lifeguard/neuronal membrane protein 35. *J Neurosci* 25: 6765-6774

Luft AR, Buitrago MM, Ringer T, Dichgans J, Schulz JB (2004) Motor skill learning depends on protein synthesis in motor cortex after training. *J Neurosci* 24: 6515-6520

Simons M, Schwärzler F, Lütjohann D, von Bergmann K, Beyreuther K, Dichgans J, Wormstall H, Hartmann T, Schulz JB (2002) Treatment with simvastatin in normocholesterolemic patients with Alzheimer's disease: a 26-week randomised, placebo-controlled, double-blind trial. *Ann Neurol* 52: 346-350

Simons M, Krämer E-M, Macchi P, Rathke-Hartlieb S, Trotter J, Nave K-A, Schulz JB (2002) Overexpression of the Myelin Proteolipid Protein leads to accumulation of cholesterol and Proteolipid Protein in endosomes/lysosomes: implications for Pelizaeus-Merzbacher disease. *J Cell Biol* 157: 327-336

Wick A, Wick W, Waltenberger J, Weller M, Dichgans J, Schulz JB (2002) Neuroprotection by hypoxic preconditioning requires sequential activation of vascular endothelial growth factor receptor and Akt. *J Neurosci* 22: 6401-6407

Xia XG, Harding T, Weller M, Uney JB, Schulz JB (2001) Gene transfer of the JNK interacting protein-1 protects dopaminergic neurons in the MPTP model of Parkinson's disease. *Proc Natl Acad Sci USA* 98: 10433-10438



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Mikael Simons

Group Leader Membrane Biology (Molecular and Cellular Neurobiology)

- 1991-1997 Medical School, University of Heidelberg
- 1993-1996 MD thesis (Laboratory of K. Beyreuther, ZMBH, University of Heidelberg)
- 1997-1999 Residency in Neurology, Department of Neurology, University of Tübingen
- 1999-2000 Post-Doc (Laboratory of J. Trotter, Department of Neurobiology, University of Heidelberg)
- 2000-2004 Residency in Neurology, Department of Neurology, University of Tübingen
- 2004 Facharzt/Specialty qualification in Neurology
- 2005 Habilitation in Neurology, University of Tübingen
- 2004 Junior group leader, Centre for Biochemistry and Molecular Cell Biology, University of Göttingen
- Junior research group leader (SFB 523), Max Planck Institute for Experimental Medicine

Major Research Interests

Mechanisms of myelin biogenesis; neuron and glia interactions; membrane trafficking in oligodendrocytes; mechanisms of remyelination in multiple sclerosis; amyloid precursor protein processing in Alzheimer's disease

Selected Recent Publications

Trajkovic K, Hsu C, Chiantia S, Rajendran L, Wenzel D, Wieland F, Schwille P, Brügger B, Simons M (2008) Ceramide triggers budding of exosome vesicles into multivesicular endosomes. *Science* 319(5867): 1244-7. PMID: 18309083 [PubMed - in process]

Trajkovic K, Dhaunchak A S, Goncalves J, Wenzel D, Bunt G, Nave K A, Simons M (2006) Neuron to glia signalling triggers myelin membrane exocytosis from endosomal storage sites. *J Cell Biol* 172: 937-48

Fitzner D, Schneider A, Kippert A, Möbius W, Willig K I, Hell S W, Bunt G, Gaus K, Simons M (2006) Myelin basic protein-dependent plasma membrane reorganization in the formation of myelin. *EMBO J* 25(21): 5037-48

Simons M, Schwärzler F, Lütjohann D, von Bergmann K, Beyreuther K, Dichgans J, Wormstall H, Hartmann T, Schulz J B (2002) Treatment with simvastatin in normocholesterolemic patients with Alzheimer's disease: a 26-week randomised, placebo-controlled, double-blind trial. *Annals of Neurology* 52: 346-350

Fassbender K, Simons* M, Bergmann C, Stroick M, Lütjohann D, Keller P, Runz H, Kühl S, Bertsch T, von Bergmann K, Hennerici M, Beyreuther K, Hartmann T (2001) Simvastatin strongly reduces levels of Alzheimer's disease amyloid peptides A β 40 and A β 42 *in vitro* and *in vivo*. *Proc Natl Acad Sci USA* 98: 5856-5861; *equal contribution to first authorship

Simons M, Krämer E M, Thiele C, Stoffel W, Trotter J (2000) Assembly of myelin by association of the proteolipid protein to galactosylceramide and cholesterol rich membrane domains. *J Cell Biol* 151: 143-153



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Judith Stegmüller

Group leader, MPI for Experimental Medicine

- 1998 Diploma, University of Heidelberg
- 2002 Ph.D. University of Heidelberg
- 2003 - 2008 Postdoc, Harvard Medical School, Boston
- Since 2008 Independent group leader at the Max-Planck-Institute of Experimental

Major Research Interests

Growing evidence implicates intrinsic mechanisms such as the ubiquitin proteasome systems (UPS) in brain development and disease. Our focus lies on the role of the UPS in axon growth and regeneration. We are particularly interested how E3 ubiquitin ligases regulate these processes. To further enhance our understanding of the UPS in the central nervous system, we are also seeking to identify novel brain-specific E3 ligases and to determine their role in various aspects of neuronal development.

To address these research objectives, we apply molecular and cell biological and biochemical techniques. We also use mouse models to gain comprehensive insight into the ligases of interest and to complement *in vitro* studies with meaningful *in vivo* experiments.

Selected Recent Publications

Stegmüller J, Huynh MA, Yuan Z, Konishi Y, Bonni A (2008) TGFbeta-Smad2 signaling regulates the Cdh1-APC/SnoN pathway of axonal morphogenesis. *J Neurosci.* Feb 20;28(8): 1961-9

Stegmüller J, Konishi Y, Huynh MA, Yuan Z, Dibacco S, Bonni A (2006) Cell-intrinsic regulation of axonal morphogenesis by the Cdh1-APC target SnoN, *Neuron* 50(3): 389-400

Lasorella A, Stegmüller J, Rothschild G, Gardavaccaro D, de la Torre-Ubieta L, Pagano M, Bonni A, Iavarone A (2006) Degradation of Id2 by the anaphase promoting complex couples control of cell cycle exit and axonal growth, *Nature* 442(7101): 471-4

Stegmüller J, Bonni A (2005) Moving past proliferation: new roles for Cdh1-APC in postmitotic neurons, *Trends Neurosci.* 28(11): 596-601

Konishi Y, Stegmüller J, Mastuda T, Bonni S, Bonni A (2004) Cdh1-APC controls axonal outgrowth and patterning in the mammalian brain, *Science* Feb13;303(5660): 1026-30



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Professor, Director of the Department of Medical Psychology and Medical Sociology

- 1993: Professor of Medical Psychology, Institute of Medical Psychology (IMP), Munich University (LMU)
- 1998 - 2002 Vice-chairperson of the German Society of Medical Psychology
- since 1998 editorship of the section "Quality of life and disease coping" of the "Zeitschrift für Medizinische Psychologie"
- 1999 Professor of the Dorothea-Erxleben Foundation, Magdeburg University
- 2001 Associate Professor of Gerontopsychology at Geneva University and Head of the Department of Neurogerontopsychology at the Unit of Psychogeriatrics at Geneva University Hospital
- 2001 - 2005 Member of the board of the Swiss Society of Psychology
- 2004 Director of the Department of Medical Psychology, Georg August University of Göttingen
- 2004 - 2005 Member of the board and vice-treasurer of the Academia Multidisciplinaria Neurotraumatologica
- since 2004 editor of the series "Psychomed Compact", UTB textbooks series
- 2005 Director of the Department of Medical Psychology and Medical Sociology, Georg August University of Göttingen

Major Research Interests

Medical Psychology

- Cross-cultural Outcome
- Cognitive Neuroscience
- Neuropsychology
- Quality and communication improvement in medicine

Medical Sociology

- Assessment of the Consequences of Technology in Medicine
- Professionalisation

Selected Recent Publications

Bruggimann L, Annoni JM, Staub F, v. Steinbüchel N, van der Linden M, Bogousslavsky J (2006) Chronic posttraumatic stress symptoms after nonsevere stroke. *Neurology* 66(4), 513-516

v. Steinbüchel N, Lischetzke T, Gurny M, Eid M (2006) Assessing quality of life in older people: Psychometric properties of the WHOQOL-BREF. *European Journal of Ageing* 3, 116-122

v. Steinbüchel N, Petersen C, Bullinger M, and the QOLIBRI Group (2005) Assessment of health-related quality of life in persons after traumatic brain injury – development of the Qolibri, a specific measure. *Acta Neurochirurgica* 93, 43-49

v. Steinbüchel N, Richter S, Morawetz C, Riemsma R (2005) Assessment of subjective health and health-related quality of life in persons with acquired or degenerative brain injury. *Current Opinion in Neurology* 18, 681-691

Wittmann M, Burtscher A, Freis W, von Steinbüchel N (2004) Effects of brain-lesion size and location on temporal-order judgement. *Neuroreport*, 15 (15): 2401-2405

Kagerer F, Wittmann M, Szlag E, v. Steinbüchel N (2002) Cortical involvement in temporal reproduction: Evidence for differential roles of the hemispheres. *Neuropsychologia* 40 (3), 357-66

Wittmann M, v. Steinbüchel N, Szlag E (2001) Hemispheric specialisation for self-paced motor sequences. *Cognitive Brain Research* 10 (3), 341-344



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- M.D., 1972, Bulgarian Medical Academy
- Research Associate in Neurochemistry, 1973-1988, Bulgarian Academy of Sciences, Sofia
- Ph.D. (Neurochemistry), 1985, Bulgarian Academy of Sciences, Sofia
- Postdoc, 1988-1989, Max Planck Institute for Biophysical Chemistry, Göttingen
- Habilitation (Neurochemistry), 1989, Bulgarian Academy of Sciences, Sofia
- Assistant Research Professor, 1989 -1991, Institute of Molecular Biology, Bulg. Acad. Sci., Sofia
- Senior Research Scientist, 1991-2002, Dept. Mol. Cell Biol., MPI for Biophysical Chemistry, Göttingen
- Recognition of Habilitation (Developmental Biology), Faculty of Medicine, University of Göttingen
- Since 2002 Research Group Leader, Max Planck Institute for Biophysical Chemistry, Göttingen

Major Research Interests

Composed of six cellular layers, the mammalian neocortex is a modular structure with many functional areas in which the neurons have specific morphology, number, connections and unique physiological properties. Our group is interested in understanding the molecular and cellular mechanisms involved in specification of the immense diversity of the cortical neurons in order to be generated in a correct time, number and place during development. We have recently identified sets of genes with a differential expression between distinct domains and layers of the embryonic mouse cortex. To study the function of selected candidates in the transcriptional control of neurogenesis, we combine approaches for targeted gene inactivation or gene activation in transgenic mice using the conventional and conditional knock-out strategies with biochemical, morphological, gene expression, tissue culture methods and techniques for gene transfer in isolated brain or living mouse embryos.

With one gene, the transcription factor Pax6, we are further ahead in understanding its function. Pax6 is a critical gene for neocortical development, endowing the pluripotent radial glial progenitors with neurogenic ability and controlling the cortical patterning, including layer and area formation. Our current research focuses in unraveling genetic mechanisms by which Pax6 regulates these developmental processes with a special emphasis on its role in the control of neuronal subtype identity. We address these questions by studying the function of genes recently identified by us to act as Pax6 targets or Pax6 protein partners controlling its neurogenic function. We further aim to get insight into Pax6 dependent mechanisms involved in generation of stem/progenitors cells and their regenerative properties in neurogenic zones of the adult brain.

Selected Recent Publications

Berger J, Berger S, Tuoc TC, D'Amelio M, Cecconi F, Gorski JA, Jones KJ, Gruss P, Stoykova A (2007) Conditional activation of Pax6 in developing cortex of transgenic mice causes progenitor apoptosis. *Development* 134: 1311-1322

Fimia GM, Stoykova A, Romagnoli A, Giunta L, Di Bartolomeo S, Nardacci R, Corazzari M, Fuoco C, Ucar A, Schwartz P, Gruss P, Pieacentini M, Chowdhury K, Cecconi F (2007) AMBRA1 regulates autophagy and development of the nervous system. *Nature* 447: 1121-1125

Mühlfriedel S, Kirsch F, Gruss P, Chowdhury K, Stoykova A (2007) Large scale microarray analysis of differential gene expression of the E16 mouse cerebral cortex. *Eur J Neurosci* 26: 33-50

Remedios R, Huilgol D, Saha B, Hari P, Bhatnagar L, Kowalczyk T, Hevner RH, Suda Y, Aizawa S, Ohshima T, Stoykova A, Tole S (2007) A novel stream of amygdaloid cells from the caudal telencephalon reveals a developmental link between the amygdala and the neocortex. *Nature Neurosci* 9: 1141-1150

Zembrzycki A, Griesel G, Stoykova A, Mansouri A (2007) Genetic interplay between the transcription factor *Sp8* and *Emx2* in the patterning of the forebrain. *Neural Development* 30 2: 8



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Walter Stühmer

Professor of Neurophysiology, Director at the Max Planck Institute for Experimental Medicine

- 1978 - 1980 PhD with Dr. F. Conti in Camogli, Italy
- 1980 - 1983 Post Doc in the Department of Physiology and Biophysics in Seattle, USA, with Dr. W. Almers
- 1983 - 1992 group leader at the Max Planck Institute for Biophysical Chemistry in Göttingen with Dr. E. Neher
- 1992 - present Director of the Department Molecular Biology of Neuronal Signals at the Max Planck Institute for Experimental Medicine in Göttingen

Major Research Interests

The principal aim of the department "Molecular Biology of Neuronal Signals" is the study of signaling within cells and between cells. To this end, molecular biology, genetics and electrophysiology are used to elucidate structure-function relationships of membrane-bound proteins, especially ion channels and receptors. Specific tools such as antibodies and toxins are developed and used to interfere with signaling pathways relevant for cell cycle control, ion selectivity and the secretion of cells in culture and in primary cells.

Selected Recent Publications

Herrero-Herranz E, Pardo LA, Bunt G, Gold R, Stühmer W, Linker RA (2007) Re-expression of a developmentally restricted potassium channel in autoimmune demyelination: Kv1.4 is implicated in oligodendroglial proliferation. *Am J Pathology* 171: 589-598

Stühmer W, Alves F, Hartung F, Zientkowska M, Pardo LA (2006) Potassium channels as tumour markers. *FEBS Letters* 580: 2850-2852

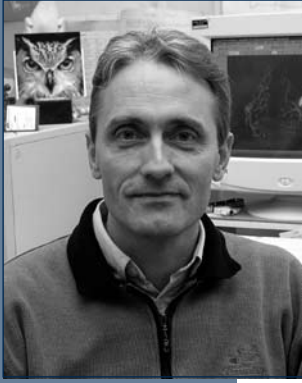
Weber C, Mello de Queiroz F, Downie F, Suckow A, Stühmer W, Pardo LA (2006) Silencing the activity and proliferative properties of the human Eag1 potassium channel by RNA interference. *J Biol Chem* 281: 13030-13037

Pardo LA, Contreras-Jurado C, Zientkowska M, Alves F, Stühmer W (2005) Role of voltage-gated potassium channels in cancer. *J Membr Biol* 205: 115-124

García-Ferreiro RE, Kerschensteiner D, Major F, Monje F, Stühmer W, Pardo LA (2004) Mechanism of block of hEag1 K⁺ channels by imipramine and astemizole. *J Gen Physiol* 124: 301-317

Becherer U, Moser T, Stühmer W, Oheim M (2003) Calcium regulates exocytosis at the level of single vesicles. *Nature Neurosci* 6: 846-853

Jenke M, Sánchez A, Monje F, Stühmer W, Weseloh RM, Pardo LA (2003) C-terminal domains implicated in the functional surface expression of potassium channels. *EMBO J* 22: 395-403



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Andreas Stumpner

Professor of Neuroethology

- Dr. rer. nat., University of Erlangen, Germany, 1988
- Postdoctoral fellow, Andrews University, Berrien Springs, USA, 1990 - 1991
- Habilitation, University of Göttingen, 1997
- Guest professor, University of Zurich, Switzerland, 2002 - 2003
- Since April 2003 Professor of Zoology at the University of Göttingen

Major Research Interests

My research focuses on how a small nervous system recognises specific frequencies and temporal patterns (in the context of acoustic communication in insects, mainly in Orthoptera). Understanding these processes bears implications also for understanding function and evolution of the same performances of the vertebrate brain. I see the strength of the acoustic and invertebrate system *a)* in the precise temporal and spectral stimuli one can deliver and the clear (innate) responses on the behavioural and neuronal level, *b)* in the comparative potential (song recognition in groups of related species and differences in neuronal layout to related non-singing or non-hearing groups) allowing to understand what mechanisms might have played a role in evolution and how evolution of songs and recognition systems depend on each other, *c)* in the identified neurone-approach allowing to find homologous neurones in related species and indicating evolutionary changes on the cellular level and *d)* the potential to directly test hypotheses in behavioural experiments.

Recent findings from intracellular studies in bushcrickets are: Central neurons receive lateral frequency-dependent inhibitions. After blocking such inhibitions the frequency tuning broadens considerably. Species-specificity of a neuron in related species depends on specific inhibitions, not on specific excitations. And homologous neurons in more distantly related species may differ considerably in their properties..

Selected Recent Publications

Stumpner A, Allen GR, Lakes-Harlan R (2007) Hearing and frequency dependence of auditory interneurons in the parasitoid fly *Homotrixa alleni* (Tachinidae: Ormiini). *J Comp Physiol A* 193: 113-125

Stumpner A, Molina J (2006) Diversity of intersegmental auditory neurons in a bush cricket. *J Comp Physiol A* 192: 1359-1376

Molina J, Stumpner A (2005) Effects of pharmacological treatment and photo-inactivation on the directional responses of an insect neuron. *J Exp Zool* 303A: 1085-1103

Hennig M, Franz A, Stumpner A (2004) Processing of auditory information in insect. *Microsc Res Tech* 63: 351-374

Stumpner A (2002) A species-specific frequency filter through specific inhibition, not specific excitation. *J comp Physiol A* 188: 239-248

Stumpner A (1999) Comparison of morphology and physiology of two plurisegmental sound-activated interneurons in a bushcricket. *J Comp Physiol A* 185: 199-205

Rust J, Stumpner A, Gottwald J (1999) Singing and hearing in an ancient bush-cricket. *Nature* 399: 650



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Victor Tarabykin

Group Leader at the Max Planck Institute for Experimental Medicine

- MD, Russian State Medical University, Moscow 1993
- PhD in Molecular Biology with S.Lukyanov, Russian Academy of Sciences, Moscow 1996
- Postdoctoral fellow with P.Gruss at the Max Planck Institute for Biophysical Chemistry, 1996 - 2001
- since 2002 Research Group Leader at the Max Planck Institute for Biophysical Chemistry; Department Molecular Cell Biology, Göttingen

Major Research Interests

During development, several populations of progenitor cells in the dorsal telencephalon generate a large variety of neurons. These neurons acquire distinct morphologies and physiological properties and serve distinct functions in the mammalian cerebral cortex.

We are interested in the cellular and molecular mechanisms underlying cell fate specification in the mouse cerebral cortex. We focus on the mechanisms controlling the generation of neurons of different cortical layers. We apply a combination of genetic, molecular and cell biological approaches. We have identified several genes that control cortical development. One of them, Sip1 is a transcription factor implicated in Mowat-Wilson syndrome (MWS) in humans. MWS patients suffer from intellectual disability, microcephaly and seizures. We inactivated the gene specifically in cortical precursors. This resulted in the degeneration of the entire hippocampus. We have shown that in the hippocampus Sip1 controls activity of non-canonical Wnt pathway.

Another gene we identified, Satb2 is a transcription factor of a novel type that interacts with special chromosomal regulatory elements, Matrix Attachment Regions. Satb2 is an important determinant of neurons of superficial cortical layers. In order to study its role in neural development we produced several mouse mutants where Satb2 expression is altered. There are several other genes that have been identified in the lab whose function in the cortical development remains to be revealed.

Selected Recent Publications

Miquelajauregui A, Van de Putte T, Polyakov A, Nityanandam A, Boppana S, Seuntjens E, Karabinos A, Higashi Y, Huylebroeck D, Tarabykin V (2007) Smad-interacting protein-1 (Zfhx1b) acts upstream of Wnt signaling in the mouse hippocampus and controls its formation. *Proc Natl Acad Sci U S A.* 31;104(31): 12919-24

Britanova O, Depew MJ, Schwark M, Thomas BL, Miletich I, Sharpe P, Tarabykin V (2006) Satb2 haploinsufficiency phenocopies 2q32-q33 deletions while loss suggests a fundamental role in the coordination of jaw development. *Am J Hum Genet* 79(4): 668-78

Britanova O, Alifragis P, Johnes K, Gruss P, Tarabykin V (2006) Tangential migration of cortical projection neurons: a novel mode of migration. *Dev Biol* 298(1): 299-311

Guillemot F, Molnar Z, Tarabykin V, Stoykova A (2006) Molecular mechanisms of cortical differentiation. *Eur J Neurosci* 23(4): 857-68

Molnar Z, Metin C, Stoykova A, Tarabykin V, Price D, Frances F, Meyer G, Dehay C, Kennedy K (2006) Comparative aspects of cerebral cortical development. *Eur J Neurosci* 23(4): 921-34

Britanova O, Akopov S, Lukyanov S, Gruss P, Tarabykin V (2005) Novel transcription factor Satb2 interacts with matrix attachment region DNA elements in a tissue-specific manner and demonstrates cell-type-dependent expression in the developing mouse CNS. *Eur J Neurosci* 21: 658-68

Tarabykin V, Stoykova A, Usman N, Gruss P (2001) Cortical upper layer neurons derive from the subventricular zone as indicated by Svet1 gene expression. *Development* 128: 1983-1993



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Stefan Treue

Professor, Director of the German Primate Center

- Head of the Cognitive Neuroscience Laboratory
- Ph.D. 1992, Massachusetts Institute of Technology
- Postdoctoral Fellow, MIT, 1992 - 1993
- Postdoctoral Fellow, Baylor College of Medicine, Houston, Texas, 1993 - 1995
- Work Group Leader, Laboratory of Cognitive Neuroscience, University of Tübingen, 1995 - 2001
- Professor of Animal Physiology, University of Tübingen, 2000 - 2001
- Professor of Cognitive Neuroscience and Biological Psychology, University of Göttingen, 2001

Major Research Interests

Research at the Cognitive Neuroscience Laboratory is aimed at understanding the neural basis of visual perception. Vision is an active process that is far more than a passive registration of our environment. Rather, on its way from the eyes to and through the cortex, visual information is modulated by numerous processes that enhance some aspects while diminishing others. One of these processes is attention, i.e. the ability to filter out unwanted information and concentrate the brain's processing abilities on relevant information.

The accurate representation of visual motion in the environment is one of the most important tasks of the visual system. Correspondingly, research in the laboratory concentrates on this ability as a model for sensory information processing in general.

We use various techniques. While our emphasis is on electrophysiology, i.e. the recording of the activity of neurons in the visual cortex of macaque monkeys and measuring human perceptual abilities with psychophysical methods, we also use theoretical approaches and functional brain imaging.

Using these techniques, we have been able to elucidate how motion information is represented in primate cortical area MT and how attention changes that representation and correspondingly the percept of the visual environment.

Selected Recent Publications

Womelsdorf T, Anton-Erxleben K, Pieper F, Treue S (2006) Dynamic shifts of visual receptive fields in cortical area MT by spatial attention. *Nature Neuroscience* 9: 1156-1160

Martinez-Trujillo JC, Treue S (2004) Feature-based attention increases the selectivity of population responses in primate visual cortex. *Current Biology* 14: 744-751

Martinez-Trujillo JC, Treue S (2002) Attentional modulation strength in cortical area MT depends on stimulus contrast. *Neuron* 35: 365-370

Treue S (2001) Neural correlates of attention in primate visual cortex. *Trends in Neurosciences* 24 (5): 295-300

Treue S, Hol K, Rauber HJ (2000) Seeing multiple directions of motion - Physiology and psychophysics. *Nature Neuroscience* 3 (3): 270-276

Treue S, Martinez Trujillo JC (1999) Feature-based attention influences motion processing gain in macaque visual cortex. *Nature* 399 (6736): 575-579

Treue S, Maunsell JHR (1996) Attentional modulation of visual motion processing in cortical areas MT and MST. *Nature* 382 (6591): 539-541



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Andreas Wodarz

Professor of Stem Cell Biology

- Diploma Biology, University of Cologne, 1990
- Dr. rer. nat. Developmental Biology, University of Cologne, 1993
- Postdoc, Howard Hughes Medical Institute, Stanford University, 1994 - 1997
- Junior Group Leader, Heinrich Heine University Düsseldorf, 1997 - 2004
- Habilitation in Genetics, Heinrich Heine University Düsseldorf, 2001
- Appointed as Head of the Department of Stem Cell Biology at the University of Göttingen, 2004

Major Research Interests

At the center of my research interests is the question of how neural stem cells divide asymmetrically to produce another stem cell and a progenitor cell that will differentiate and give rise to neurons and glia cells. One important aspect of asymmetric cell division is the establishment of an intrinsic polarity which is the prerequisite for the asymmetric localization of proteins and mRNAs that serve as cell fate determinants. Our model system for the asymmetric division of stem cells is the embryonic neuroblast of *Drosophila*. Here we study the function of genes that control cell polarity, asymmetric localization of cell fate determinants and orientation of the mitotic spindle. The knowledge obtained in the *Drosophila* system has stimulated intense research on the participation of the orthologous genes and proteins in the asymmetric division of vertebrate stem cells.

Selected Recent Publications

Wodarz A, Stewart DB, Nelson WJ, Nusse R (2006) Wingless signaling modulates cadherin-mediated cell adhesion in *Drosophila* imaginal disc cells. *J Cell Sci* 119: 2425-2434

Wodarz A (2005) Molecular control of cell polarity and asymmetric cell division in *Drosophila* neuroblasts. *Curr Opin Cell Biol* 17: 475-481

von Stein W, Ramrath A, Grimm A, Müller-Borg M, Wodarz A (2005) Direct association of Bazooka/PAR-3 with the lipid phosphatase PTEN reveals a link between the PAR/aPKC complex and phosphoinositide signaling. *Development* 132: 1675-1686

Wodarz A, Ramrath A, Grimm A, Knust E (2000) *Drosophila* atypical protein kinase C associates with Bazooka and controls polarity of epithelia and neuroblasts. *J Cell Biol* 150: 1361-1374

Wodarz A, Ramrath A, Kuchinke U, Knust E (1999) Bazooka provides an apical cue for Inscuteable localization in *Drosophila* neuroblasts. *Nature* 402: 544-547



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Fred Wolf

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- Head of the Research Group “Theoretical Neurophysics”, Department of Nonlinear Dynamics, Max-Planck-Institut für Strömungsforschung, Göttingen, since 2004.
- Visiting Scholar, Kavli Institute for Theoretical Physics, UC Santa Barbara (USA), Fall 2001, 2003, 2004
- Research Associate, Max-Planck-Institut für Strömungsforschung, Göttingen, 2001 - 2004
- Amos de Shalit Fellow, Racah Institute of Physics and Interdisciplinary Center for Neural Computation, Hebrew Univ., Jerusalem (Israel), 2000
- Dr. phil. nat., J.W. Goethe Universität, Frankfurt , 1999

Major Research Interests

- Theoretical neuroscience and nonlinear dynamics
- Dynamics and synchronization in cortical neural networks
- Function and development of the visual cortex
- Sensory processing in the auditory system

The brains of humans and animals arguably are among the most complex systems in nature. Over the past decade, theoretical neuroscience - the use of quantitative theories, mathematical modelling and advanced quantitative data analysis methods for the study of brain function - has started to provide powerful new approaches for understanding the neuronal basis of perception, learning, memory, and other higher brain functions. This is because, even during the neuronal processing of the most elementary sensory stimulus large ensembles of interacting nerve cells distributed throughout the brain are activated, the collective operations of which are often hard to understand by means of purely qualitative reasoning.

The primary focus of our research in theoretical neuroscience is self-organization in the dynamics of cortical networks. In particular, we have developed novel approaches to model and predict the dynamics and neuronal plasticity of the visual cortex. To quantitatively connect theory and experiment in this system, we recently also designed methods that enable to quantify the organization of visual cortical functional architecture with high precision. Another important focus of our work is the mathematical analysis of the dynamics of large and complex networks of pulse-coupled neuron models. The concepts and tools for the representation of the dynamics of cortical circuits developed enable a rational and transparent design of models of higher cortical functions such as the processes underlying perceptual learning phenomena.

Selected Recent Publications

Schnabel M, Kaschube M, Loewel S, Wolf F: Europhysics Random Waves in the Brain: Symmetries and Defect Generation in the Visual Cortex. Journal (in press)

Naundorf B, Wolf F, Volgushev M (2007) Hodgkin and Huxley model - still standing? Nature 445: E2-E3

Timme M, Geisel T, Wolf F (2006) Speed of synchronization in complex networks of neural oscillators: analytic results based on Random Matrix Theory. Chaos 16: 015108, 2006

Wolf F, Naundorf B, Volgushev M (2006) Unique features of action potential initiation in cortical neurons. Nature 440(7087)

Wolf F (2005) Symmetry, Multistability, and Long-Range Interactions in Brain Development. Phys. Rev. Lett., 95: 208701

Naundorf B, Geisel T, Wolf F (2005) Action potential onset dynamics and the response speed of neuronal populations. Journal of Computational Neuroscience, 18(3): 297-309

Wolf F (2005) Symmetry Breaking and Pattern Selection in Visual Cortical Development. Methods and Models in Neurophysics, Les Houches, Session LXXX, 2003, p. 575-639, Chow CC, Gutkin B, Hansel D, Meunier C, Dalibard J (eds.), Elsevier



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Fred Wouters

Professor, Laboratory for Molecular and Cellular Systems

- Dr. (Ph. D.) 1997, Faculty of Chemistry, University of Utrecht, The Netherlands
- Postdoctoral fellow, Imperial Cancer Research Fund (ICRF), London UK, 1997 - 2000
- Postdoctoral fellow, European Molecular Biology laboratory (EMBL), Heidelberg, 2000 - 2001
- Appointed as group leader at the European Neuroscience Institute, Göttingen 2001
- PD (habilitation) 2006, Physiology, Göttingen University

Major Research Interests

The focus of our research is the regulation and role of the neuronal cytoskeleton in the modulation of neuronal shape and motility during chemotactic processes. The growing neuronal growth cone probes its environment for the chemical composition of its substrate and the presence of neighbouring cells. The former information is sampled by cell adhesion receptors in focal adhesion structures that, next to their sensing function also perform a structural function in that they provide the cell with a means to exert force on its substrate. We are primarily interested in the signal transduction processes that regulate these effects and the cross-talk between the different motility systems.

The main interest areas in this question are; 1. The role and molecular mechanism of lipid raft-resident cell adhesion molecules in the remodelling of the membrane cytoskeleton, 2. Dynamic control of growth cone protein content by local proteolysis and chaperone function during chemotactic responses, 3. Role and mechanism of the neuronal exocyst complex as critical landmarks for dendritic/axonal neuritogenesis.

Our group has a related interest in the pathophysiological mechanism of neurodegeneration by intracellular aggregation of the tau protein, as occurs in Alzheimer's disease. As tau is an intrinsically unstructured protein that can undergo remarkable conformational changes upon binding to microtubules and in the Alzheimer-related aggregation condition, it presents an ideal model system for the biophysical analysis of protein conformational change and protein interactions. Our research depends on the development and application of advanced microscopy techniques, primarily; fluorescence lifetime imaging microscopy (FLIM), and Förster resonance energy transfer (FRET) microscopy, in combination with a range of GFP-based optical biosensors and novel bioconjugation approaches for organic dyes, and protein biochemical/molecular biological techniques to resolve and quantify biochemical reactions and conditions in living cells.

Selected Recent Publications

Iliev AI, Djannatian JR, Nau R, Mitchell TJ, Wouters FS (2007) Cholesterol-dependent actin remodeling via RhoA and Rac1 activation by the *Streptococcus pneumoniae* toxin pneumolysin. *Proc Natl Acad Sci USA* 104: 2897-2902

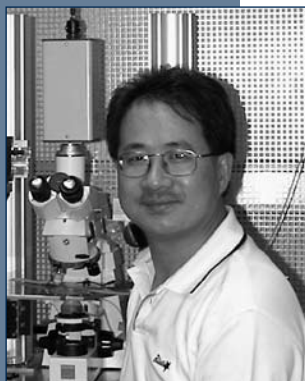
Esposito A, Dohm CP, Kermer P, Bahr M, Wouters FS (2007) alpha-Synuclein and its disease-related mutants interact differentially with the microtubule protein tau and associate with the actin cytoskeleton. *Neurobiol Dis* 26: 521-531

Esposito A, Dohm CP, Bahr M, Wouters FS (2007) Unsupervised fluorescence lifetime imaging microscopy for high content and high throughput screening *Mol Cell Proteomics* 6: 1446-1454

Hillebrand M, Verrier SE, Ohlenbusch A, Schafer A, Soling HD, Wouters FS, Gartner J (2007) Live cell FRET Microscopy: homo- and heterodimerization of two human peroxisomal ABC transporters, the adrenoleukodystrophy protein (ALDP, ABCD1) and PMP70 (ABCD3). *J Biol Chem* 282: 26997-27005

Pommereit D, Wouters FS. (2007) An NGF-induced Exo70-TC10 complex locally antagonises Cdc42-mediated activation of N-WASP to modulate neurite outgrowth. *J Cell Sci* 120: 2694-2705

Esposito A, Gerritsen HC, Wouters FS (2007) Optimizing frequency-domain fluorescence lifetime sensing for high-throughput applications: photon economy and acquisition speed. *J Opt Soc Am A* 24: 3261-3273



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- Dr. med. (M. D.) University of Bonn, 1987
- Internship, Department of Neurology, University of Bern, Switzerland, 1988
- Postdoctoral fellow, Department of Physiology, University of Bern, Switzerland, 1989 - 1994
- Postdoctoral fellow, Department of Physiology, University of Oxford, UK, 1993
- Postdoctoral fellow, The Nobel Institute of Neurophysiology, Karolinska Institute, Stockholm, Sweden, 1994 - 1996
- Research Group Leader, Center of Physiology and Pathophysiology, University of Göttingen, since 1997
- Habilitation, University of Göttingen, 2003

Major Research Interests

The neuronal developmental disorders associated with Rett syndrome, the classic autism and other autistic spectrum diseases (ASD) are correlated with a disruption of functional synaptic maturation during postnatal development. Such developmental dysregulation causes cognitive, social and motor retardations. Most ASD patients achieve normal developmental milestones until 6-18 months of age when they enter a period of regression with loss of acquired cognitive, social and motor skills. The main interest of our research group is to analyze disease-related changes of the expression of receptor subunits, the properties of ion-channels and dysfunction synaptic transmission within intact neuronal network in mutant mice models, such as MECP2, neuroligin, neurexin and neurebechin mutants. Using an integrative approach, we aim to clarify the functional consequences of identified molecular disturbances in functional synaptic maturation and identify the changes in neuromodulation. In addition, we elucidate the potency of various strategies of protection and restoration including pharmacotherapies using the mutant mice models for ASD.

Selected Recent Publications

Medrihan L, Tantalaki E, Sargsyan V, Aramuni G, Dudanova I, Missler M, Zhang W (2008) Early defects of GABAergic synapses in the brainstem of a MeCP2 mouse model of Rett syndrome. *Journal of Neurophysiology* 99 (1): 112-121

Varoqueaux F, Aramuni G, Rawson R, Mohrmann R, Gottmann K, Zhang W, Südhof TC, Brose N (2006) Neuroligins control synaptic function and network activity but not synaptogenesis. *Neuron* 51: 741-754

Zhang W, Rohlmann A, Sargsyan V, Aramuni G, Hammer R, Südhof TC, Missler M (2005) Extracellular domains of α -neurexin are important for regulating synaptic transmission by selectively affecting N- and P/Q-type Ca^{2+} -channels. *Journal of Neuroscience*. 25(17): 4330-4342

Missler M, Zhang W, Rohlmann A, Kattenstroth G, Hammer R, Gottmann K, Südhof TC (2003) α -Neurexins are Required for Coupling Ca^{2+} -Channels to Synaptic Vesicle Exocytosis. *Nature* 423: 939-948

Zhang W, Barnbrock A, Gajic S, Pfeiffer A, Ritter B (2002) Differential ontogeny of GABA_B receptor-mediated pre- and postsynaptic modulation of GABA and Glycine transmission in respiratory rhythm-generating network of mouse. *The Journal of Physiology* 540(2): 435-446

Ritter B, Zhang W (2000) The GABA_A-mediated inhibition matures during first postnatal week in brain stem of mouse. *European Journal of Neuroscience* 12: 2975-2984

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