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Advanced Biochemistry

Master of Science



EBERHARD KARLS
UNIVERSITÄT
TÜBINGEN



Welcome

Welcome to the University of Tübingen and to the Interfaculty Institute of Biochemistry (IFIB). We are happy that you chose our master program and congratulations on passing the competitive selection process.

You are now at the second stage of your university education. A period to discover and develop your preferences in the field. At the end of the Master, you will choose whether to continue along the academic track with a doctorate or to enter the job market. We strive to support you with this important decision.

You are also no longer a beginner in the field, and we believe the structure of the master's degree should strongly reflect this and support your further maturation towards a full-fledged biochemist. In our program, we have only a small number of compulsory elements, but a large variety of choices you can make. Importantly, a significant part of the program will be dedicated to preparing you for future career decisions.

We are looking forward to having you in the IFIB and wish you all the best for the start of your master program here in Tübingen.

Rober Feil, Dean of Studies

Ralf-Peter Jansen, Head of Examination Board

Markus Wolters, Study coordinator

cover picture: albarst



Gerhard Groebe

Overview

The Tübingen *Master of Advanced Biochemistry* is a consecutive, full-time two-year program composed of different modules (see table below). Students have to earn a minimum of 120 ECTS credits in total, each of which corresponds to approximately 30 hours of work including self-directed study. All elements are explained in detail later in this guide. The lecture *Advanced Biochemistry* and the seminar series *Current Topics* are theoretical modules while *courses*, *labs*, and the *master thesis* are combined theoretical and practical elements.

Modules of the Master	Credit points (ECTS)
Advanced Biochemistry	9
Current Topics	3
3x Courses	3 x 6
3x Labs	3 x 15
Advanced Qualifications	15
Master thesis	30
	120

The Tübingen *Master of Advanced Biochemistry* gives you a high level of flexibility. You do not have to take the modules in a specific order or at a specific time. Nevertheless, we recommend organizing your degree roughly according to the order above. The theoretical elements are a good starting point to gain an overview of the fast-moving field of biochemistry and are thus well placed at the beginning. They may also serve as preparation for the laboratory work later. *Courses* are designed to be a transition from theory to more practical work, since they contain components of both theory and experimental work. Attending the “Advanced Biochemistry” lectures already in your first year is also a good way of getting to know your peers. *Labs* require more organization from your part and are your stepping stone from pre-designed courses to the more independent master thesis that concludes the *Master of Biochemistry*.

Aims

Upon completion of the Master of Advanced Biochemistry in Tübingen, you will have gained significant lab experience in diverse areas of life science, mainly via three lab rotations and your master thesis. This will enable you to make a well-informed choice regarding your future career, in research or elsewhere. In contrast to fast-track programs, you will get exposed to several research areas which will allow you to pick the one that suits you the most for your further steps.

In addition to having improved your lab skills, you will have deepened and broadened your understanding of the biochemical basis of life and disease. This new knowledge will stem from the advanced lecture series, the course seminars, and your lab rotations.

Besides biochemical practice and theory, you will have improved three transfer skills: presenting, writing, and organization. You will get qualified feedback on your presentation and writing skills during all modules. You can also take optional skill courses to improve the above. The large variety of lab projects will increase your capacity to organize and exert your work.

A Master Plan

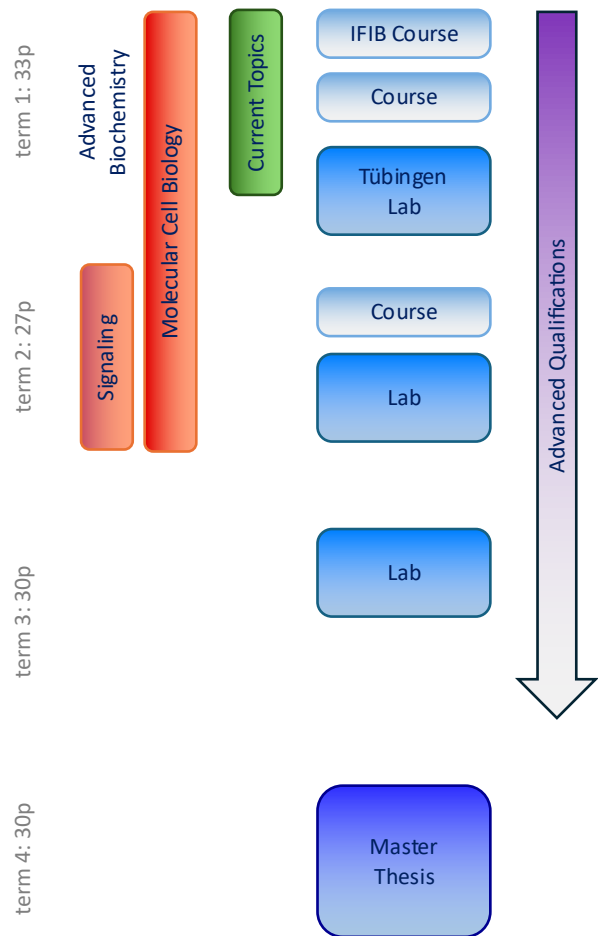
This page shows you one possible and straightforward way to complete the *Master of Advanced Biochemistry*. Many variations on this path are possible and depend on your individual preferences.

Year 1 - Introduction, Courses, first Labs

The first year kicks off with a joint lecture series by research group leaders of the institute. Departing from the basics you have acquired in your previous degree, this series takes you into the advanced topics of biochemistry. In parallel, you will choose your first *courses* and your first *lab* placements.

In parallel to the lecture series, you will take a course on current topics, which includes a series of presentations by invited national and international guest speakers presenting their latest discoveries. In addition, you will most likely select a third course and a second lab placement.

In principle, you can also attend the *Advanced Biochemistry* lectures and *Current Topics* during later semesters. However, we recommend starting early.



Year 2 - Course, Labs & Master Project

As you enter the second year, the elements of the program will become longer and more comprehensive culminating in the master thesis at the end of the degree. You may decide to do one or more of the research placements outside the institute (including abroad). We will support you with national and international placement. At the end of the last semester, you will write up the data collected during your thesis project and present the results to your host group.

During the entire degree, you will be able to acquire transferable skills ranging from numerical proficiency, use of scientific software, self-management and communication to interpersonal skills (purple track in the above scheme). Those courses are for instance offered by the University's transdisciplinary course program. Furthermore, IFIB faculty members can advise you in the decision about your future professional career.

Requirements

Please refer to our webpage (<https://www.ifib.uni-tuebingen.de/studium/master.html>) for details on our entrance requirements. In short, you need to have a bachelor in Biochemistry or a related life science with excellent results, in-depth lab experience, and fluency in English (equivalent to at least B2 or proficient user). So far, you should have executed appropriate experiments and critically evaluated the results, most commonly in your bachelor thesis. You should be self-motivated, driven, and curious.

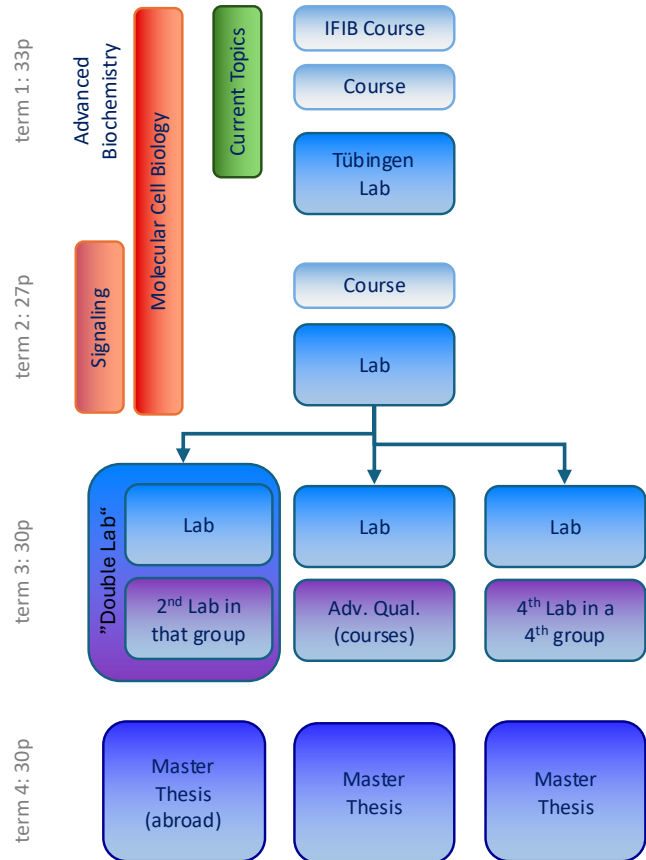
Varieties

Here are a few ideas on how to adapt the template presented on the previous page to your personal preferences. Many variations are possible. Please discuss them with us.

In case you want to delve deeper into a special topic or in case you think one lab block is not enough to gain all the experience you want in a certain topic, you can fuse a *Lab* block together with the *Advanced Qualifications* module to form a "Double Lab" accounting for 2x15 ECTS. Do not combine more than two labs in one research group to make sure that you will get exposed to a variety of research topics.

If you want to do one of your *Labs* abroad, a double lab may be useful. It takes time to organize a stay in another country. To go through all this work for a single lab element may not be efficient for you. Furthermore, the receiving group leader may be more willing to accommodate you, if you stay longer.

Another possibility to gain international experience is to do your master thesis abroad. This element of the master's degree is naturally longer, thus the ratio of preparation and lab time is better than for a single lab. On the other hand, it is an element that impacts on your final grade, so you need to be careful to select a good lab and an interesting project.



A word of caution: Flexibility vs. study progress

You have seen already that our programme is very flexible and allows you to adjust it to your needs and interests.

Keep in mind, however, that this flexibility can come at the cost of your study progress. If you plan to do lab rotations outside Tübingen, you probably need to move to another town or even country. This often requires time that you will need on top of your regular studies. Accordingly, the high degree of flexibility in our programme can result in study times longer than 4 semesters.

We have a study limit of 8 semesters (excluding official breaks caused by sickness, child care, etc), because we believe that there should be some focus on getting your degree. Nevertheless, extended or multiple stays outside Tübingen should be possible during this time. If you feel that your intended course of studies might get in conflict with the study limit, please contact the coordinator. We will find a solution.

Advanced Biochemistry (Module: BCH-M-ABC)

The lecture series *Advanced Biochemistry* (ABC) picks up where you left off with the basic lecture series of the bachelor degree and takes you deeper into advanced research topics. Reflecting this, the study material will be more recent articles and less textbooks, and the hypotheses will be more controversial and less established.

This module of the master is composed of 2 parts: Molecular Cell Biology and Signaling. The Molecular Cell Biology series runs during 2 terms (winter + summer) while the Signaling lectures are all in the summer semester.

Advanced Biochemistry is a relay of lecturers of IFIB. This lecture series will give you the chance to get to know the faculty members of biochemistry and experience their distinct research angles.

Element Overview

Goals	You acquire cutting-edge knowledge of the biochemistry governing cellular dynamics and communication. You critically evaluate recent discoveries.
Content	Extracellular matrix, cellular attachment, pathogen receptors, cellular uptake processes, endocytosis, endosomal escape mechanisms, targeting/transport in cells, organelle structure and maintenance, cytoskeleton, motor proteins, signaling pathways, oncobiology, hormones
Format	• Molecular Cell Biology (2 hours/week in winter and in summer term) • Signaling (2 hours/week in summer term)
Duration	2 semesters in total
Recommended semester	• Molecular Cell Biology: 1st + 2nd semester • Signaling: 1st or 2nd semester
Frequency	Each lectures block is offered once per year
Participants	20-35
Evaluation	Oral examination with two Biochemistry professors or lecturers, at least one from the IFIB or biochemistry section of the ZMBP, total duration: 45 min
Responsible	Dean of Studies in Biochemistry
Organizers	Professors / lecturers of IFIB
Workload	90 h contact time + 180 h self-directed study
Credit points	9 CP

Molecular Cell Biology - Advanced Biochemistry



Molecular Cell Biology (winter)	
Rapaport	Transport of proteins and lipids
Schäffer	Cytoskeleton and biophysics
Deindl	Structural biology of tyrosin kinases and chromatin

The approximate overview of topics is indicated in the scheme. Each lecturer is responsible for their block of presentations, typically taking place every Tuesday morning.

The biochemical basics of cellular transport processes will be reviewed before proceeding to the topical research questions of the field.

Molecular Cell Biology (summer)	
Filarsky	Epigenetics / Metabolic signaling / Immune evasion
Jansen	Cytoskeleton / Motors / Mitosis
Schwarzer	Visualization and Drug-targeting of Nuclear Import and Export

Unless you have started in summer, we recommend taking the winter semester lectures first, as it is partly the basis for later lectures. However, the order can also be reversed.

Details regarding the individual lectures and teaching material will be made available and updated on the learning platform ILIAS.

Signaling - Advanced Biochemistry

Signaling (summer)	
Feil	Cardiovascular Signaling
Schulze-Osthoff	Signaling in Apoptosis and Cancer
Stafforst	Nucleic Acid Signaling Nucleic Acid Therapeutics

In summer, in parallel to the 2nd part of the Molecular Cell Biology lectures, there is an additional series on the biochemistry of signaling processes.

In this block, you will get to know the remaining group leaders of the IFIB.

Signaling is a central cutting-edge field involved in most aspects of biochemistry. Together with Molecular Cell Biology they both represent very active areas of biochemical research. We hope you

will enjoy our lectures and are looking forward to your feedback.

Examination

Advanced Biochemistry is one of the graded elements of the *Master of Biochemistry*. After you have completed the lecture series, there will be an approximately 45-minute oral examination by two lecturers on general biochemistry, the lecture series, and three guest talks of external speakers of your choosing.

See <https://uni-tuebingen.de/de/46394>.

Current Topics in Biochemistry (Module: BCH-M-CT)

Biochemistry is a fast-moving field. Thus, we want to dedicate an entire element of the master's course to topical issues. In addition, as master students you have already gained some experience, so we want to discuss the various subjects in a more interactive format rather than a typical frontal lecture.

Goals	You critically summarize and present a recent technology / discovery. You give constructive feedback to improve the work of your peers.
Content	Current findings, problems, and debates in biochemistry Modern experimental approaches
Format	Various interactive seminars and presentations
Duration	2 terms
Recommended semester	1 st or 2 nd term
Frequency	• 10-12 invited speakers (typically Mondays) per term • Weekly Current topic seminars every winter term
Evaluation	Participation and attendance Your presentation will receive a score (only for your orientation).
Responsible	IFIB lecturers
Workload	30 h seminars + 60 h preparation / research
Credit points	3 CP

In one part of *Current Topics* you will give a presentation, introducing novel research, currently discussed topics in society or science, scientific ethics, sustainability or other topics. This will allow you to train your presentation skills for difficult or complex subjects and to discuss them with your fellow students and peers.

In another part of *Current Topics* you have to attend and participate in at least 20 of the weekly institute seminars (usually Mondays during semester time). Lecturers and students at the institute invite top class researchers from Germany and abroad to inform us about their recent discoveries. Attendance will give you an impression of current research, provides an opportunity to participate actively by asking questions, and find a lab rotation.

Additionally, you will choose 3 lectures of the invited speakers to be part of your ABC exam.

Courses (Modules: BCH-M-EC-BCH, BCH-M-EC2, BCH-M-EC3)

Courses are intended as both a transition from large-scale practical courses to independent research and to connect with your classmates and build your network. *Courses* last about a month and consist of about 90 hours of contact time plus 90 hours of self-directed study. They typically consist of a theoretical part in the form of lectures and/or seminars and a practical part. You will select a total of at least three *courses* and receive six credit points for each successfully completed one.

IFIB Courses (at least 1 from this pool)

Module: BCH-M-EC-BCH, -EC2 and -EC3

- Cell Biochemistry of Organelles (Rapaport)
- Cell Signaling (Feil)
- Chemical Biology (Schwarzer)
- Post-transcriptional Control / RNA (Jansen, Stafforst)
- Microscopic Imaging Techniques (Wolters, Feldhaus)
- Epigenetics and gene regulation in infection biology (Filarsky)
- Systems Immunology (Wimmers)
- Anti-Cancer Therapy (Schulze-Osthoff, Eßmann, Wimmers)

term 1: 33p

Advanced
Biochemistry

Molecular Cell Biology

Current Topics

IFIB Course

Course

Tübingen
Lab

Course

Lab

term 2: 27p

Signaling

term 3: 30p

Lab

Courses from contributing investigators

Module: BCH-M-EC2 and -EC3

- Imaging from Probe Development to in vivo Application (Beziere)
- Cell Biochemistry with Fluorescent Fusion Proteins (Rothbauer)
- Immunology (Weber & colleagues)
- Advanced Infection Biology (Wagner)
- Osteogenesis and Wound Closure (Ehnert, Nüssler)
- Pathobiochemistry (Weigert)
- Structure-based Drug Design (Thiel)
- Introduction to Metabolic Imaging (Martins)
- Yeast Cell Biology (Ewald)

term 4: 30p

Master
Thesis

(Additional courses might be offered irregularly.)

Apart from the requirement of at least one IFIB *course*, you are completely free to choose according to your personal preferences and aims. You could, for example, focus on medical topics like oncology, pharmacology and pathobiochemistry. Or, you could enhance your technical skills with bioinformatics, microscopy, and biochemical techniques.

Courses are evaluated combining different aspects of the work done. Many will take into account your lab protocol, have an exam to gauge your grasp of the underlying theory, and evaluate your participation and lab work.

Timing of Courses

In designing the master degree, we had to choose between unavoidable trade-offs. On the one hand, a master with smoothly timed but compulsory elements. On the other hand, a master that gives you many choices but where timing problems may occur. We favour the latter option since it is important that you can develop your own personal profile and since we believe you can organize your own schedule.

Some courses are offered both in winter and summer term. Most of them have no additional requirements and can be completed within a four-week time window. Please check the current course description below.

Most of them also have a page with details on the learning platform *ILIAS*, which you can access as a student of our program.

IFIB Course: Cell Biochemistry of Organelles (BCH-5500)

Goals	Experience in handling of yeast cell cultures; subcellular fractionation of organelles; experience in fluorescence microscopy of mitochondria and peroxisomes; cell-free translation systems; critical presentation and discussion of data
Content	<u>Topics</u> : mitochondrial biogenesis and morphology; peroxisome dynamics and function <u>Methods</u> : subcellular fractionation, fluorescence microscopy, native gel electrophoresis, yeast genetics, in vitro translation
Format	Lecture, seminar, practical course
Frequency	Summer term
Participants	4-8
Evaluation	Audited protocol (not graded) Oral presentation of the results and oral examination (=> grade)
Organizers	Doron Rapaport

IFIB Course: Cell Signaling (BCH-5570)

Goals	Understanding of selected signal transduction modules that are important for cardiovascular function in mammals Evaluation of the relevance of biochemical research to improve our understanding of human health and disease Getting experience in handling and analysis of mammalian cells and stem cells in particular Design and performance of well-controlled experiments; documentation and interpretation of data; critical presentation and discussion of data
Content	<u>Topics</u> : signal transduction in mammalian cells, with a focus on the cardiovascular system; visualization of signaling molecules in real time in living cells; cell death and survival; transgenic mice as models for human diseases; stem cell biology; regenerative medicine <u>Methods</u> : biochemical analysis of signaling proteins; live cell imaging (e.g. with FRET-based biosensors); analysis of gene expression (e.g. immunohistochemistry on mouse tissue sections); cell growth assays; generation of genetically-modified mice; Cre/lox recombination system; culture of mammalian cells, in particular stem cell culture and differentiation; iPS cell technology
Format	Lecture/seminar, final seminar Practical course
Frequency	Summer term
Participants	4-8
Evaluation	Protocol (graded, 25%) Performance at work (graded, 25%) Oral examination (graded, 50%)
Organizers	Robert Feil, Susanne Feil, Hannes Schmidt

IFIB Course: Chemical Biology (BCH-5560)

Goals	Exercise in advanced solid-phase peptide chemistry. Handling and synthesis of building blocks for solid-phase peptide synthesis. Application of chemoselective reactions for protein engineering. Development of strategies for protein semisynthesis.
Content	<u>Topics:</u> posttranslational modifications of proteins, chemical labeling of proteins, protein chemistry <u>Methods:</u> solid-phase peptide synthesis, organic chemistry, protein semi-synthesis, methods for purification and analyses of semisynthetic proteins
Format	Lecture, seminar, practical course
Frequency	Summer term (from summer 2024 onwards)
Participants	2-8
Evaluation	Protocol (10-15 pages including introduction, results and discussion) 50% Oral examination (20 minutes) 50%
Organizers	Dirk Schwarzer

IFIB Course: Modern Genetic Engineering (BCH-1290)

Goals	Overview of the most important methods of modern genetic engineering and their application in plants
Content	<u>Methods applied:</u> PCR mutagenesis, Gateway cloning, DNA sequencing, transient protein expression, transformation and PCR analysis, synthetic gene design <u>Methods discussed:</u> generation of genetically modified organisms, virus-induced gene silencing, amiRNA technology, TALEN, lambda-red
Format	Lectures, seminar, practical course
Frequency	Winter term
Participants	0-6 (together with 10 BSc students)
Evaluation	60 % oral exam 40 % seminar talk and participation in discussion (Bonus/Malus: Labbook) Details will be announced at the beginning of the course
Organizers	Andrea Gust, Elisabeth Fuß

IFIB Course: Posttranscriptional Control of Gene Expression (BCH-5510)

Goals	Experience in handling of RNA; design and evaluation of experiments for quantitative measurement of RNA and proteins; design of control experiments for quantitative measurements; critical presentation and discussion of data
Content	<u>Topics:</u> mRNA stability; translation and its control; RNA-binding proteins; RNA modification; RNA editing <u>Methods:</u> In vitro transcription; Probe generation; Northern blot; qRT-PCR; RNA editing assays
Format	Lecture, seminar, practical course
Frequency	Winter term
Participants	4-8
Evaluation	Audited protocol and labbook (not graded but bonus possible) Oral presentation of the results (not graded but bonus possible) Oral examination (=> grade)
Organizers	Ralf-Peter Jansen, Thorsten Stafforst

IFIB Course: Epigenetics and Gene Regulation in Infection Biology (BCH-5640)

Goals	Advanced understanding of the basic principles underlying epigenetics and gene regulation in eukaryotic organisms, with an additional focus of their relevance in the context of infection biology. Gain insights in state-of-the-art methodology to investigate epigenetics and chromatin biology. Design conduct and control complex molecular and cell biology experiments. Critical presentation and discussion of experimental data.
Content	<u>Topics:</u> Apicomplexan parasites with a focus on the malaria causing parasite <i>Plasmodium falciparum</i> . Genome editing. Epigenetics and chromatin biology. Non-coding RNAs. <u>Methods:</u> Molecular cloning; CRISPR/Cas9 gene editing; Western blot; <i>P. falciparum</i> cell culture; Parasite growth assays; Flow cytometry; Fluorescence microscopy; Protein co-localization studies
Format	Lecture / Seminar / Practical course
Frequency	Summer term
Participants	4-6
Evaluation	Performance during the course (25%) Project/Seminar presentation with discussion (75%) Audited protocol (required to pass)
Organizers	Michael Filarsky

IFIB Course: Systems Immunology, Omics Analyses, and Applied Bioinformatics (BCH-1345)

Goals	Understand the principles and applications of scRNA-seq and Olink technologies in immunology. Perform basic plotting, data analysis and pipelines for scRNA-seq and Olink datasets using R. Utilize generative AI and LLMs to assist in code generation and data analysis. Independently analyze real-world datasets and present findings effectively.
Content	This intensive 4-week module introduces undergraduate students to the use of systems biology and omics tools in investigating immunological questions. Students will learn the fundamentals of analysing single-cell RNA sequencing (scRNA-seq) and Olink proteomics datasets using real-life data from previous studies. The course emphasizes computational skills, conceptual understanding, and the application of generative AI and large language models (LLMs) to enhance data analysis and code generation.
Format	This course employs modern teaching philosophies centred around active learning, student engagement, and the integration of advanced computational tools like gen AI and LLMs. Lectures incorporate interactive elements, while practical sessions encourage independent learning through online resources and AI-assisted coding. The capstone project fosters critical thinking and real-world application of skills, promoting deeper understanding and retention.
Frequency	Winter term
Participants	Up to 2 Master students
Evaluation	Graph competition (20%) Multiple-choice test (30%) Capstone project presentation (50%)
Organizers	Florian Wimmers

IFIB course: Cellular Mechanisms of Anti-Cancer Therapy (BCH-5525)

Goals	Understanding of the major signaling pathways that promote carcinogenesis and therapy resistance Understanding of the major metabolic alterations in tumor cells Obtain insights in novel tumor therapies and conduct of clinical trials Design and performance of well-controlled experiments; documentation and interpretation of data; critical presentation and discussion of data
Content	<u>Topics</u> : Principles of tumor biology; oncogenes and tumor suppressor genes; driver mutations, cancer epigenetics; control of cell death and survival; apoptosis and senescence; tumor metabolism; translational oncology and personalized medicine; molecular targeted cancer therapies, tumor immunotherapies. <u>Methods</u> : biochemical analysis of signaling pathways; live cell imaging; immunocytochemistry; flow cytometry; analysis of gene expression; assays for detection of apoptosis and senescence, cell cycle analysis; culture of mammalian cells
Format	Seminar, final seminar, practical course
Frequency	Winter term
Participants	Up to 8
Evaluation	Lab protocol Oral exam
Organizers	Prof. K. Schulze-Osthoff, PD Dr. F. Eßmann, Dr. F. Wimmers

“Cross-Campus” Course: Microscopic Imaging Techniques (BCH-5590)

NOTE: This is an intensive course with work on weekends!

Goals	Understand the technology of different microscopes. Understand the pros and cons of different imaging techniques and how to choose the appropriate ones. Ability to work with digital images and perform analysis of imaging experiments. Apply the gained knowledge by building your own optical imaging device with Fischer Technik.
Content	In the first week, basic optic principles present in all microscopes are covered in seminars, demonstrations and hands-on experiments. Furthermore, students learn how to use Fiji/ImageJ to work with and analyse digital images. From the second week on, demonstrations of different microscopic techniques (wide field, fluorescence, life cell/FRET, confocal microscopy) provide detailed theoretical and practical knowledge. If available, further techniques like 2-photon or super-resolution microscopy will be covered. Additionally, students build and characterise an optical imaging device with Fischer Technik. Some newly developed imaging techniques, that are not available on campus, are covered by seminars.
Format	Seminars, demonstration visits to groups on campus/MPI/clinics, practical course
Frequency	Summer term
Participants	4-8
Evaluation	Seminar presentation (25%), Methods Discussion (25%), Project presentation (50%)
Organizers	Christian Feldhaus; Markus Wolters

External Course: Modulating Osteogenesis and Wound Closure *in vitro* (BCH-5600)

Goals	Experience in handling of cell cultures (isolation, expansion and differentiation of primary human osteoblasts) Design and evaluation of experiments for determining the effect of biological substances/epigenetic drugs on osteoblast growth and function Team work (groups of 2); Presentation and discussion of the experimental results in English
Content	<u>Topic:</u> Evaluating the effect of various biological substances/ epigenetic drugs on osteoblast growth and function (osteogenesis) <u>Methods:</u> Isolation and culture of primary human osteoblasts, viability tests, proliferation measurement, enzyme activity measurement, histological stainings (e.g. matrix mineralization), scratch assay for measuring wound closure
Format	Lecture/seminar, practical course
Frequency	Each term
Participants	2-4
Evaluation	Written summary (2-5 pages) incl. materials, methods & results (30%) Performance at work (30% of final grade) Talk/poster presentation (40%)
Organizers	Sabrina Ehnert, Andreas Nüssler

External Course: Cell Biochemistry with Fluorescent Fusion Proteins (BCH-5580)

Goals	One of the major challenges in cell biology is to understand cellular processes in response to external stimuli. The main goal is to localize cellular components and understand their interactions. Fluorescent fusion proteins are key factors to study the cellular distribution and dynamics of proteins in living cells using time lapse analysis and High-Content Analysis based on automated microscopy. To develop cellular models for compound or siRNA screening, fluorescent fusion proteins have to be generated and expression of fluorescent fusion proteins has to be validated in a number of disease relevant cell lines.
Content	In this course students will learn different aspects on how to generate fluorescent fusion proteins in appropriate vector systems (PCR, cloning, sequence analysis) as well as how to use them in various cellular and biochemical assays (cell proliferation, immunoprecipitation, Western blotting). You will learn cultivation of various human cell lines (sterile cell culture), different techniques to transfect/transduce human cell lines with expression vectors and how to analyze transfected cells on molecular and biochemical level. Students will have access to various imaging techniques including live cell analysis and immunofluorescence.
Format	Practical course, seminar
Frequency	Each term
Participants	2-4
Evaluation	Oral presentation at the start (1/3 of grade) Audited protocol (1/3 of grade) Oral examination (1/3 of grade)
Organizers	Ulrich Rothbauer

External Course: Imaging from Probe Development to *in vivo* Application (BCH-5610)

Goals	Participants will gain a solid theoretical and practical foundation in clinically relevant imaging techniques like positron emission tomography, magnetic resonance, and computer tomography.
Content	The module deals with the radiochemistry of imaging probes (50%) and with their <i>in vivo</i> detection (50%). Participants will synthesize precursors for radiolabeling, learn how to radiolabel, and finally apply these imaging probes in small live laboratory animals. The module also involves detailed data analysis. Methods: Organic synthesis (Schlenck technique) and analysis (NMR, MS) of precursors, radiolabeling of precursors, PET, MR, SPECT, CT and optical imaging, detailed data analysis of acquired data.
Format	Lectures, practical lab work, data analysis in seminars, literature seminar
Frequency	Summer term
Participants	4-8
Evaluation	Practical evaluation, exam, literature seminar evaluation
Organizers	Nicolas Beziere

External Course: Introduction to Metabolic Imaging and Medical Diagnostics (BCH-5630)

Goals	We aim in this module to address several topics in metabolism, physiology, and medical imaging applications. The students will get acquainted with relevant metabolic pathways in cancer cell biology, develop new sensors to access reverted physiology and diagnostic medical applications. The programme will also extend to metabolic imaging <i>in vivo</i> and <i>in vivo/ex vivo</i> analysis. This module will account for the contribution of experts in metabolism, metabolomics, probe development, oncology, cancer stress, neurology, and data analysis.
Content	<u>Topics:</u> Synthesis of organic and inorganic metabolic probes, introduction to cancer metabolism, implement concepts in multimodal metabolic imaging (PET, MRI, Optical, photoacoustics) and applications Students will perform - practical study from probe development to preclinical application in cancer - tissue collection and staining - ¹H NMR metabolomics analysis - training on organizing a lab book and biostatistical imaging analysis
Format	Lectures, practical lab work, data analysis workshops
Frequency	Winter term
Participants	4-10
Evaluation	Labbook organization (report) (1/3) Oral presentation followed by a short Q&A session (2/3)
Organizers	Andre F. Martins

External Course: Immunology (BCH-5620)

Goals	Skills: practical skills (experimental planning and execution, data analysis, discussion), presentation, scientific writing Knowledge: theory of immunological methods, insights into aspects of innate and adaptive immunology
Content	Students will be acquainted with the <i>theoretical background</i> for the following immunology-related techniques in the introductory lecture block (during first week of the module and separate from the lectures "Einführungsvorlesung Immunologie" and "Advanced Immunology"): - Characteristics of the main innate and adaptive immune cells - Working with blood- and skin-derived immune cells - Isolation and phenotyping of cell types - T-cell based assays and immune monitoring - Immunohistochemistry / Immunofluorescence / Immunoblot / ELISA - Flow cytometry (intracellular and surface stain; phospho-flow cytometry) - Immunological techniques in mice / immunological mouse models In the following 3.5 weeks, 1-2 students will be assigned to the following participating labs offering <i>insights into their ongoing research</i> and allowing first-hand <i>practical experience</i>. Preferences will be considered if possible. - T-cell assays and immune monitoring (C. Gouttefangeas, Immunology) - Viral vaccine development (R. Amann, Immunology) - Peptide vaccination: from discovery to drugs (J. Walz, Immunology) - gamma-delta T cells (K. Schilbach-Stückle, University Hospital) - Immune cell signaling (S. Beer-Hammer, Toxicology) - Immune responses in the skin towards infection (B. Schitteck, Dermatology) - Innate Immunity (A. Weber, Immunology) - Neonatal Immunology (C. Gille, Neonatology) Upon completion of the practical period, students will be requested to: - present their lab project in a short Powerpoint presentation held during a 1-day colloquium in the final week (all students and PIs participate) - generate a report detailing their research topic, data and discussion.
Format	Lectures, practical, colloquium
Frequency	Winter term
Participants	4-10
Evaluation	Students present their lab project in a short Powerpoint presentation held during a 1-day colloquium in the final week (all students and lab supervisors participate). Each short presentation is followed by a short Q&A session to yield a combined final grade.
Organizers	Alexander Weber
Requirements	Attendance of the Immunology lectures at the Department is recommended prior to completion of the course but not formally required. Students with prior knowledge or experience in immunology gained in Tübingen or elsewhere may be given preference should the course be oversubscribed.

External Course: Pathobiochemistry (S01PLAB01)

Goals	Experience in detection of phosphorylated proteins, design and evaluation of experiments for detection of activated signal transduction, understanding and interpretation of pathobiochemical alterations and disease-related biomarkers in body fluids, critical presentation and discussion of data, presentation and discussion of scientific publications
Content	<u>Topics:</u> molecular mechanisms of metabolic regulation in health and disease, insulin resistance and diabetic late complications, diagnostic tools for detection of pathobiochemical alterations and disease-related biomarkers in body fluids <u>Methods:</u> cell culture, transfection of cells, immunodetection of phosphorylated proteins, immunoprecipitation, point of care testing
Format	lectures, seminar, practical course
Frequency	Winter term
Participants	4-9
Evaluation	Protocol (not graded) Presentation of a recent scientific paper (graded, 30%) Oral examination (graded, 70%)
Organizers	Cora Weigert, Rainer Lehmann, Andreas Peter
Note	This course consists of three parts: S01PLAB01 (practical), S01SLAB01 (seminar) and S01VLAB01 (lecture). Only the practical part will show on your transcript and in your examination booking, but you need to attend all three parts.

External Course: Mechanisms of Microbial Pathogenicity (Bio-MIB-200, together with Bio-MIB-208)

Goals	Knowledge of molecular and cellular mechanisms of microbial pathogenicity
Content	Theory taught in lectures - How do bacteria survive in humans and cause infections? - How does our defense work and how is it circumvented by bacteria? - Which are the most topical and most urgent infectious diseases? Practical course - <i>Salmonella</i> mutagenesis - Functional analysis of the type 3 secretion system - Cell culture infection models of <i>Salmonella</i> - Blue native gel electrophoresis
Format	Practical (2-week block S4-slot) Lectures (every Friday 8-10 AM) Seminar (2nd half of summer term, Tuesday 17:00)
Frequency	Summer term
Participants	Up to 3
Evaluation	1/3 each: exam after the lecture, seminar grade, presentation after practical
Organizers	Samuel Wagner
Note	For the lecture, see also Bio-MIB-208

External course: Search and Find: Exploring Phage-Host Interactions through Transcriptomics (BCH-5650)

Goals	Students will receive a compact but application-oriented introduction to bioinformatics. They will learn how biological research questions can be addressed using existing omics datasets. By the end of the module, students should be able to perform simple bioinformatic analyses independently, interpret their results critically, and document their workflow reproducibly.
Content	The module is focussed on transcriptomics in phage-host interactions, and students will work in teams of two on a dedicated research question, for instance: • How are defense and anti-defense systems expressed during phage infections? • What are the best approaches to classify phage genes and what can be learned from these phage gene classes? • Are there indications of previously unannotated transcripts or transcription start sites in the RNA-seq data? • Are there RNA-editing events(A-to-I) in phage infections? These questions will provide students with an impression how bioinformatics can serve as an entry point to identify targets for detailed mechanistic investigation with biochemical approaches.
Format	Block-Course of lectures and bioinformatics exercises/projects
Frequency	Winter term
Participants	6-8
Evaluation	The final grade will be built on: • the small lecture given by the student in 1st week (20 %) • two oral code reviews by the student (20 %) • the GitHub repository handed in by the student team (20 %) • final oral examination about course content (40 %)
Organizers	Dr. Maik Wolfram-Schauerte
Note	Students must bring a laptop or similar device for minimal local computations and logging and working on servers.

External course: Structure-based Drug Design (BIOINF4371)

Goals	Students have a working knowledge on the pharmaceutical development process. They are familiar with protein and ligand structures, how to resolve and model them. They can identify relevant physicochemical interactions. Students have detailed knowledge of algorithmic techniques to predict protein-ligand binding and can implement them.
Content	Starting with a broad introduction of the pharmaceutical drug development process, the lecture conveys key concepts of structure-based computer-aided drug design. Required basics on pharmaceutical key concepts are discussed, followed by basic concepts for modelling of 3D structures. In the second part, physicochemical interactions between proteins and ligands are presented, forming the basis to discuss strategies to predict protein-ligand binding with a focus on algorithms for protein-ligand docking. The estimation of binding affinities between proteins and ligands <i>in silico</i> is introduced, leading to the discussion, development, and use of scoring functions.
Format	Lecture (2h/week), exercise (2h/week) and project
Frequency	Winter term, every second year (alternating with a Chemical informatics course)
Participants	Up to 5
Evaluation	Oral exam or, in case of too many students, written exam. 50% of the achievable points from the assignments and the project, individually, are required for exam admission.
Organizers	Philipp Thiel
Note	No formal requirements. Basic knowledge of protein structure, organic chemistry, and programming skills in Python are recommended. Lectures and exercises span the whole term. In winter 2026, lectures are scheduled for Monday 16-18. Take note, that this conflicts with the MoKo's. The date for the exercises will be decided with the participants during the first lecture.

External course: Transcriptional Drivers of Cancer Cell Plasticity and Identity (S07PMOLMED21)

Goals	Students: • understand the fundamental principles of epigenetic regulation and transcriptional control of cell identity and plasticity. • gain insights into the molecular biological basis of tumour heterogeneity. • become familiar with common gene-editing technologies and understand how degron-based protein degradation systems work. • acquire practical skills in cell culture, functional assays, protein and nucleic acid analysis, and chromatin immunoprecipitation. • are able to evaluate and critically reflect on experimental data, and to analyse and interpret bioinformatic data such as mRNA-Seq, ChIP-Seq and scRNA-Seq.
Content	<u>Lecture:</u> 1. Epigenetic and transcriptional regulatory mechanisms of cell identity 2. Tumour heterogeneity in solid tumours 3. Introduction to next-generation sequencing (NGS) and library preparation 4. Gene editing: CRISPR/Cas9 and the FKBPv-dTAG-Degron system 5. Molecular basis of cell cycle regulation and apoptosis 6. Fundamentals of sequence data analysis: pre-processing, alignment and visualisation. <u>Seminar:</u> 1. Analysis and interpretation of qRT-PCR results 2. Chromatin immunoprecipitation (ChIP): data analysis and methodological critique 3. Analysis of proliferation assays using live-cell imaging 4. Analysis of apoptosis assays: fluorescence-based measurements 5. Introduction to the analysis of ChIP and mRNA-Seq data 6. Single-cell RNA-Seq: data analysis and biological interpretation <u>Practical:</u> 1. Cell culture techniques: cell seeding, medium changes, treatment 2. Protein extraction, BCA assay, SDS-PAGE, Western blot 3. RNA isolation, cDNA synthesis, quantitative PCR (qPCR) 4. Chromatin isolation, shearing, DNA purification, ChIP-qPCR 5. Real-time monitoring of cell proliferation and apoptosis using imaging systems
Format	Lecture, seminar, practical course
Frequency	Winter term
Participants	Up to 2
Evaluation	Final report
Organizers	Prof. Dr. Steven A. Johnsen
Notes	Location of the course is in STUTTGART: Robert Bosch Centrum für Tumorerkrankungen (RBCT) Auerbachstraße 112, 70376 Stuttgart Students must bring a lab coat, notebook to take notes and an external harddrive or USB drive. In Winter 2026 this course will only grant you 4 ECTS.

Labs (Modules: BCH-M-LABTUE, BCH-M-LAB2, BCH-M-LAB3)

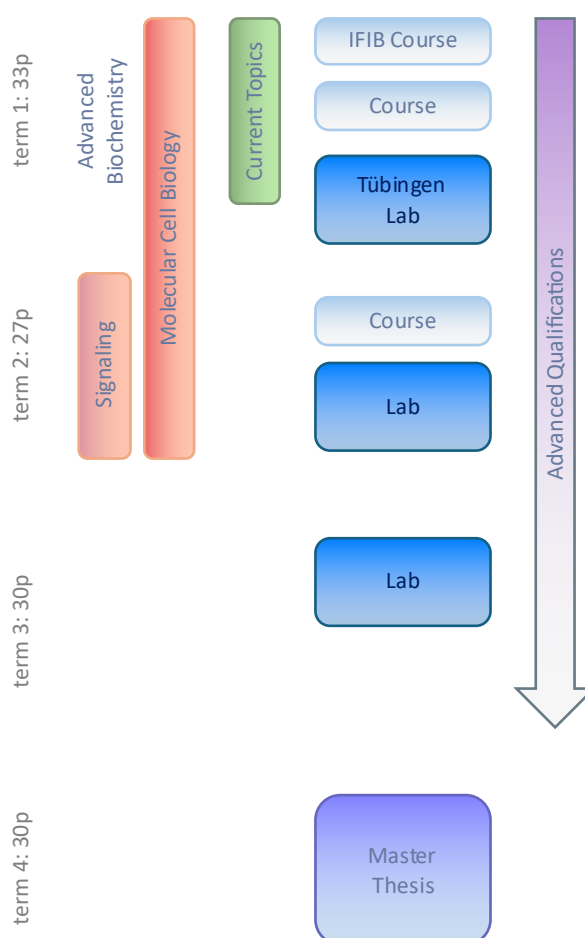
Labs (or lab rotations) are short research projects in biochemistry. They consist of about two months of lab work followed mostly by writing up a summary/protocol and a final presentation. You will arrange your own *Labs* by contacting selected principal investigators. All labs must be formally registered.

We are thinking of *Labs* as a way for you to test yourself in different research areas of biochemistry. This will help you determine which topic is most suited to your preferences and skills. After having explored different fields of biochemistry you will be in a better position to decide where to do your Master thesis. For the same reason you should not do more than two *Labs* with the same group.

Labs often cannot be graded reliably because the results depend not only on your skills and work ethics but also on external factors. Also, *Labs* done abroad may shorten your effective working time at the bench and thus decrease your data output. Hence, we have decided that these elements of the *Master of Advanced Biochemistry* would not be graded. We believe this will give you more flexibility to test various topics of modern biochemical research.

In total, you will select 3 *Labs*, one of which has to be in an academic lab in Tübingen. With 3x 15 ECTSs, or 45 ECTSs in total, the *Labs* will be the largest part of your master. You can do one or more labs in other German laboratories or even abroad. If you want to spend more time on a topic, you can fuse a lab together with the *Advanced Qualifications* module into a continuous lab block for a total of about four months.

When looking for a lab, keep in mind that at your stage you require good supervision. This can be harder to find in large, occasionally more impersonal groups. We are collecting your feedback on labs to build a database to help you profit from previous students' experience when looking for labs. This can be especially important for industry labs who are often inaccessible without personal contacts.



Goals	You review published data in preparation for experimental work. You practice the flow from hypothesis, via experimental design including controls, to analysis of the results and interpretation. You accurately document your work and present your data and ideas.
Duration	8 weeks
Evaluation	Audited protocol and oral presentation of the results and/or the theoretical background (all not graded)
Responsible	MSc Biochemistry coordinator
Organizers	IFIB group leaders, other group leaders after confirmation
Credit points	15 CP

Advanced Qualifications (Module: BCH-M-AQ)

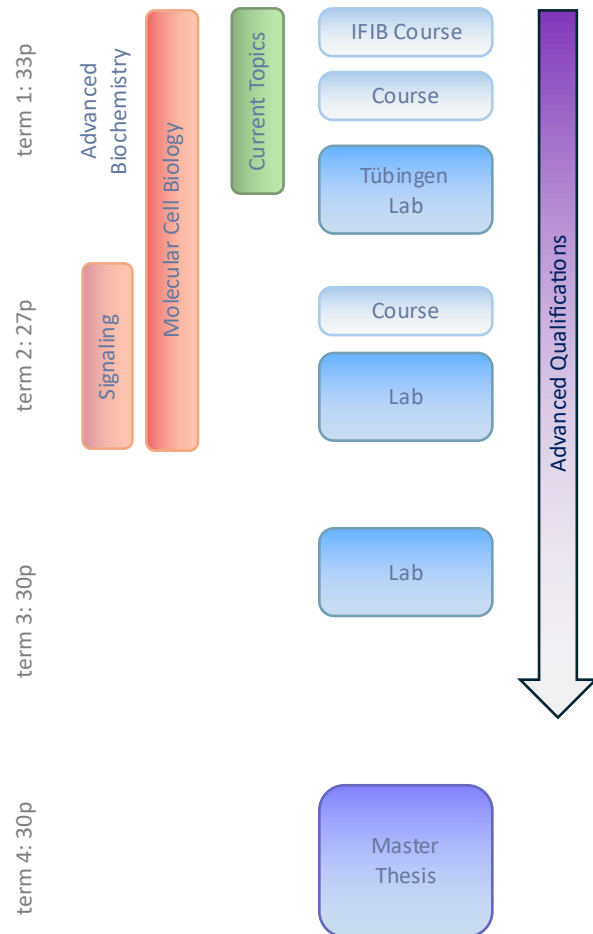
The *Advanced Qualifications*-module gives you even more flexibility to adapt your studies to your interests or future ambitions.

Please note, that this is not a Soft-Skills-module, like you might have had in your Bachelor's degree.

As mentioned before, you can use this module to do a double lab. Or you can use it for a fourth single lab rotation.

However, you are not restricted to filling this module with lab rotations. Of the 15 ECTS in this module, at least 10 must be earned from courses or lab work in the field of natural sciences or medicine. A maximum of 5 ECTS can be gained from other academic fields, like language or business classes.

If you are interested in an international study exchange, you can use this module for courses you took abroad.



Master Thesis (Module: BCH-M-MS)

Your *Master Thesis* is the last and final module and concludes the two-year degree program with us. To start your thesis, you must have finished all previous modules.

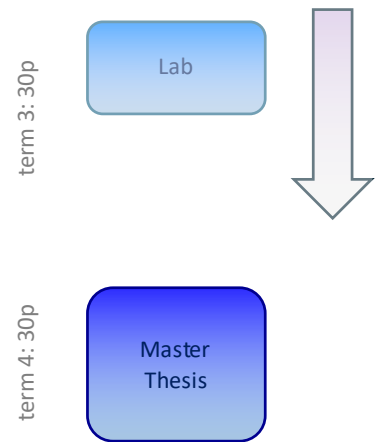
After having brushed up on the theoretical foundations in the *ABC lecture* and *Current Topic* series, after having done more organized practical training in several *Courses* and then in more self-organized laboratory placements, the *Master Thesis* will be your most independent and in-depth piece of research to date.

This is reflected in its setup. The longer time allotted gives you more time to dig deeper into the topic of your choice. More than in the *Labs* before, we expect you to actively develop your research plan before conducting experiments. The write up will also be more extensive and evaluated more thoroughly.

After doing several *Labs*, you will have gained a good overview of several biochemical research areas. Carefully choose which thesis topic is most suited, as it will influence future choices regarding a doctorate or your job search. The topic of the *Master Thesis* has to be clearly separate from any previous projects in the same lab, if any.

As with the *Labs*, the *Master Thesis* can also be done abroad if you wish. As it needs to be formally graded and influences your final master grade, please consult us during your decision process. You will also need a second supervisor in the IFIB.

We hope that the *Master Thesis* will bring you a step closer to becoming an independent and fully fledged biochemical researcher.



Goals	You survey published data to develop a novel hypothesis or to address unanswered questions You devise an experimental plan incl. suitable methods, controls, and timing You write up your work accurately and present your new findings.
Format	Practical work
Duration	6 months
When?	4 th semester
Prerequisite	All other modules (ABC exam, Current Topics, Courses, Lab Rotations, Advanced Qualifications)
Evaluation	Oral presentation of results (not graded) Written thesis (⇒ graded by the direct supervisor and, if necessary, a 2 nd reviewer)
Responsible	Head of the examination committee
Organizers	IFIB group leaders, course organizers, other labs after confirmation
Credit points	30 CP

Final Grade

As you progress from undergraduate studies via the master with us to a doctorate or employment outside academia, grades become less important and experimental skills, experience, contacts, and publications gain in weight. Nevertheless, the final grade of the *Masters in Advanced Biochemistry* will be of some influence.

The final grade is composed in equal part of the grades of the *Advanced Biochemistry* exam, your three *Courses*, and the *Master Thesis*. If you completed four courses, the best grades will be used automatically. It will thus be based in equal parts on the oral examination after the lectures, the mean grade of your *courses*, and the overall grade of your master thesis. On average you get one to two grades per term.

