

The Biobank of the Medical Faculty Würzburg

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It is our great pleasure to provide you with this brochure introducing a key instrument for the progress of medical research at our faculty: The Interdisciplinary Bank of Biomaterials and Data Würzburg - the »ibdw«.

Initialliy the **ibdw** was one amongst the first five centralized German biobanks funded by the Federal Ministry of Education and Research (BMBF) in the framework of the National Biobank Initiative. Today, together with 11 other German biobanks, the ibdw represents one key element of the German Biobank Alliance (GBA), a framework funded by the BMBF until end 2020.

The **ibdw** is an independent interdisciplinary institution of the medical faculty under joint responsibility of the University Hospital and the University of Würzburg.

With the implementation of the **ibdw** it is now possible to systematically collect, portion, and long-term store liquid biosamples (like blood, serum, urine) as well as tissue samples that have been donated by our patients and study participants fulfilling highest quality standards.

Medical research with biological material linked to related clinical data will enrich current medical knowledge. It will foster improvements in the early recognition, diagnosis, treatment, and prevention of known and hitherto unknown diseases.

We invite you to get some deeper insight into the **ibdw** – its objectives and principles, technical capacities, achieved targets and, of course, the team representing the core of the ibdw.

Enjoy!

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Welcome to the **ibdw**



The **ibdw** is the central bank of biological materials and data of the Medical Faculty of Würzburg. It is a joint institution under the responsibility of both the University Hospital and the University of Würzburg. The **ibdw** is governed by its own steering committee and is not associated with a specific department.

The **ibdw** stores liquid biosamples (e.g., blood, serum, blood cells, urine, and cerebrospinal fluid) and solid biosamples (e.g., tissue specimen, biopsies). These samples are long-term stored maintaining highest quality standards for future medical research.

All biological materials hosted by the ibdw are donated on a strictly voluntary basis by patients and study participants for (bio-)medical research.

To protect a donor's privacy all identifying information is replaced by unique number-/letter codes (»double pseudonymized«).



The Central Biobank Würzburg

Support of Future Research

The **ibdw** is a valuable resource for the progress in medical research

Research based on human biological material and related analytical and clinical data that have been or will be obtained are crucial to foster progress in medical research.

The **ibdw** stores human tissue and liquid biosamples over many decades maintaining highest quality standards and safety conditions.

By provisioning high-quality biosamples the **ibdw** contributes to evidence-based modern medicine.
Thus, in the future, everyone will possibly benefit.

On the following pages we would like to introduce you to the **ibdw** of the Medical Faculty Würzburg and give you some insight into the various working fields of a Biobank.

Donated human biological material is labeled, registered, portioned in small 750 µl tubes (»aliquots«), and stored in fully automated cryo-repositories.

Which data is stored together with the samples?

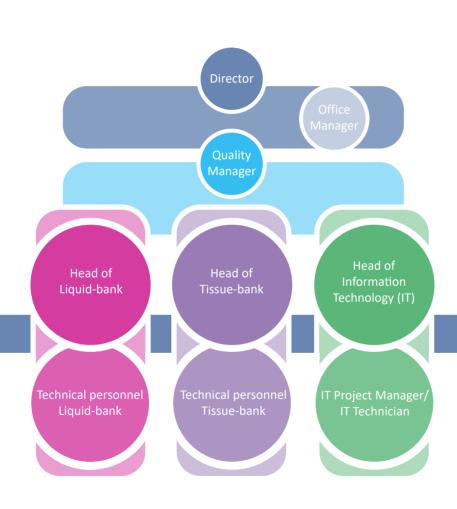
- General personal data (e.g. age, sex)
- Medical data (e.g. main diagnosis)
- Analytical data derived from biological material

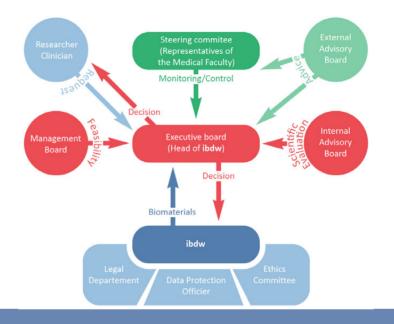


- the central liquid-bank
- the central tissue-bank
- the central database

Organisational Structure

The **ibdw** is composed of a core database and two centralized biosample repositories – one for liquid biosamples and the other for solid/tissue biospecimen – and a limited number of specialized decentralized subunits, each of them adhering to **ibdw** standards.





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National and international scientists and research groups as well as collaborating industry partners may request data and human biological materials hosted by the **ibdw** for biomedical research projects. However, any application requires prior ap-

proval by an independent ethics committee.

External advice and internal monitoring ensure highest quality standards of the **ibdw**

Quality Assurance

Tibdw with its quality management system accomplishes all requirements of the standard DIN EN ISO 9001:2015.

The certification procedure regularly analyses the implementation of a quality policy, the comprehensive documentation of all **ibdw's** procedures as well as its processes and testing measures.

8 | Workflow – Tissue Samples

ZOM A: Level -1 ZOM A1 Level -3 Sketch of Department of Surgery







Workflow – Tissue Samples

Human tissue taken during routine surgery is transferred to the rapid section laboratory where tissue samples are cut and temporarily stored in liquid nitrogen.

Human tissue taken during routine surgery that is not required for further diagnostic evaluation may be donated to the ibdw's tissue bank for biomedical research purposes.

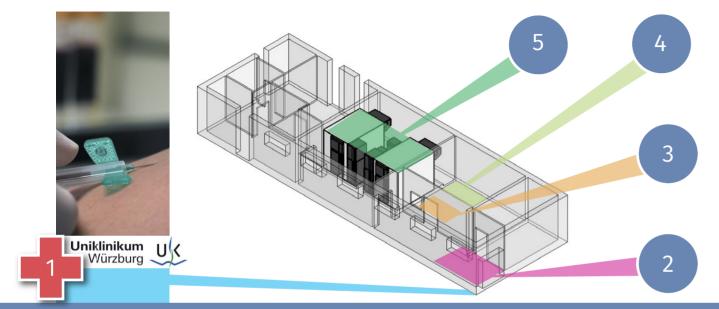
Time is an important factor when collecting tissue samples. It takes only a few minutes from the extraction of the material by the surgeon, the pathologist's evaluation in the rapid section laboratory, and the final storage of the specimen in our central tissue repository. Timestamps at each step of this process serve to monitor and document sample transport and -quality. Human tissue samples are analyzed in the rapid section laboratory. Residual tissue is prepared and transferred to the **ibdw** tissue repository in individually labeled tubes.

Tissue tubes are registered in the central tissue repository. A unique 2D-barcode engraved in the tube-bottom allows identification, allocation, and retrieval of each tissue sample.

Subsequently the tissue tubes are stored in online monitored freezers at -80 °C. The location of each single tissue tube is stored in the ibdw database.

HE-stained reference sections of each tissue sample hosted by the **ibdw** provide information about the type and quality of each individual tissue sample.

10 | Workflow – Liquid Biosamples



Workflow – Liquid Biosamples

Liquid biosamples are collected during routine blood drawing in wards or outpatient departments of the University Hospital. Sketch of the **ibdw**-building illustrating the interior layout of the liquid repository.

The individual steps of biosample processing are indicated by numbers and different colors.

The workflow from storage to release of individual biosamples is quite complex.

Each step of this process is managed and controlled by competent staff together with a high degree of automation that comprises labelling, tracking, storage and retrieval of biosamples.



The registration process of biosamples includes verification of the temperature-track, transport time, and sample integrity.



Some liquid biosamples are centrifuged prior to processing. This procedure separates liquid biosamples into individual components.



Material from routine blood vials is automatically transferred into $750\,\mu$ l tubes by a robot. 96 of such tubes fit on a single carrier rack.



Carrier racks with 96 tubes are transferred into our fully automated cryorepositories at constant -80 °C temperature.



The design of our cryo-repositories follows the principle of a chest freezer. The temperature is maintained at constant -80 °C.

Sensitive moving parts are localized in a special chamber at -20 °C. Here, the $750\,\mu$ l tubes can be reorganized/picked and/or compiled and released for analytical testing. During these processes the system keeps humidity below 25 % thus preventing icing of the tubes .

The name »Kiwi Tube Store« for the cryo-repositories is derived from the operation method of the robot handling individual tubes.



Similar to the running bird from New Zealand that "pecks" its prey with his long beak, the robot's gripper arm selects individual 750 µl tubes for reorganization or release.



Automation

A high degree of automation ensures a reliable high quality of the biosamples

Routine vials are divided into multiple 350 µlportions (»aliquots«). Thus, each aliquot contains
sufficient material for most analytical tests.
The advantage: Utilizing only one aliquot for
analytical testing leaves the other aliquots of the
same sample untouched. This strategy guarantees
constant high quality of the biosamples.

The fully automated pipetting-device processes (»aliquots«) up to eight routine vials simultaneously.

Each »aliquot« has a unique identifier represented by a scanner readable two-dimensional barcode engraved in the bottom of each 750 µl tube. A single cryo-repository contains 196 movable shelves that are divided into 28 slots each. Each section provides storage space for a carrier rack holding up to 96 sample tubes (»aliquots«).

Thus, one cryo-repository may contain up to 526.848 sample tubes.

As a backup in case of technical failure each cryo-

repository has two independent cooling units. If the main unit breaks down, the backup unit automatically takes over. If both cooling units fail simultaneously, an automated emergency system based on liquid nitrogen will be activated immediately to ensure low storage-temperatures

Freezing Concepts

»Kiwi«: Like the running bird from New Zealand the robot »pecks« single tubes

14 | Ultra-low Temperature Storage (-180°C)



Biosamples for sensitive analytical methods are stored at ultra-low temperatures at -180 °C

Based on fee-for-service a limited number of biosamples for sensitive analytical methods (e.g. proteomics, metabolomics) can be stored at ultra-low temperatures (-180 °C).

This low temperature is achieved by filling the storage containers with liquid nitrogen. The samples are not in direct contact with the liquid nitrogen itself but stored in the vapor that hovers above.



Ultra-low Temperature Storage (-180°C)



As described before single carrier racks, each of them holding 96 tubes are registered in the **ibdw** database and placed manually in each slot of a storage tower. Each tank contains 32 storage towers resulting in a total of 34.000 tubes per tank. Due to the ultra-low temperatures transfer and release of single tubes have to be carried out very carefully. For security reasons our staff is equipped with special protective garment, visor masks and gloves.

Sophisticated technology is the basis for maintaining a constant and high quality of the stored biosamples. The engine-room of the **ibdw** building contains almost all high-tech equipment required for heating, ventilation, air conditioning, water supply, and air compression.

All technical systems are redundant by design ensuring constant functioning in case of power breakdown or failure of air conditioning.

The backbone of all cross-linked technical systems is formed by a total of about 12.5 km of power-and network-cabling.

The capacity of the **ibdw's** ventilation system would be able to inflate 420,000 footballs within one hour.

An emergency cooling system with 3,500 liters of liquid nitrogen safeguards our stored biosamples in case of any severe cooling failure.

The Engine-Room of the **ibdw**

ibdw's air ventilation would be able to inflate 420,000 footballs within one hour

»Broad Consent« – The consent of the **ibdw**

The hospital's **broad consent** procedure facilitates the collection of human biosamples long-term stored for (bio-)medical research not bound to specific research purposes.

Human biosamples collected under the Broad Consent are also called faculty biosamples.

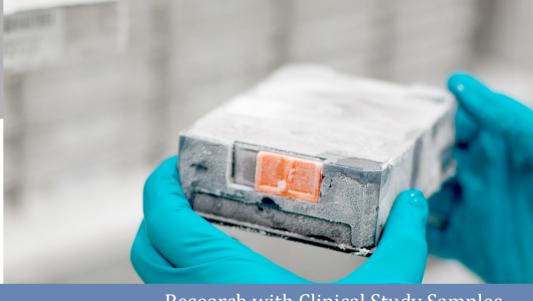


Your precious clinical **ibdw's** cryorepositories

Unlike broad consent samples, samples from clinical studies are based on a project-specific informed consent.

The number of biosamples and storage-duration are given by the study protocol. The study-PIs are owner of the respective biosamples.

For biosamples collected in the frame of a clinical study specificially tailored labels are generated in accordance with the study-PIs.



Research with Clinical Study Samples

Research with »Broad Consent«-Biosamples

These faculty biosamples are not bound to specific research purposes. They can be used for (bio-)medical research worldwide under clearly specified conditions

The University Hospital of Würzburg is the owner of these biosamples.

A standard sample collected under Broad Consent conditions includes:

- 2 Serum-tubes à 4.7ml
- 1 EDTA tube à 2.8 ml
- 1 Tube for Spontaneous urine

Labels for faculty biosamples are generated by the hospital's laboratory order entry system (Lauris)

available throughout all departments of the University Hospital.

Together with its research partners, the ibdw has developed a simplified procedure for researchers to obtain easy access to broad consent biosamples. Further information on the easy access procedure can be found on page 18 of this brochure.

Reciept, storage and release of study samples are associated with costs.

We will gladly support you in the planning of your biosample collection and best storage conditions for the success of your research project.

Please contact our office: ibdw@ukw.de

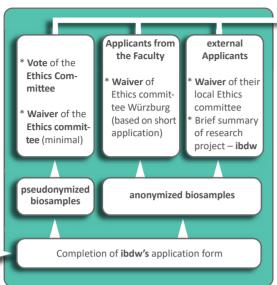


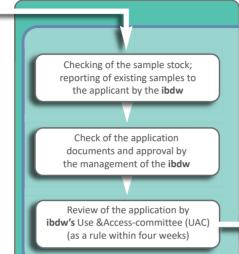


Sample collection of the medical faculty

Sample pool of the faculty, from which I need certain biosamples for my research project

Send a **stock inquiry** about available biosamples via e-mail to the ibdw office (ibdw@ukw.de)





PROJECT-/STUDY-**SAMPLES** Accounting ESPONSIBILITY OF THE IBDW I have collected biosamples Shipping of samples by the **ibdw** for my project/study and or pick up by the applicant these are hosted by the ibdw Inquiry about required samples ibdw will provide the samples to ibdw's office (ibdw@ukw.de) within a maximum of 14 days

Human samples of a clinical study are hosted by the ibdw and are ordered on demand.

Study samples are delivered by the **ibdw** within 14 days after the order.

Samples are shipped by the **ibdw** but can also be picked up by the study/project staff

Subsequently, the ibdw sends an invoice for the delivered samples and, if necessary, for shipping

Request and Delivery of Biosamples

from the **ibdw**.

The **ibdw** has designed a straight forward process that enables researchers to obtain easy access to biosamples.

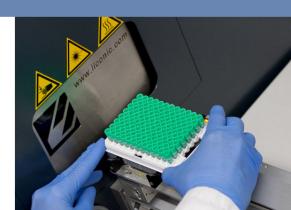
In the case of research taking advantage of broad consent biosamples, the ibdw office can check whether the required samples are available.

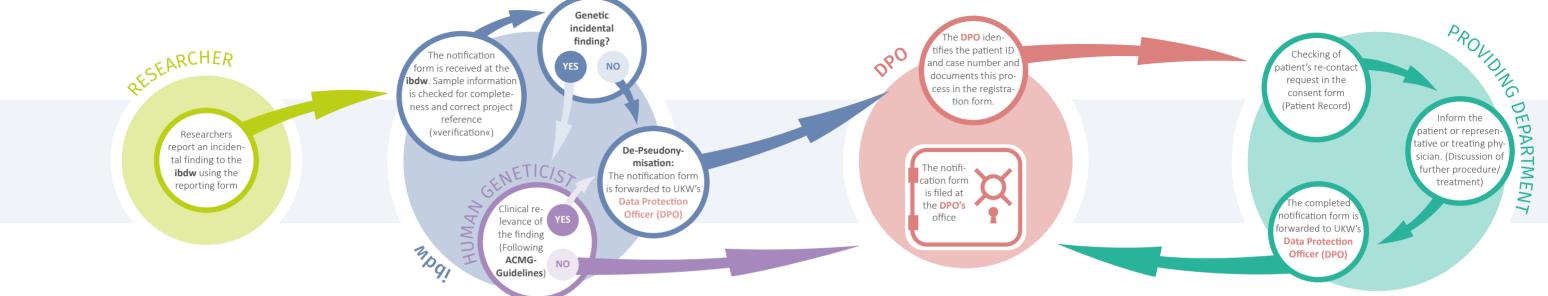
Thereafter researchers enter the required details in the sample application form provided by ibdw. When applying for pseudonymized biosamples, an ethics vote and a study protocol must be submitted. When applying for anonymized biosamples, a waiver of the responsible by the ethics committee and a brief description of the project are required.

Subsequently, the ibdw management board checks the application documents and the availabilty of the requested biosamples. If the request complies with the specifications and there are no objections from the ethics committee, the ibdw management acknowledges correctness and feasibility of the application.

Within the following 7-22 working days the application is reviewed, a decision is made by the ibdw's Use & Access committee.

After UAC approval, the ibdw delivers the reguested biosamples within a maximum of 14 days.





Management of Incidental Findings

Incidental findings can be a challenge for researchers

Incidental findings are often regarded as a challenge for the researcher(s). If additional or random incidental findings are made during a research project with »Broad Consent« samples from the **ibdw**, it is possible to contact the respective donor mediated by the ibdw. the custodian (i.e. the institutional DPO). and the clinic involved.

The management board of the ibdw together with the ethics committee, the legal department and the institutional data protection officer (DPO) of the university hospital developed a workflow (including documentation-templates) with a sequential notification process shown in the scheme above.

After verification if the respective biosample(s) do in fact originate from the **ibdw**, in case of non-genetic incidental findings for de-identification of the donor, the master-workflow involves a trusted third party - in our case the institutional DPO.

The DPO informs the head of the respective department where the patient donated her/his biosamples.

The providing department (1) will check the medical records for the broad consent form (consent to re-contact or right not to know), (2) discuss the potential clinical relevance of the finding with specialists, and (3) finally organize an appointment with the patient/donor. During this appointment in hospital the patient/donor will be informed and will recieve medical advice.

If you have any further questions regarding the topic of incidental findings in human samples please contact our office: ibdw@ukw.de

ibdw offers additional comprehensive services

The key task of the **ibdw** is to collect, store and retrieve human biosamples based on a broad consent scheme. The hosted biosamples comprise all kinds of body fluids as well as tissue specimen.

In addition the **ibdw** offers project-specific processing, storage and retrieval of human biosamples as a service for the Medical Faculty.

Storage

- Sample registration
- Sample processing and preperation (if necessary)
- Semi-automatied storage of »fresh frozen« tissue specimen and biopsies
- Automated storage of liquid biosamples at -80 °C
- Semi-automatic storage of liquid biosamples at -180 °C

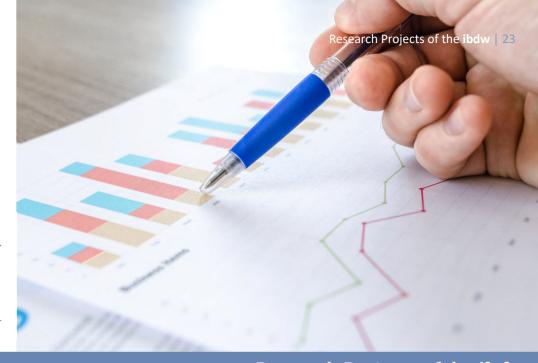
Training and Logistics

- Training of sample providers (medical and nursing staff, study staff)
- Setup of specific primary sample labels in the Lauris system of the UKW
- Planning of biosample logistics including sample tracking and quality controlled storage, documentation and reporting for studies/projects
- Quality controlled isolation of DNA/RNA from human liquid samples and tissues



As a key member of the German Biobank Alliance (GBA), the **ibdw** regularly participates in ring-trials.

By these trials techniques and measurement procedures in the field of tissue and liquid sample biobanking are assessed with regard to quality and reproducibility. These trials serve as a benchmark in order to optimize and harmonize biobank-processes



Services of the **ibdw**

All **ibdw**-processes are controlled and monitored according to DIN EN ISO 9001:2015.

Backup and emergency concepts also secure an optimal environment for biosamples and data.

Delivery

- Delivery of human biological materials, including the preparation and special stainings of sections of tissue samples, if this is is applied for (Standard: HE staining for reference sections)
- Delivery of quality-controlled purified DNA/RNA in single tubes or on 96-Well carriers (rack)



One kind of ring trial comprises the processing the processing of a same source material by different biobanks. The results are sent to a reference laboratory where the data are analysed and compared.

Stakeholder engagement is a critical factor for the success of a biobank. Biobanks must take into account the views of various interest groups and integrate these different view-points into their daily work.

Stakeholders of the **ibdw** include representatives from medicine, research and science, the press, funding organisations, the biobank community, industry and, in particular, donors, namely patients and test persons.

As a corner stone of the GBA, **ibdw** significantly contributes to the development of strategies and concepts to integrate the different stakeholder view-points in modern biobanking.

Research Projects of the **ibdw**

In addition to its range of services, **ibdw** carries out research in its own projects



May 2011

Founding of the **ibdw** in the frame of the BMBF-funded National Biobank Initiative

September 2011

National pilot-approach: implementation of a broad consent for donation, long-term storage and thematically unrestricted use of human biological material for (bio-)medical research purposes

December 2012

Enacting of the statutes of the **ibdw**

June 2013

Grand opening & first public »Open Day« of the ibdw

January 2014

Quality-controlled storage of the first tissue samples

Cooperation contract with the Blood Donor Service of the Bavarian Red Cross (»BioKEP«)

December 2014

Quality-controlled storage of the first liquid biosamples in the automated cryorepositories

August 2016

ISO certification of the ibdw according to DIN EN ISO 9001:2015 (Certificate No: Z13305)

History and Development of the **ibdw**



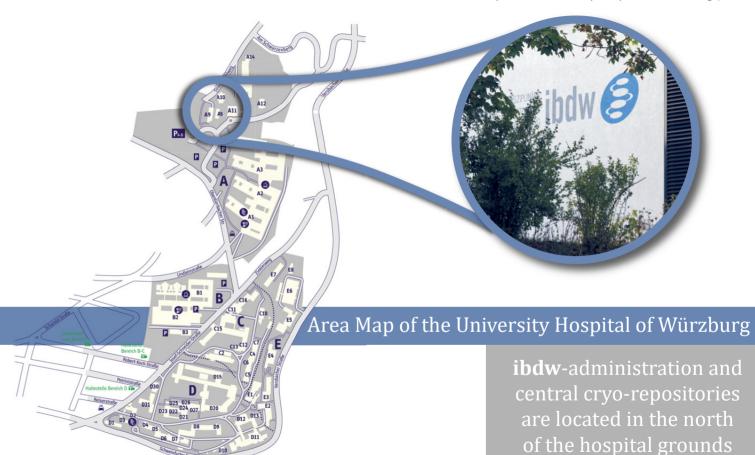
ibdw becomes member of the BMBF-funded German Biobank Node »GBN«

May 2019

ibdw becomes member of the BMBF-funded German Biobank Alliance »GBA«

August 2019

Re-certification of the ibdw according to DIN EN ISO 9001:2015 (Certificate No: Z13305R1)





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Alliance Partner Biobank Node bbmri.de











