



BIOBANKING IN TIMES OF COVID19 – PRE-ANALYTICAL PROCESSES

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BBMRI-ERIC QUALITY MANAGEMENT SERVICES FOR BASIC AND APPLIED RESEARCH



KNOWLEDGE HUB

Find everything you need to know about quality management in biobanking and biomedical research in one place: The BBMRI.QM Knowledge Hub consists of a pool of experts that can help you with any of the following:

- International standards relevant for biobanking and biomedical research
- Quality management tasks in EU projects
- Quality management contribution to national and international research project proposals
- Specific questions you may have
- Biobanking in times of COVID-19



TRAINING & SUPPORT

Whether you are looking for training opportunities or practical support with your quality management – quality experts can provide you with a solution tailored to your needs.

Within the BBMRI.QM network, we offer inhouse and online training, university courses, summer schools, short courses, workshops, and consulting services covering different quality management systems:

- Web conference series about the biobanking standard ISO 20387:2018
- International standards relevant for biomedical research
- General quality management systems
- Integrated management systems



AUDITING

Request an audit from BBMRI-ERIC:

If you run a non-certified biobank and/or you want to know if the samples stored fulfill certain quality requirements, get the support IBBMRI.QM. We offer peer-review-style audits on request. Take the next QM improvement step together with us!

 BBMRI-ERIC Self-Assessment Survey for biobanks

What is the BBMRI-ERIC Self-Assessment Survey? How does it work? More details...

BBMRI-ERIC Audit





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BBMRI.QM WEB CONFERENCE

Web conference #2 will take place on 7 April

BIOBANKING IN TIMES OF COVID-19, PRE-ANALYTICAL PROCEDURES

Aim

This web conference is aimed at biobanking experts and researchers. We will follow up on the open questions raised in the first web conference and, in addition, we will focus on pre-analytical procedures in the hospital and biobanking setting.

"Routine processing of COVID-19 samples from the bedside to the laboratory and to the biobank" will be addressed, and we will discuss whether researchers have any "additional requirements for the sample quality", when thinking of any scientifically driven downstream analysis.

Together with Lukasz Kozera, colleagues and partners from the biobanking community, I will address your questions and together we will try to give answers while incorporating your expertise.

The web conference will take place on 7 April 2020, 10:00-12:00 am CEST.

Join us!

Web conference #1 took place on 1 April 2020:

BIOBANKING IN TIMES OF COVID-19, RISKS AND OPPORTUNITIES FOR BIOBANKS

Aim and Agenda

Follow this link

Recording

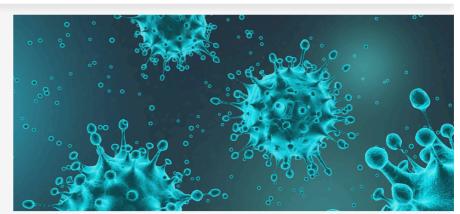
Follow this link

Presentations

Petr Holub - BBMRI-ERIC Directory and Negotiator for COVID

Maurice Bradsley - EVAg BBMRI COVID19

Andrea Ganna – COVID-19 Rapid Global Reset Response Genetics





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Documents

WHO

- Laboratory biosafety guidance related to coronavirus disease (COVID-19): interim guidance, 19 March 2020
- Laboratory testing for coronavirus disease (COVID-19) in suspected human cases: interim guidance, 19 March 2020
- Laboratory testing of human suspected cases of novel coronavirus (nCoV) infection: interim guidance, 10 January 2020
- Guidance for laboratories shipping specimens to WHO reference laboratories that provide confirmatory testing for COVID-19 virus: interim guidance, 2 March 2020

CDC/USA

- Interim Guidelines for Collecting, Handling, and Testing Clinical Specimens from Persons for Coronavirus Disease 2019 (COVID-19)
- Interim Laboratory Biosafety Guidelines for Handling and Processing Specimens Associated with Coronavirus Disease 2019 (COVID-19)

ZHEIJANG UNIVERSITY SCHOOL OF MEDICINE/CHINA

- Handbook of COVID-19 Prevention and Treatment

UCSF/USA

– UCSF Guidelines for Research Biospecimens during the COVID-19 Outbreak



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Biobanking in Times of COVID-19, #1 Risks and Opportunities (April 1, 2020) #2 Pre-Analytical Processes (April 7, 2020)

Questions and Answers

Q: Can Biobanks around the world be included in the COVID BE
A: Yes, the BBMRI-ERIC offers to include biobanks in the Direc
eric.eu/menu/main/app-molgenis-app-biobank-explorer/bioba
presentation of Petr Holub on BBMRI-ERIC website

Q: A general question: I believe it would be also feasible to loo genetic information that you have collected. Standardized metapatient stratification (after positive testing) and thereafter for t responder). If someone is interested I shall be happy to share in A: contact: bijon.chatterji@biocrates.com

Q: What type of samples are you collecting right now? Since w studies will be carried out. We are not sure what type of samples serum?

A: See presentation of Lukasz Kozera 7 April 2020 on the BBMF

Q: It should be usefull to know about other biological samples safe?

A: In any case handling biological sample is recommended to b Guideline https://apps.who.int/iris/bitstream/handle/10665/3: eng.pdf?sequence=1&isAllowed=y

Q: Any partnerships with any African countries for samples/data since the prevalence of the disease there is low?

A: African biobanks can be included in the Directory see https://directory.bbmrieric.eu/menu/main/app-molgenis-app-biobank-explorer/biobankexplorer

Q: What should those of us who run biobanks in countries where COVID19 has not really taken hold (yet!)? What should we do to prepare for as biobankers? Advice please.

A: Follow the web conference on 7 April 2020 and see the presentation of Lukasz Kozera and Helmuth Haslacher on the BBMRI-ERIC website

Q: How will BBMRI work with countries, especially China, that cannot send samples outside their borders?

A: Chinese biobanks can be included in the Directory see https://directory.bbmrieric.eu/menu/main/app-molgenis-app-biobank-explorer/biobankexplorer

Q: Could you also please share the links wher we could try to find all the documents related to the safety regulations and the quality of samples and their associated data?

A: See a list of documents https://www.bbmri-eric.eu/services/bbmriqm-covid and https://www.bbmri-eric.eu/services/standardisation/

Q: Is there support availble to enable researchers and biobanks to put in the right paperwork (contracts, MTA's etc) in place to actually get those samples send from the biobanks to the researchers?

CLINICAL ASPECTS AND BIOBANKING

Q: What about asymptomatic patients in high risk population?

AW 07 April 2020

ISO 20387:2018

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Annex A (normative) Documentation requirements

Annex B (informative) Implementation guidance for Annex A

Annex C (informative) Quality management system options

Bibliography



INTERNATIONAL STANDARD

ISO 20387

> First edition 2018-08

Biotechnology — Biobanking — General requirements for biobanking

Biotechnologie — «Biobanking» — Exigences générales relatives au «biobankina»



Reference number

© ISO 2018

ISO 20387:2018 WEB CON SERIES





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ON THE NEW BIOBANKING STANDARD ISO 20387:2018 "BIOTECHNOLOGY - BIOBANKING - GENERAL REQUIREMENTS FOR BIOBANKING"

From April 2019 to February 2020 the BBMRI.QM team provided our members and observers with an in-depth training on the new international standard "General Requirements for Biobanking, ISO 20387:2018".

This BBMRI-ERIC training and education programme has been presented as an online course, split into 22 sessions in which the individual chapters of the standard were discussed. Renowned experts gave comprehensive

presentations on requirements, 225 registrants from our member presentations regularly. The web and can be accessed at any time individual chapters.

If you are part of the BBMRI-Quality Management team:

Quality Management team:

Andrea Wutte / Quality Management team:

How have we been doing so far?

presentations on requirements, If you are part of the BBMRI-ERIC community and would like to have access to the recordings of our web conference series, please contact our 225 registrants from our member Ouality Management team:

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UNDERSTANDING COVID-19

Tell a Friend

A rapid, comprehensive, and relevant update on the current state of COVID-19, with attention to the epidemiology, clinical presentation, and emerging therapeutic options.

4/7/2020

When: April 7, 2020

9:00am PST/12:00pm EST

Where: United States