

# Biomedical models Hub

Final results

DG JRC

29<sup>th</sup> National Contact Point meeting

12/11/2025

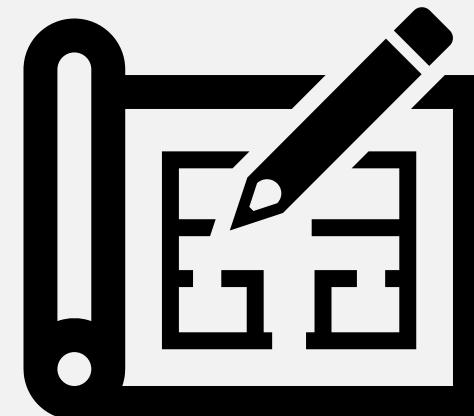


# BimmoH Project Results

**Leveraging Artificial Intelligence technologies to publish the largest dataset of human biology-based models used in biomedical research keeping it updated over time**

## SECTIONS

- 1. Introduction**
- 2. Objectives**
- 3. Method**
- 4. Results**

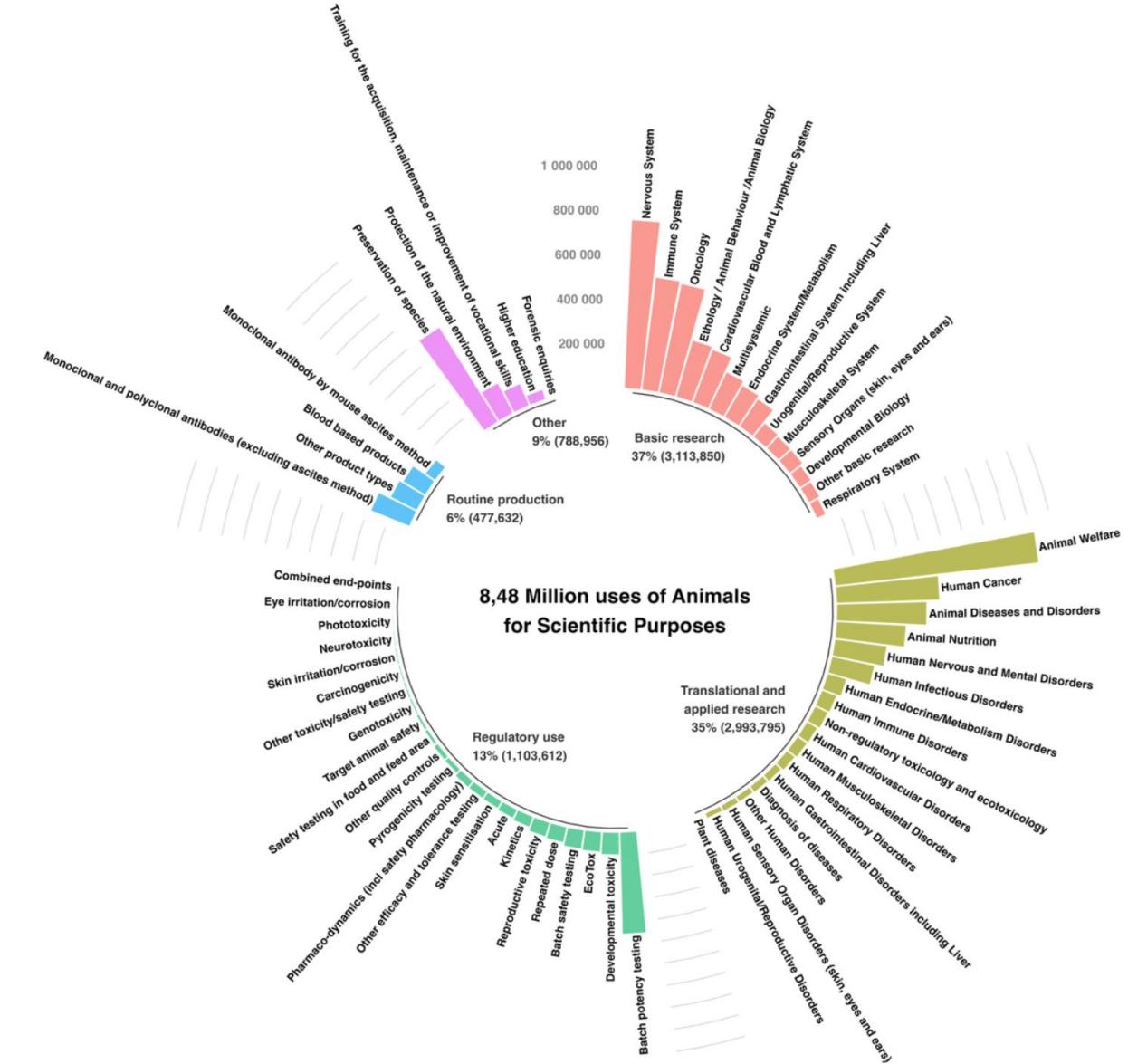


# Introduction



# Promoting alternatives in biomedical research

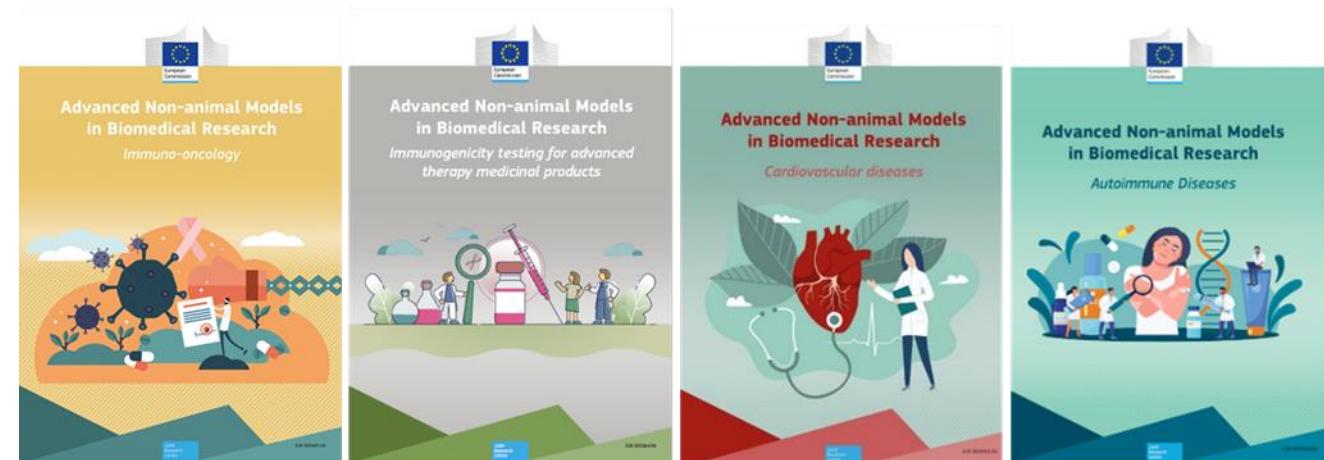
- Half of the uses of animals takes place in biomedical research
- Projects needs to be authorised by member states based on justification that there is no equivalent alternative
- Researchers must ensure that the 3Rs are assessed



# Biomedical reviews of non-animal models

Identify and describe specific research contexts where animal models have been put aside in favour of novel non-animal techniques (2013-2019)

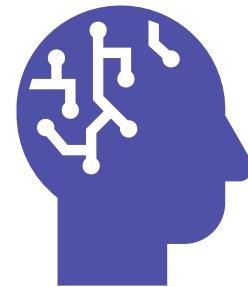
- Respiratory tract diseases
- Breast cancer
- Neurodegenerative diseases
- Immuno-oncology
- Immunogenicity testing for advanced therapy medicinal products
- Cardiovascular diseases
- Autoimmune diseases



# Automated approaches for systematic reviews

**Recent progress in machine learning, artificial intelligence and natural processing becoming mature enough to be implemented**

- Automated Literature Search and Screening (NLP)
- Meta-analysis and Data Interpretation (ML)
- Semi-automation and Human-in-the-loop Approaches (ML/AI)
- Active Learning and Reinforcement Learning (AI)



# Pilot Project



**Initiative or scheme designed to test and explore the feasibility, effectiveness, or usefulness of a particular idea, concept, policy, or program within a limited scope or timeframe.**

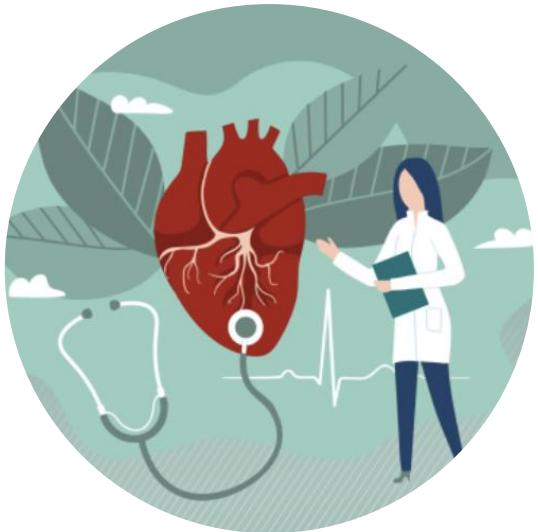
- Use of automated approaches to mine the vast body of published literature
- Enable the creation and maintenance of an up-to-date, state of the art knowledge source collating Non-Animal Models applied to biomedical research
- Allow the extension of models already collected by EURL ECVAM both in time and in scope



# BimmoH's Objectives



# BimmoH's primary goals



Extending biomedical reviews



Focusing on human-based biology models



Supporting stakeholders



# Definitions

## ***BimmoH Dataset***

Database of scientific articles references making use of human-based biology models selected by BimmoH's Machine Learning Classifier and indexed with BimmoH vocabularies.

## ***Machine Learning classifier***

Subset of Artificial Intelligence algorithms which build a model based on training data, to make predictions or decisions without being explicitly programmed to do so.

## ***Non-Animal model***

Research method that studies biological processes without making use of animals

## ***Human biology-based model***

Systems or methods used to replicate or simulate human biological processes, diseases, or drug responses for research purposes, based on human material or data.

## ***BimmoH Vocabularies***

List of key terms along with their definitions, used to index articles present in the BimmoH dataset.



# Human biology-based model identification

In BimmoH, a model is defined as any system capable of receiving an input and that allows recording an output. This includes human and/or animal models based on data or biological or biochemical materials, and may be:

- ***In vitro***: Living biological components outside a living organism.
- ***In silico***: Computational-based models.
- ***In chemico***: Chemical systems used for testing and analysis.

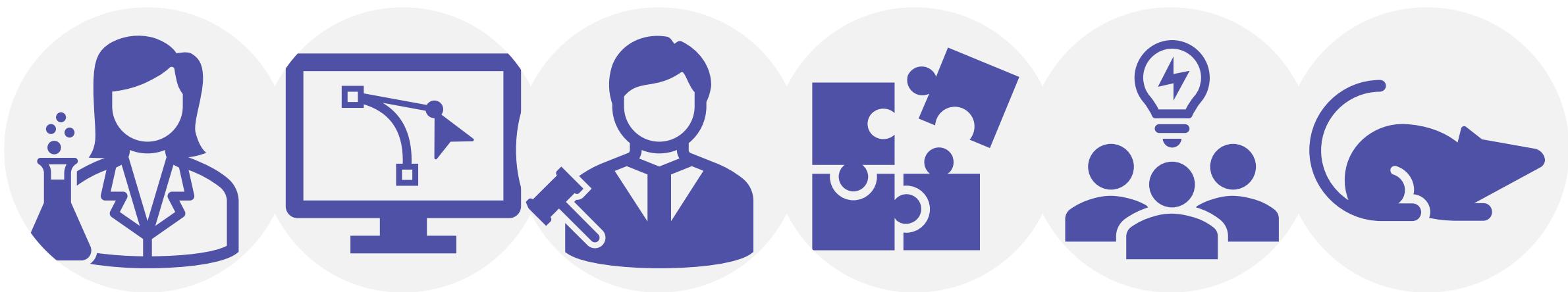
Studies using biological materials or data solely for analysis (i.e., biomarker detection in histological studies, omics studies, etc.) are not considered as models. Humans are not considered models for ethical reasons. They are test subjects under the governance of clinical trial regulations.

This definition was used by biomedical research experts to create select relevant scientific articles to train the Artificial Intelligence classifier and to validate its performance.



# User communities support

BimmoH's final goal is to promote further human-based biology models with the aim of increasing the translatability of basic and applied research while reducing the use of animal models.



Academic  
Researchers

Biomedical  
Data  
scientists

Regulators  
Policy  
Makers

Industry  
Innovators

Investment  
Analysis

Three Rs  
Organisations



# Requirements

1. Include articles that utilise human biology-based models, even if they also incorporate animal models
2. The database must be comprehensive, covering a broad range of diseases and conditions.
3. Support advanced search capabilities, including filters and keywords.
4. Provide detailed metadata for each model, by including study methodology.
5. Provide direct link access to original publications.
6. Ensure data consistency and accuracy through regular updates and expert validation.
7. Allow for the export of data and search results in various formats.
8. Offer training materials and resources for new users.
9. Ensure high performance and responsiveness of the database search functionality.
10. Develop a responsive and intuitive web interface for easy navigation.
11. Include visual aids and infographics to enhance data presentation.



# Methodological approach



# SWOT analysis and Stakeholder workshop

## Included requirements

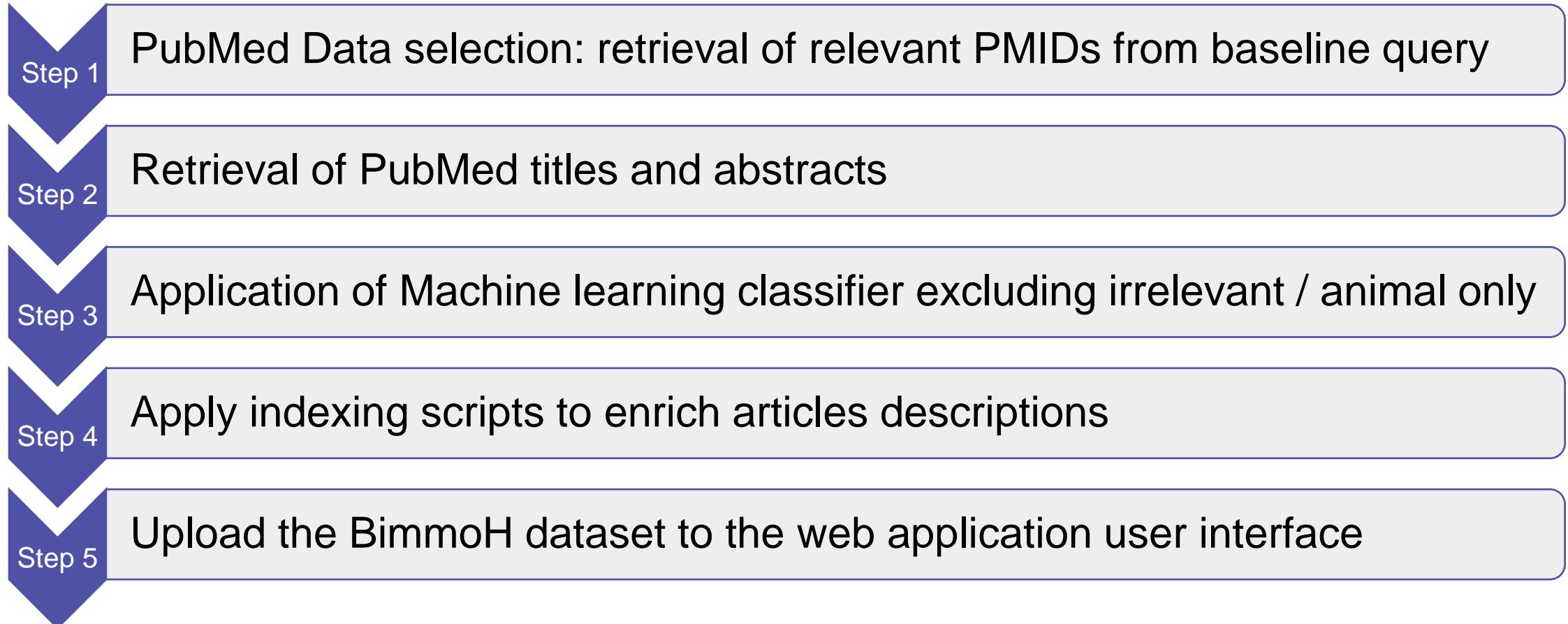
- Include both reviews and articles
- Go beyond identified disease areas to cover the whole biomedical research
- Include past and future article references (regular updates) compared to biomedical reviews
- Include articles even if animal models are used alongside human-biology based models
- Limit the analysis the title and abstract selection allowing wide screening

## Excluded or postponed

- Provision of full papers (but DOI links are included)
- Assessing Material and Methods section
- Provision of bibliographic and citation data
- Provision of information on human-based model (performance, validation, guideline adherence, etc.)
- Perform manual curation (inclusion/exclusion) of the AI collected data
- Identification of animal derived material



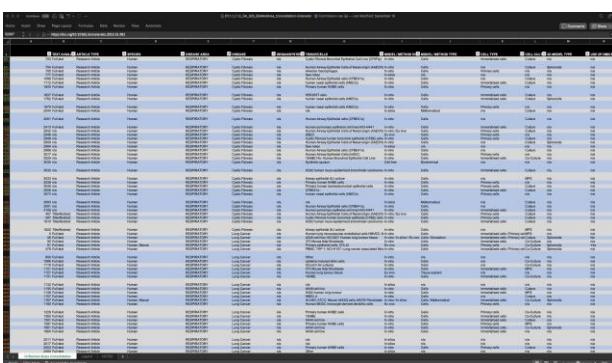
# Data process pipeline



# Data preparation

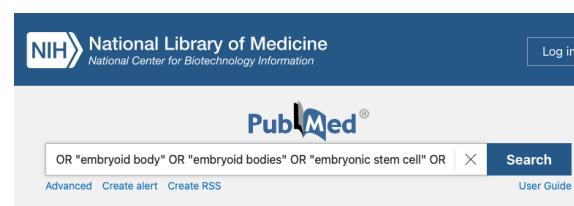
## Biomedical reviews data consolidation

- Single Excel file containing of seven biomedical reviews models
- Harmonisation of data (Disease, Organ, Tissue, Model type, Cell type, Omics use, Title, Abstract)



## PubMed baseline dataset creation

- PubMed query identifying Human-biology based models and excluding non relevant articles (clinical trial, case studies, etc...)
- Retrieving 4.8 million candidate articles to classify
- Using PubMed API to retrieve articles PMID



## PubMed data content and access

- PubMed APIs use for single article information retrieval
- Standard information (Title, Abstract, Author, DOI, PMID, etc.)

PMID - 40026989  
OWN - NLM  
STAT - In-Process  
LID - 1663-9812  
IS - 1663-9812 (Print)  
IS - 1663-9812 (Electronic)  
IS - 1663-9812 (Linking)  
VI - 16  
DI - 925  
T1 - Organoids: their emerging essential role in pathological mechanisms and drug discovery of diabetes and its complications.  
PG - 16502000  
LID - 16502000  
LID - 16502000 [doi]  
AB - Diabetes mellitus is a common disease with a high global prevalence, which often leads to complications, including diabetic retinopathy and nephropathy. Current treatments are mainly pharmacological and dietary. However, 3D organoid models have been used in the study of diabetes and have broad application prospects in the field of pharmacological research. We summarized the organoid models of diabetes and its complications. We analyzed the mechanism and its complications, and describe their sources, establishment and maturation methods, and their applications. We also summarized the contributions of organoids to the contribution of organoids in diabetes and its complications in terms of mechanistic studies, drug screening, and cell therapy. We also summarized the hope of providing a feasible direction for personalized medicine and precision treatment of diabetes and its complications. In addition, we discuss the strengths and weaknesses of organoid models in the study of diabetes and its complications. Nowadays, people strongly advocate personalized medicine and precision medicine. Organoid models have many advantages that are unmatched by any conventional experimental models. By combining organoid technology with high-throughput technologies, "patient-specific" drug screening can be realized, and personalized medicine. Organoids will also become a potential source of transplantable tissues and functional cell types for cellular replacement therapy. With the development of organoid technology, organoid assembly and vascularization techniques, organoids will gradually mature and improve. In conclusion, the 3D organoid system greatly complements the existing modeling system and may play a significant role in future basic and clinical research.

CT - China © 2025 Xu, Zhang, Geng, Lu and Sun.  
FAU - Xu, Xiaoyu  
AU - Xu, Xiaoyu  
AB - State Key Laboratory for Quality Eurance and Sustainable Use of Dao-di Herbs, Beijing, China  
AD - Lab of New Medicinal Plant Development, Peking Union Medical College and Chinese Academy of Medical Sciences, Beijing, China  
AD - Key Laboratory of Chinese Medicine Quality Control and Resource Utilization of Chinese Herbal Medicine, Ministry of Education, Beijing, China  
AU - Zhang, Yunqi  
AU - Zhang, Y



# Classification (identifying relevant articles)

## Developing a classifier based on supervised Machine Learning

- Low data requirement for training
- Allowing task specific optimisation
- Production of predictable and stable results
- Allowing transparency and interpretability
- Low computational cost
- Ease of update over time



# Indexing of selected articles

## Anatomy, Histology, and Cells

**Anatomical classification**  
lung, liver, skin, etc.

**Histological characteristics**  
epithelial, connective tissue, etc.

**Cell types**  
fibroblasts, neurons, hepatocytes, etc.

## Clinical Conditions, Disease, and Pathophysiology

**Specific diseases**  
diabetes, cancer

**Clinical conditions**  
inflammation, fibrosis

**Pathophysiological processes**  
immune dysregulation

## Models

**in vitro**  
living biological components outside a living organism

**in silico**  
computer-based simulations or predictions

**in chemico**  
purely chemical or biochemical test systems



# BimmoH Dataset – JRC Data Catalogue

## Data coming from PubMed:

- PMID
- DOI
- Authors information
- Bibliographical details
- Title
- Mesh terms

## Data coming from BimmoH:

- Model
- Disease
- Human anatomy
- Omics
- Cell line type
- Animal mentions

## Data coming from OpenAlex:

- Open Access availability
- Abstract (Key terms)



# BimmoH – Web User Interface

Available soon at <https://bimmoh.eu>

**BimmoH / BioMed Model Hub**  
A human biology based curated database designed to structure and consolidate information about models to support biomedical research.

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**Custom Filters**

**Anatomy, Histology, and Cells**

breast

✓ breast (278)

Show More... 1 of 972

AND OR NOT NONE

**Clinical Conditions, Disease and Pathophysiology**

cancer

✓ cancer (278)

Show More... 1 of 706

AND OR NOT NONE

**Models**

tissue

✓ tissue (278)

engineered tissue (0)

primary tissue (0)

bioprinted tissue (0)

scaffold-free tissue (0)

**Custom Filters**

Year

Open Access

Article Type

Author

MeSH

Human Cell Lines

Omics Techniques

Animal

**Free Text Search**

Search

Type to search

Reset filters

Find: All Words At Least One

**Free Text Search**

Find: All Words At Least One

immunotherapy

Fas (APO-1/CD95) receptor expression and new options for immunotherapy in childhood medulloblastomas.

Characterization of a new MUC1 monoclonal antibody (VU-2-G7) directed to the glycosylated PDTR sequence of MUC1.

The role of apoptosis in normal ontogenesis and solid human neoplasms.

Expression of SART3 antigen and induction of CTLs by SART3-derived peptides in breast cancer patients.

Phagocytosis of breast cancer cells mediated by anti-MUC-1 monoclonal antibody, DF3, and its bispecific antibody.

Identification of fibroblast growth factor-5 as an overexpressed antigen in multiple human adenocarcinomas.

Spontaneous cytotoxic T-cell responses against survivin-derived MHC class I-restricted T-cell epitopes *in situ* as well as *ex vivo* in cancer patients.

**Query History**

human\_anatomy

blood, breast, cell, lymphocyte, mononuclear, t-lymphocyte

T-cell, breast, cell, cellular, donor, epithelium, granulocyte, macrophage, monocyte

T-cell, breast, cell, cti, fibroblast, growth factor, lymphocyte, prostate, pulmonary, renal

T-cell, breast, cell, immune

cancer, rejection, tumour

adenocarcinoma, cancer

adenocarcinoma, cancer, carcinoma, lesion, melanoma, tumor

cancer, leukemia, melanoma, neoplasm, tumor

stem cell, tissue

cell line, tissue

In vitro, tissue

cell line, tissue

culture, tissue

cell line, tissue

ex vivo, tissue

Yes - Journal Article 1999 [Link](#)

Yes - Journal Article 2000 [Link](#)

Yes - Journal Article 2000 [Link](#)

Yes - Journal Article 2001 [Link](#)

No - Journal Article 2001 [Link](#)

No ZR-75-1 Journal Article 2001 [Link](#)

No - Journal Article 2001 [Link](#)

No - Journal Article 2001 [Link](#)

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**Reset filters**

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-  [Open Access](#) [?](#) [▼](#)
-  [Article Type](#) [?](#) [▼](#)
-  [Author](#) [?](#) [▼](#)
-  [MeSH](#) [?](#) [▼](#)
-  [Human Cell Lines](#) [?](#) [▼](#)
-  [Omics Techniques](#) [?](#) [▼](#)
-  [Animal](#) [?](#) [▼](#)

**Anatomy, Histology, and Cells** [?](#)  
[Anatomy, Histology, and Cells search](#)  
[Show All](#) 0 of 972

**Clinical Conditions, Disease and Pathophysiology** [?](#)  
[Clinical Conditions, Disease and Pathophysiology search](#)  
[Show All](#) 0 of 706

**Models** [?](#)  
[Models search](#)  
 [In silico](#)  [In chemico](#)  [In vitro](#)  
[Show All](#) 0 of 121

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 All Words  At Least One Word  Search Phrase [Search](#)

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- [Open Access](#) [?](#)
- [Article Type](#) [?](#)
- [Author](#) [?](#)
- [MeSH](#) [?](#)
- [Human Cell Lines](#) [?](#)
- [Omics Techniques](#) [?](#)
- [Animal](#) [?](#)

**Anatomy, Histology, and Cells** [?](#)

breast

[breast \(278\)](#)

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AND  OR  NOT  NONE

**Clinical Conditions, Disease and Pathophysiology** [?](#)

cancer

[cancer \(278\)](#)

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**Models** [?](#)

tissue

[In silico](#)  [In chemico](#)  [In vitro](#)

[tissue \(278\)](#) [engineered tissue \(0\)](#) [primary tissue \(0\)](#)

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AND  OR  NOT  NONE

**Free Text Search** [?](#)

immunotherapy

Find:

All Words  At Least One Word  Search Phrase

**Search**

**Reset filters**

**Query History**

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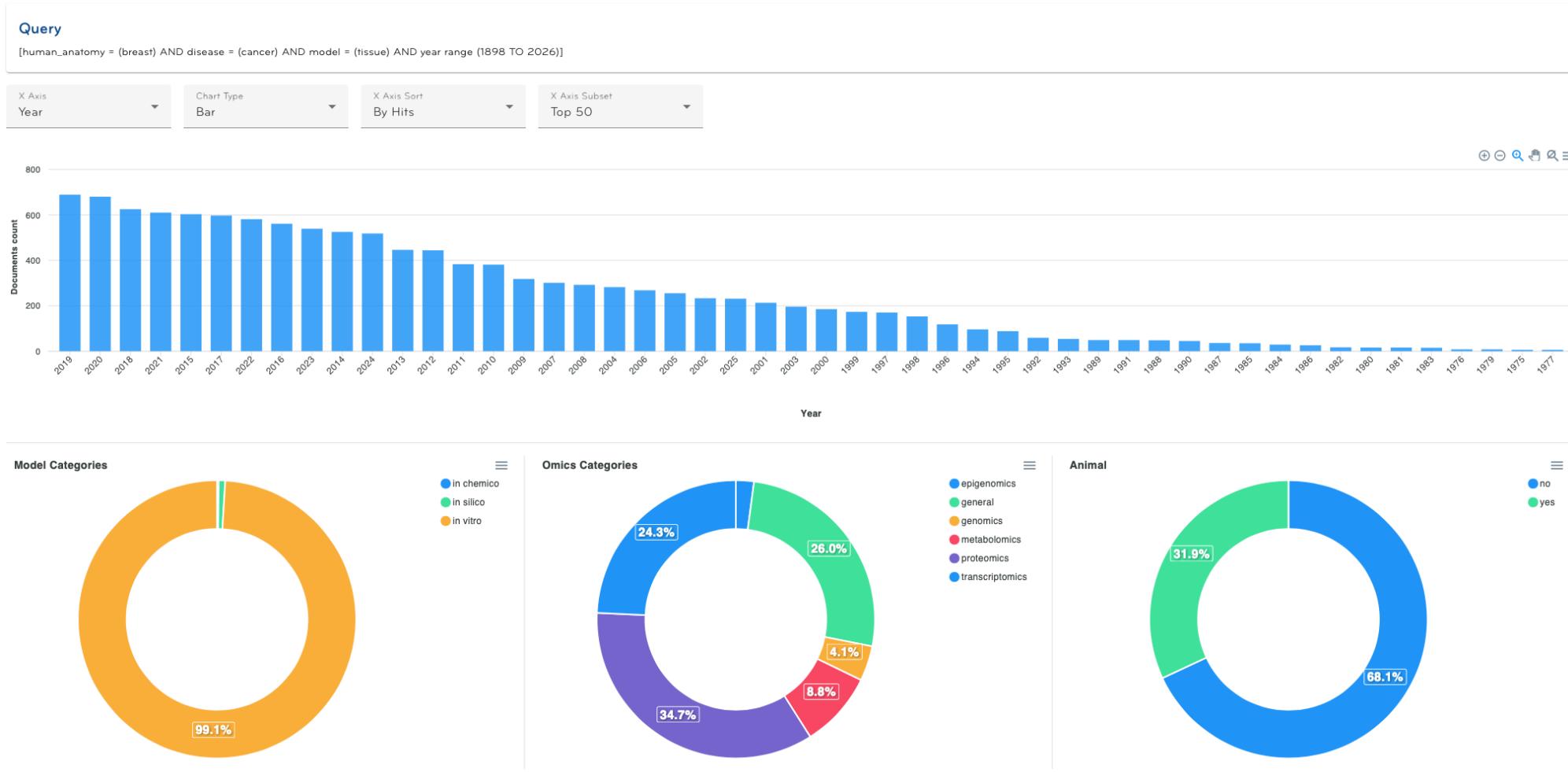


# BimmoH – Web User Interface

<input type="checkbox"/>	Title / Abstract / Author	Title	Anatomy, Histology, and Cells	Clinical Conditions and Pathophysiology	Models	Omics Techniques	Animal	Human Cell Lines	Article Type	Year	<a href="#">Link</a>	
<input type="checkbox"/>		Fas (APO-1, CD95) receptor expression and new options for immunotherapy in childhood medulloblastomas.	CNS, body, brain, breast, cell, cellular, colon, colorectal, embryonic, epithelium, glial, growth factor, hepatocellular, immunological, lung, membrane, nerve, nervous, neuroectoderm, nk, nucleus, pancreas, prostate, thymus, tonsil	cancer, carcinoma, leukemia, lymphocytic, malignancy, medulloblastoma, melanoma, necrosis, neoplasm, neoplastic, tumor	stem cell, tissue	-	Yes	-	Journal Article	1999	<a href="#">Link</a>	
<input type="checkbox"/>		Characterization of a new MUC1 monoclonal antibody (VU-2-G7) directed to the glycosylated PDTR sequence of MUC1.	breast, cell, membrane, ovarian, ovary	cancer, carcinoma, malignant, tumor	cell line, tissue	-	Yes	-	Journal Article	2000	<a href="#">Link</a>	
<input type="checkbox"/>		The role of apoptosis in normal ontogenesis and solid human neoplasms.	T-cell, body, brain, breast, cell, cellular, chromatin, colon, cortex, ctl, cytoplasm, gastric, glial, hepatocellular, lymphocyte, membrane, mitochondria, nucleus, ovarian, pancreas, renal, reticulum, stromal, t-lymphocyte, thymocyte, thymus, thyroid	Hodgkin, anaplastic, cancer, carcinoma, hyperthermia, infection, lymphoma, malignant, necrosis, neoplasm, neoplastic, sarcoma, tumor, viral	In vitro, tissue	-	Yes	-	Journal Article	2000	<a href="#">Link</a>	
<input type="checkbox"/>		Expression of SART3 antigen and induction of CTLs by SART3-derived peptides in breast cancer patients.	blood, breast, cell, lymphocyte, mononuclear, t-lymphocyte	cancer, rejection, tumour	cell line, tissue	-	No	-	Journal Article	2001	<a href="#">Link</a>	
<input type="checkbox"/>		Phagocytosis of breast cancer cells mediated by anti-MUC-1 monoclonal antibody, DF3, and its bispecific antibody.	T-cell, breast, cell, cellular, donor, epithelium, granulocyte, macrophage, monocyte	adenocarcinoma, cancer	culture, tissue	-	No	ZR-75-1	Journal Article	2001	<a href="#">Link</a>	
<input type="checkbox"/>		Identification of fibroblast growth factor-5 as an overexpressed antigen in multiple human adenocarcinomas.	T-cell, breast, cell, ctl, fibroblast, growth factor, lymphocyte, prostate, pulmonary, renal	adenocarcinoma, cancer, carcinoma, lesion, melanoma, tumor	cell line, tissue	-	No	-	Journal Article	2001	<a href="#">Link</a>	
<input type="checkbox"/>		Spontaneous cytotoxic T-cell responses against survivin-derived MHC class I-restricted T-cell epitopes <i>in situ</i> as well as <i>ex vivo</i> in cancer patients.	T-cell, breast, cell, immune	cancer, leukemia, melanoma, neoplasm, tumor	ex vivo, tissue	-	No	-	Journal Article	2001	<a href="#">Link</a>	



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And  Or    



Search by tta  

{ "condition": "and", "rules": [ { "field": "tta", "operator": "contains any", "value": "" } ] }

Import query 

Search by title or author or abstract 

No results

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# Conclusion



# BimmoH dataset - Statistics



**More than 750,000 articles  
with a 90% precision**

This is the largest dataset of scientific articles making use of human biology-based models kept updated over time



# Thank you



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