



OpenAIRE and Open Science

Elli Papadopoulou

OpenAIRE Greek National Open Access Desk
ATHENA Research & Innovation Center



MICCAI 24th | September 29, 2021



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MICCAI x OpenAIRE session

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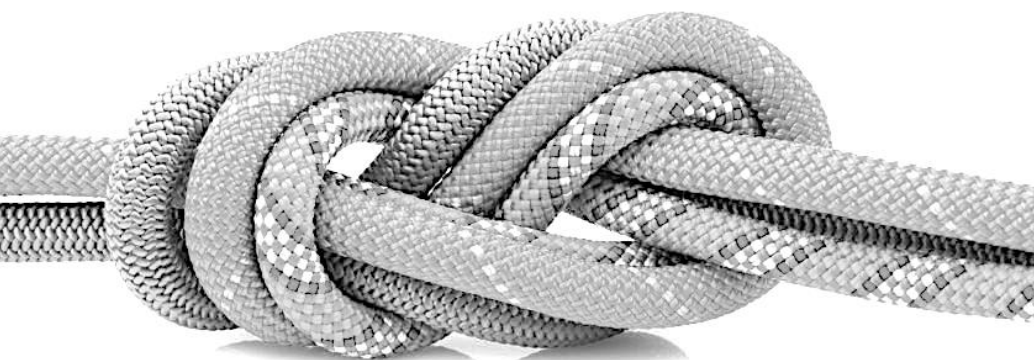
Or use QR code



Open Science



What is Open Science



Spiros Athanasiou et al, 2020. National Plan for Open Science.

Open Science is the **new standard of practices, means and collaboration** for producing and distributing scientific output and research results, with a direct scientific, economic and societal impact

Core principles

1. **Collaboration**
2. **Open Access**
Read and access scientific information
3. **FAIR* principles**
Discovery, Interoperability, Reuse
4. **Documentation**
Transparency, Accuracy

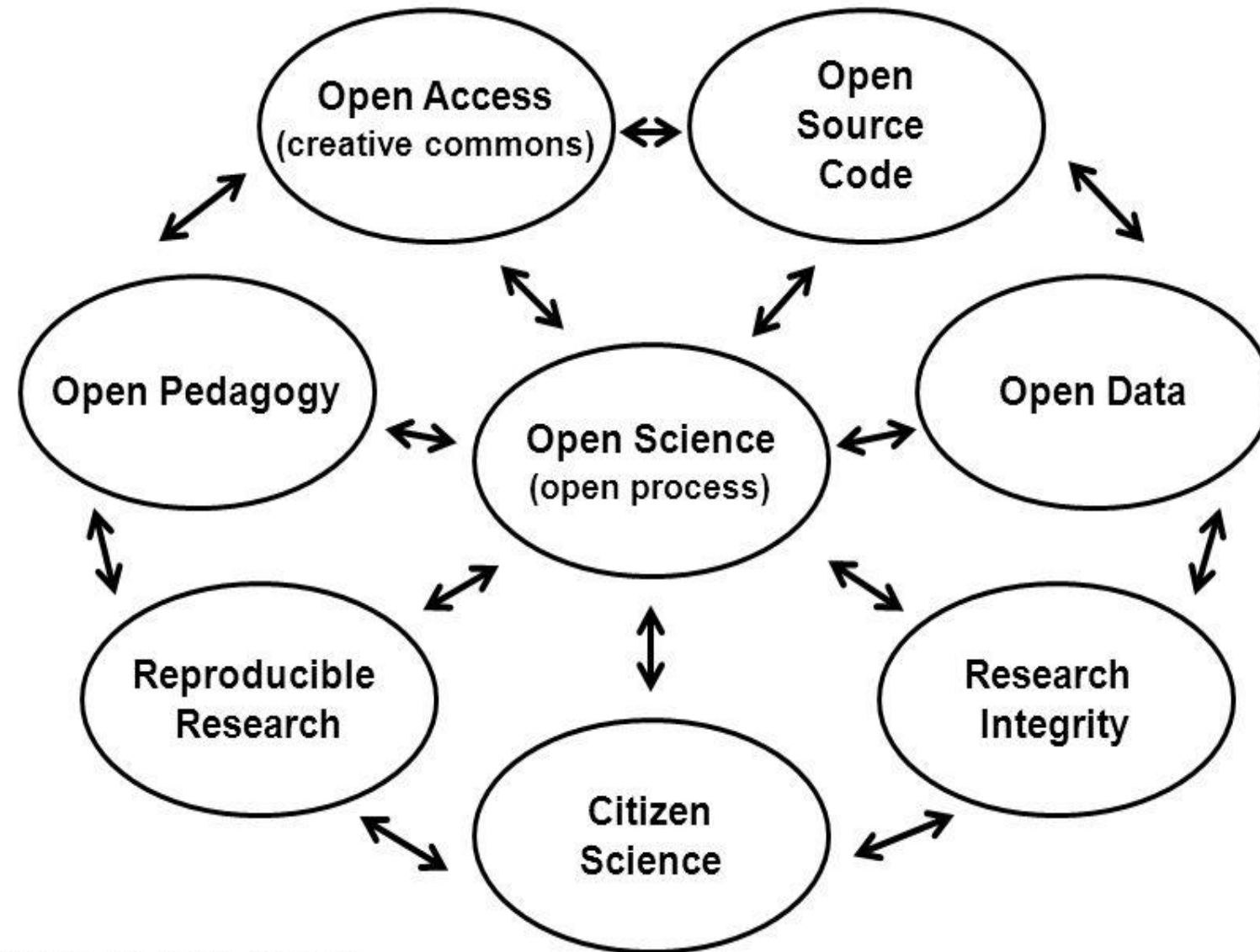
Areas of action

Scientific outputs

Infrastructures and services for research

Training and new skills

Open Science Ecosystem



With thanks to John Jungck

Why choosing Open Science?

Research

- Reach wider audience
- Re-use research outputs
 - Validate research
- Prevent information and data loss

Economy

- Stimulate innovation
- Strengthen regional and national markets
 - New job openings

Researchers

- Promote integrity
- Increase use of citations and get more credits
- Rewards in the EOSC

Society

- Builds trust ← transparency
- Inclusivity in Science → Citizen Science
- Collaborations on national and EU level

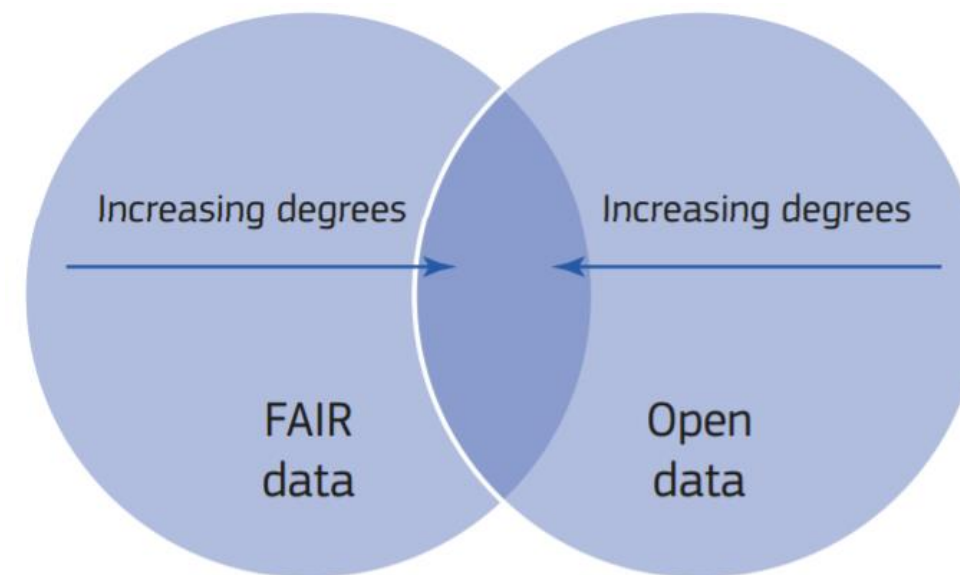


Research Data Management



EOSC - The Web of FAIR data

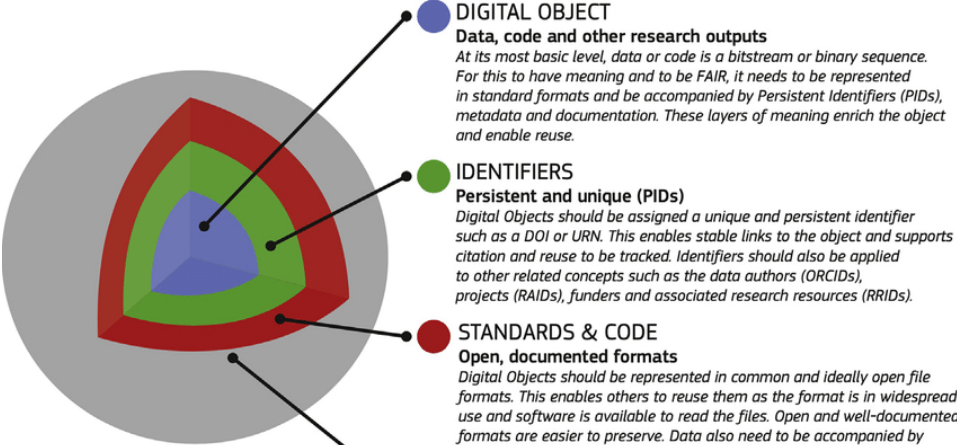
- FAIR-enabling ecosystems



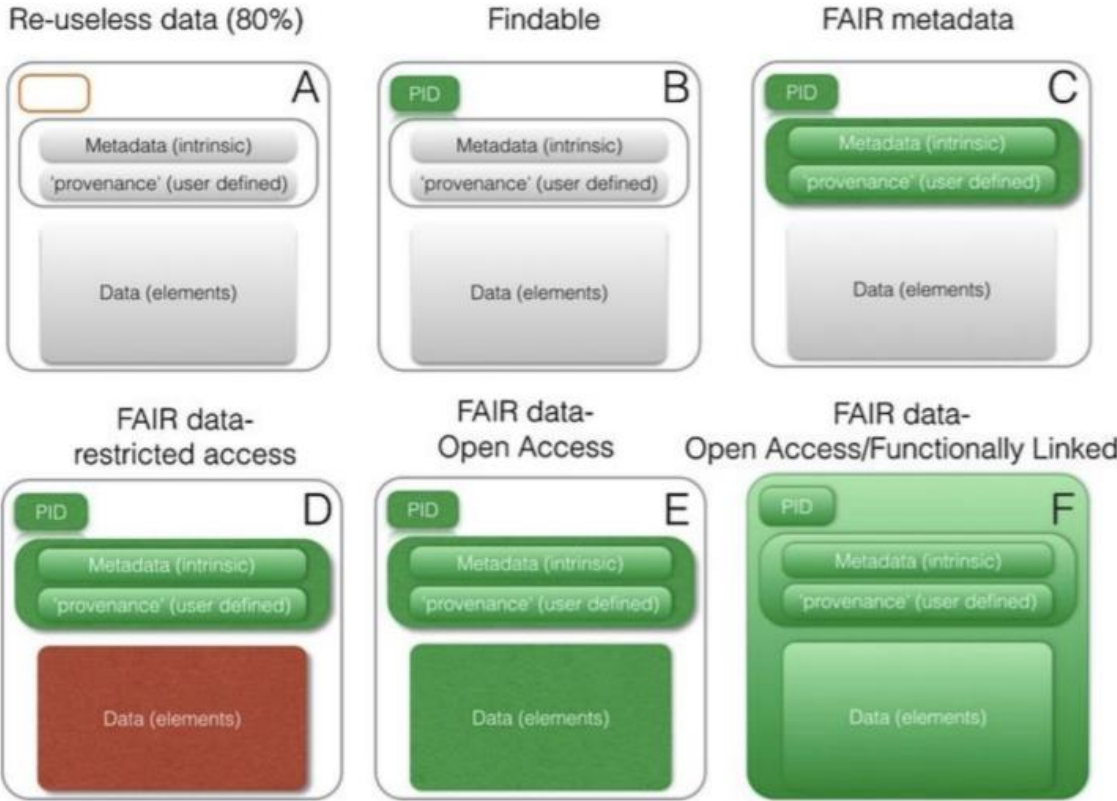
“highly distributed ecosystem requiring technical mechanisms linking resources, and social mechanisms to define specifications, standards and protocols”

FAIR principles for digital objects

Findable 	Persistent Identifiers (PIDs) 	Rich metadata 	Indexed data repositories 	PIDs in metadata
Accessible 	Standard communications protocol 	Open, free protocol 	Authentication, where necessary 	Metadata is always available
Interoperable 	Vocabularies 	Vocabularies are FAIR 	Linked metadata 	
Reusable 	Metadata have multiple attributes 	Usage license 	Provenance 	Community standards



Data as increasingly FAIR Digital Objects



<https://www.ands.org.au/working-with-data/fairdata/training>

RDM lifecycles



Table 1
Life cycle stages and identified questions

Stages	Questions
Identifying	<ul style="list-style-type: none"> What data are available? What is the current audience for these data? What potential future audiences exist for these data? Is this an isolated data set or could it be combined with other sets?
Digitizing	<ul style="list-style-type: none"> Are the data in digital format? If no, what would it take to digitize the data?
Cleaning	<ul style="list-style-type: none"> Are the data in a stable digital format that can be preserved? How many people have touched or will touch the data? What rules have been created to ensure consistent data standardization?
Describing	<ul style="list-style-type: none"> What tools am I using to standardize the data? Is there a README.txt file outlining the project? Is there a standard ontology applicable to this data set? What information would others need to use the data?
Storing and preserving	<ul style="list-style-type: none"> What access is needed to work with the data now? Who needs access now? What are the best storage options for the future? What is the intended duration of preservation?
Sharing	<ul style="list-style-type: none"> Are there any privacy concerns about these data? Who is the owner of this data set? What institutional policies apply to these data? How can sharing rights be maximized?
Analyzing	<ul style="list-style-type: none"> What analysis tools are available? What are the limitations of the data set?

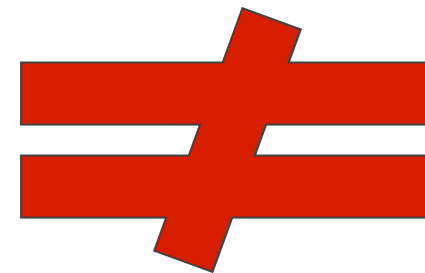
<https://datamanagement.hms.harvard.edu/about/what-research-data-management/biomedical-data-lifecycle>

Goben A, Raszewski R (2015) The data life cycle applied to our own data. J Med Libr Assoc. 103(1): 40–44. doi: [10.3163/1536-5050.103.1.008](https://doi.org/10.3163/1536-5050.103.1.008)

Data Management Plans

Deliverable and “living” document

- documents processes undertaken throughout data management lifecycle, including costs



What is not a DMP?

Research assessment method

Datasets

Title: OpenAPC

Template: Horizon 2020

This dataset describes the OpenAPC research data.

Dataset Description

1.1 Data Summary

1.1.1 What is the purpose of the data collection/generation and its relation to the objectives of the project?

["To share information", "To make informed decisions", "To combine with other data"]

Comment: Collecting and disseminating datasets on fees paid for open access publishing

1.1.2 What types of data will the project generate/collect?

["sensor data", "text mining", "observational (e.g., sensor data, data from surveys)", "derived or compiled (e.g., text mining, 3D models)"]

1.1.3 What formats of data will the project generate/collect?

[".txt files", "PDF", "RTF", "Text files - MS Word docs, .txt files, PDF, RTF, XML (Extensible Markup Language)"]

CSV

1.1.4 What is the origin of the data?

["Secondary data"]

1.1.5 What is the expected size of the data?

MB (megabyte)

1.1.6 To whom might it be useful ('data utility')?

["Researchers", "Research communities", "Decision makers", "Other"]

2.1 Reused Data

2.1.1 Will you re-use any existing data and how?

Yes

["To reproduce and validate findings", "To compare and combine with other data"]

2.1.3 Which data will be re-used?

Tables in csv format

Open Science in Europe



Article 29

Publications

Article 29.2 Grant Agreement
- Open Access
(check embargo periods)

“As open as possible,
as closed as necessary”

Article 29.3 Grant Agreement
- Open & FAIR data
- DMP “living document”

Research Data

 **Opting out
options**

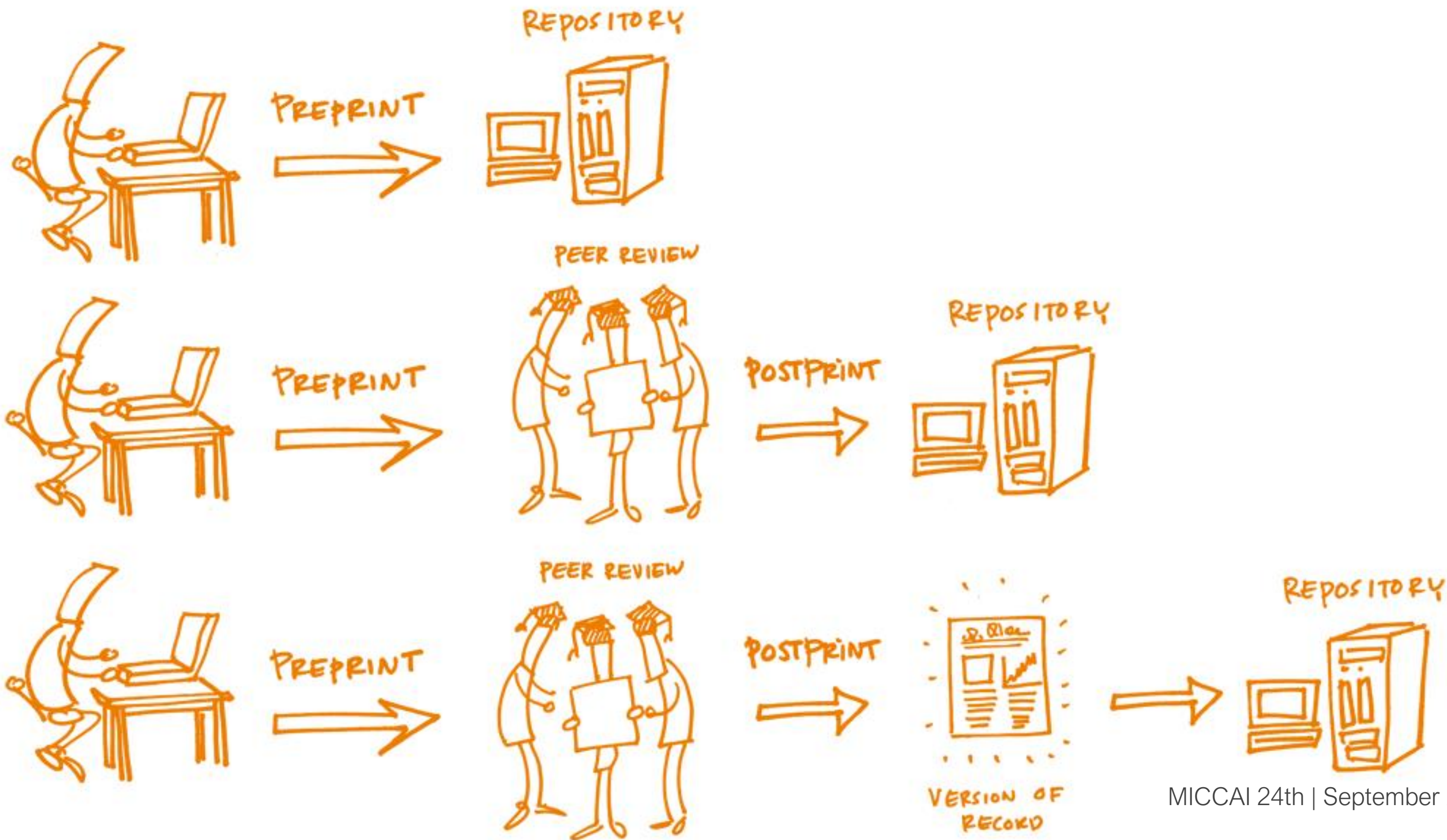


Access and preservation

- **Deposit a version of the publication in a literature repository and ensure open access**
 - Embargoes may apply: 6 months for STEM or 12 months for SSH
- **Publish research in an Open Access journal**
 - Article Processing Charges (APCs)

+ open access to metadata!

MODES OF SELF-ARCHIVING



Acknowledge funding

When you deposit, you must also ensure open access to the descriptive metadata that identify the deposited publication. This metadata must be in a standard format and must include all of the following:

- the words ["European Union (EU)" and "Horizon 2020"] ["Euratom" and Euratom research and training programme 2014-2018"];
- the name of the project, acronym and grant number;
- the publication date, and length of embargo period if applicable, and
- [a persistent identifier](#).

Research data


RESEARCH DATA - OPEN BY DEFAULT

Horizon 2020 grantees are required

take measures to ensure open access to the data underlying their scientific publications

provide open access to any other research data of their choice

Horizon 2020 grantees are encouraged to also share datasets beyond publication



1. Develop a Data Management Plan (DMP)
2. Deposit research data in data repositories
3. Provide open access to research data, if possible.
4. Make data FAIR

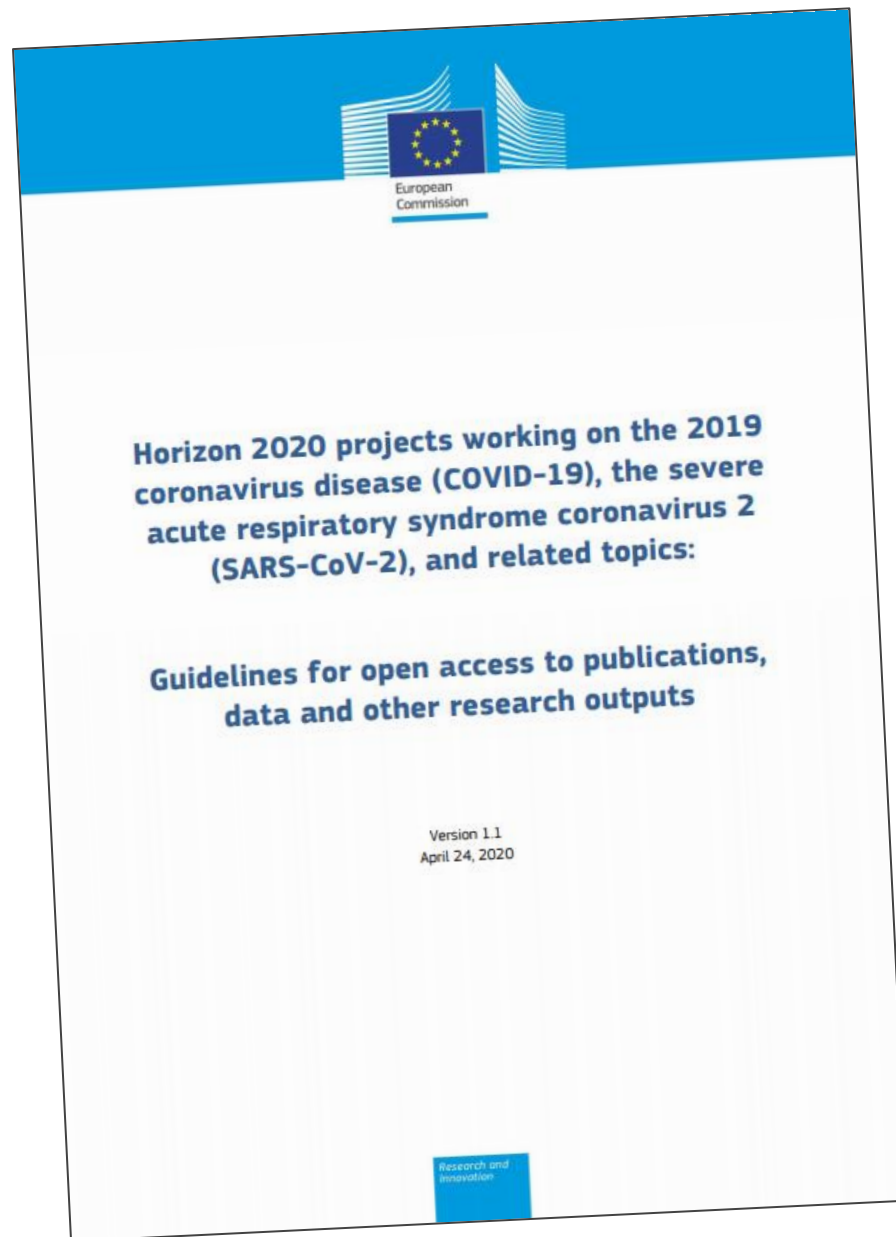
COVID-19 research

Open access data benefits millions of scientists around the world and is essential for a rapid response to the COVID-19 pandemic



Artist's impression of open access COVID-19 data sharing. Credit: Spencer Phillips, EMBL-EBI.

Open Access for COVID-19 research



- **All peer-reviewed research publications** relevant to the outbreak are made immediately open access, or freely available at least for the duration of the outbreak;
- Research findings relevant to the outbreak are shared **immediately with the WHO upon journal submission**, by the journal and with author knowledge;
- Research findings are made available via preprint servers before journal publication, or via platforms that make papers openly accessible before peer review, with clear **statements regarding the availability of underlying data**;
- Share interim and final research data, **together with protocols and standards** used to collect the data, as rapidly and widely as possible - **including with public health and research communities and the WHO**;
- Authors are clear that data or preprints shared ahead of submission will not pre-empt its publication in these journals.

https://ec.europa.eu/research/participants/data/ref/h2020/other/hi/oa-pilot/h2020-guidelines-oa-covid-19_en.pdf

OpenAIRE



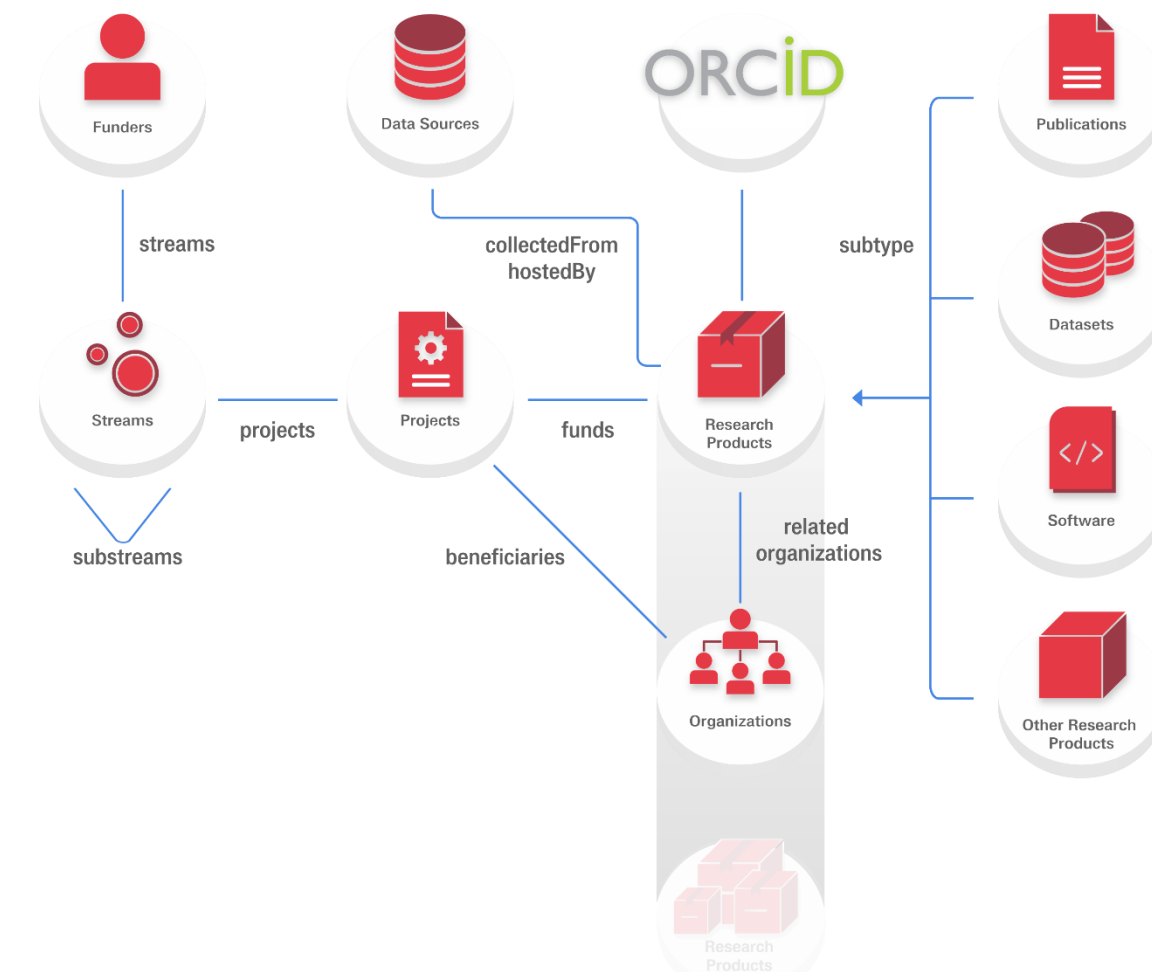


About OpenAIRE



Paneuropean electronic infrastructure: added value **horizontal services**

- **Distributed** and **participatory** electronic infrastructure
 - Based on existing national infrastructures
 - Interconnection of various actors across Europe
- **Linked** Open Science
 - Support for all types of research entities and outputs
- **Human** network
 - National nodes in more than 34 countries

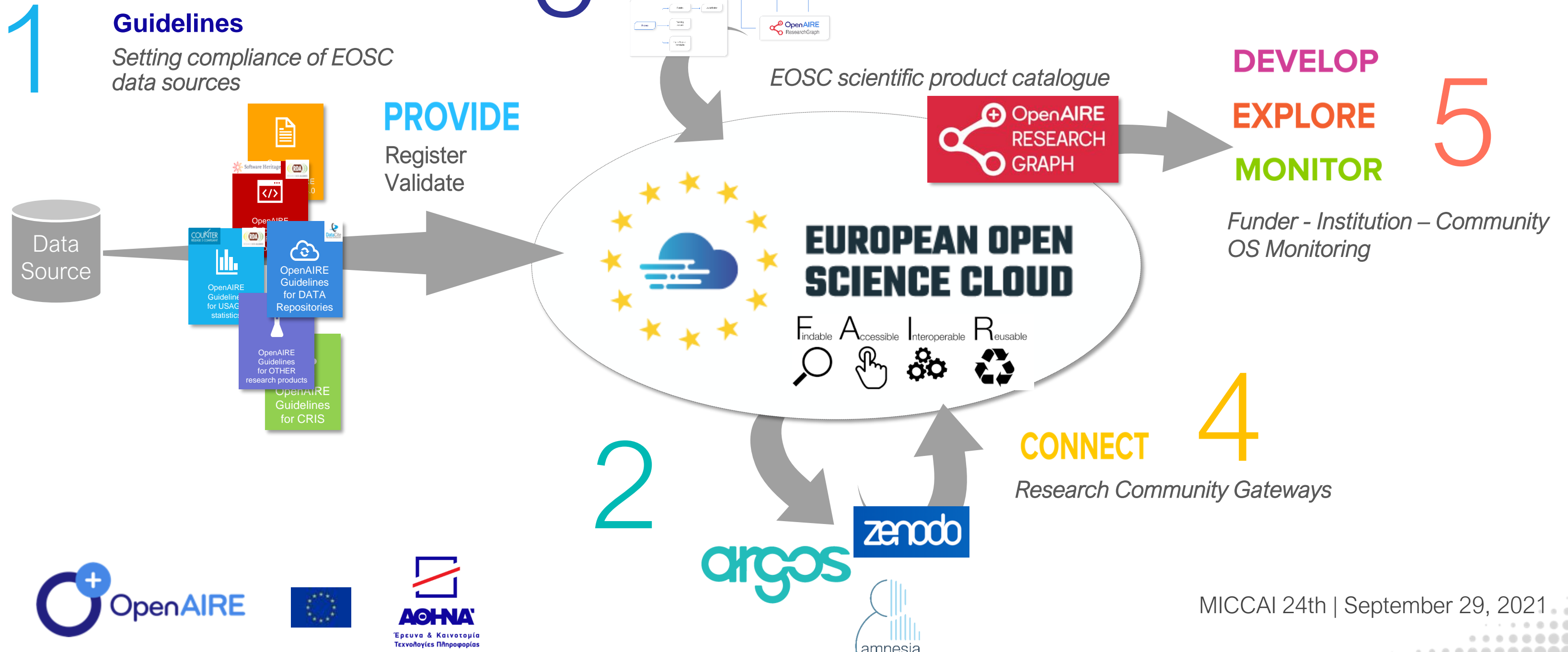


EXPLORE



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OpenAIRE services in EOSC





Policies

Aligning

Training

Empowering

Services

Connecting

OpenAIRE

OpenAIRE RESEARCH GRAPH
Putting research into context.
Making the connections.



explore.openaire.eu

USAGE ANALYTICS
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a catch-all-repository



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across content providers



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Interlinking data
and publications



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Helping you apply
the OpenAIRE Guidelines



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Discover and share research




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OpenAIRE | DEVELOP
APIs to access the
OpenAIRE Research Graph



develop.openaire.eu

argos
Plan and follow
your data



openaire.eu/argos



Resources

How to comply with H2020 mandates - publications

How can identifiers improve the dissemination of your research outputs?

Legal issues

How do I know if my research data is protected?


























How do I license my research data

Can I reuse someone else's research data

Data reuse stories & use cases

Series of stories and use cases reporting the process of data reuse, describing experiences of reuse of a variety of research data.

ALL A B C D E F G H I J K L M N P R S T U

 Austria Gerda Mc Neil University of Vienna	 Belgium Inge Van Nieuwerburgh University of Ghent	 Belgium Emilie Hermans University of Ghent	 Bulgaria Peter Stanchev Bulgarian Academy of Sciences	 Bulgaria Georgi Simeonov Bulgarian Academy of Sciences
 Croatia Jadranka Stojanovski Ruđer Bošković Institute	 Cyprus Sylvia Koukouridou University of Cyprus	 Cyprus Natasia Ioannou University of Cyprus	 Cyprus Zafeiro Marti University of Cyprus	 Czech Republic Michal Růžička Masaryk University
 Denmark Asger Vaering Larsen Syddansk Universitet	 Denmark Anne Thorst Melbye Syddansk Universitet	 Estonia Lisi Lembinen University of Tartu	 Finland Pauli Assinen University of Helsinki	 Finland Kimmo Koskinen University of Helsinki
 France Andre Dazy Couperin	 Germany Anja Oberlander University of Konstanz	 Germany Lena Dehor University of Konstanz		
 Hungary Gyongyi Karacsony University of Debrecen	 Hungary Judit Fazekas-Paragh University of Debrecen	 Ireland Anna Sigríður Guðnadóttir Lundskott University Hospital	 Ireland Niamh Brennan Trinity College Library Dublin	 Ireland Eddie Davies Trinity College Library Dublin

onymize your data

For researchers



Register & validate your repository

Researchers

Tips on H2020 OA Requirements Compliance & OpenAIRE Tools to Support Reporting
Monday, 29 April 2019

Report your publication and data to the EC

For project coordinators



Claim a publication or data to your funding

For researchers



Content Enrichment

For Repository Managers

Track the usage activity of your repository

Find a NOAD in your country!
<https://www.openaire.eu/contact-noad>



Useful Links

- OpenAIRE Helpdesk: <https://www.openaire.eu/support/helpdesk>
- OpenAIRE NOADs: <https://www.openaire.eu/contact-noads>

- OpenAIRE Factsheets: <https://www.openaire.eu/openaire-h2020-factsheets>
- OpenAIRE Research Community Dashboard: <https://connect.openaire.eu/>
- OpenAIRE webinars:
- <https://www.openaire.eu/frontpage>

Don't miss out!

Sign up to [OpenAIRE](#) and subscribe to the [Newsletter](#) to get informed about news and updates

Thank you!

Elli Papadopoulou

✉ elli.p@athenarc.gr

ORCID 0000-0002-0893-8509

🐦 @elli_lib



Benefits and concerns of biomedical data sharing

Reasons to share individual-level data

To improve science

- Enable verification, replication, and expansion of research results
- Address biases, deficiencies, and dishonesty in research
- Enable novel analyses and increase study power
- Improve meta-analyses
- Maximize data use, particularly for datasets that cannot be replicated
- Inform research design and research funding
- Improve teaching resources
- Increase primary data producers' academic profiles and collaboration opportunities

To improve health

- Inform health care planning and allocation
- Inform regulatory review
- Improve evidence base for clinical decision making
- Improve use of health care resources
- Improve patient care

Explicit moral claims

- Importance of maximizing the value and utility of data
- Promotion of scientific values
- Promotion of best practices in research conduct, analysis, and reporting
- Demonstration of respect for research participants
- Promotion of the public good

Concerns about sharing individual-level data

May hamper science

- Reputational harms of critical secondary analyses
- Consequences of flawed/poor quality secondary analyses
- Reduction of incentives for primary research
- Increased incentives to conduct short-term research rather than long-term research
- Opportunity costs of curating and sharing data

May hamper health

- Effects of flawed secondary analyses on scientific evidence base
- Burden of evaluating validity of secondary analyses
- Effects of second-guessing regulatory procedures, policies, and processes

Explicit ethical issues

- Protection of participants' privacy and confidentiality
- Validity of consent, including broad consent
- Potential harms of secondary research for research participants including discrimination and stigma
- Researchers' ability to fulfill commitments made to research participants during data collection
- Effects of moral distance and limited awareness of the context in which data were collected
- Potential impacts on public trust and confidence of conflicting analyses
- Balancing the interests of differing stakeholders in data sharing
- Making best use of limited research resources

Barriers to sharing

- Costs of developing and maintaining appropriate expertise and infrastructure
- Curation costs
- Ownership, intellectual property rights, and commercial confidentiality
- Lack of policies and processes

Bull S, et al. (2015) Views of Ethical Best Practices in Sharing Individual-Level Data from Medical and Public Health Research. Journal of Empirical Research on Human Research Ethics. <http://dx.doi.org/10.1177/1556264615594767>