DELIVERABLE D5.2

DATA MANAGEMENT PLAN

Unlocking the scientific excellence and innovation capacity of the University of Aveiro in supramolecular multicomponent biomaterials for enabling advanced biomaterials for healthcare

(Grant Agreement no. 101079482)

By the University of Aveiro

Funded by
the European Union

The SUPRALIFE project has received funding from the European Union's Horizon Europe research and innovation programme under grant agreement No. 101079482.

DELIVERABLE D5.2: DATA MANAGEMENT PLAN

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WORK PACKAGE (WP): WP5 – Dissemination, exploitation, and communication activities

DELIVERABLE: D5.2

LEAD BENEFICIARY: 1. UAVR (University of Aveiro)

DELIVERABLE TYPE: DMP – Data Management Plan

DISSEMINATION LEVEL: PU – Public

DUE DATE: M6 (30/06/2023)

DELIVERY DATE: 30 June 2023

DOCUMENT HISTORY

Version Date	Date	Description
V1.0	11 June 2023	First Version prepared of the Data Management Plan (DMP) – Draft – sent to revision by João Borges
V1.1	20 June 2023	Input by João Borges
V1.1	21 June 2023	Inputs into the first draft incorporated – sent to revision by Luísa Sal
V1.2	22 June 2023	Revision from Luísa Sal
V1.3	23 June 2023	Draft version of the DMP sent to revision by TU/e and UBx/IPB/CNRS
V1.4	29 June 2023	Revision from TU/e and UBx/IPB/CNRS
V2.0	30 June 2023	DMP Submitted

CONTENTS

LIST OF ACRONYMS AND ABBREVIATIONS

Table 1. List of acronyms and abbreviations.

Acronym / Abbreviation	Meaning / Full Text
UAVR	University of Aveiro
TU/e	Eindhoven University of Technology
UBx	University of Bordeaux
M6	Month 6
M9	Month 19
DMP	Data Management Plan
IPR	Intellectual Property Rights
CA	Consortium Agreement
FAIR	Findable, Accessible, Interoperable, Reusable
ORE	Open Research Europe
DOI	Digital Object Identifiers
D&C	Dissemination and Communication
OpenAIRE	Open Access Infrastructure for Research in Europe
OA	Open Access
DPO	Data Protection Officer
GDPR	General Data Protection Regulation

LIST OF TABLES

1. EXECUTIVE SUMMARY

This document represents the initial version of the Data Management Plan (DMP) delivered in Month 6 (M6; Deliverable 5.2) of the SupraLife project. It provides an overview of the datasets that will be produced by the project and specifies the associated conditions. The DMP will be revised and updated on Month 19 (M19; Deliverable 5.3), in line with the project progress, and to disclose the status of the project's reflections on data management.

The objective of this document is to:

- Facilitate the data management life cycle for all data to be collected, processed, or generated within the SupraLife project;
- Identify the most effective methods for gathering information on the diverse range of data to be utilized in the SupraLife project.

The SupraLife's DMP outlines all types of data, that will be generated or collected during the project, the standards that will be followed, the methods for preserving that data, and the portions of the datasets that will be shared for verification or reuse. It also aligns with the current agreements within the Consortium Agreement (CA) regarding data management and must be consistent with exploitation and Intellectual Property Rights (IPR) requirements.

The DMP is a dynamic document that will evolve throughout the project's timeframe, especially in response to significant changes such as dataset updates or modifications in Consortium policies.

This document will be used as a manual to describe the data management life cycle for the data to be collected, processed, and/or generated by the project, in line with the FAIR principles (Findable, Accessible, Interoperable, Reusable).

2. OVERVIEW OF THE DATA MANAGEMENT PLAN

The purpose of data generation and collection within the SupraLife project is to fulfill the project's main objectives:

- Significantly strengthening the science, technology and innovation capacity of the University of Aveiro (UAVR) in supramolecular biomaterials' chemistry research field to achieve scientific excellence and competitiveness on the development of advanced supramolecular multicomponent biomaterials for healthcare;
- Position the UAVR and its staff in an international competitive level in this research field, significantly enhancing its research and innovation capabilities, international visibility, capacity to successfully compete for funding and attract new research organizations and business stakeholders.

Moreover, the overarching goals of the SupraLife project can be summarized by ten primary objectives, which are comprehensively described on Table 2. Additionally, an overview of the data collection that will be generated or reused for each specific objective within the SupraLife project is provided on Table 2.

Table 2. Data collection/generation and its relation to the objectives of the SupraLife project.

Objectives (O) of the SupraLife Project	Data Collection/Generation
O1: Enhance the knowledge, scientific excellence, and research profile of UAVR and its staff in supramolecular biomaterials; Significantly increase the international visibility of UAVR and unleash its networking gaps with internationally leading R&D organizations and enterprises	Personal, scientific and project data related to: • Exchange of PhD students and post-doc researchers through short-term on-site training activities; • Exchange of staff researchers through short-term exchanges/visits; • Expert visits; • Collaborations and new partnerships with leading research organizations, enterprises, hospitals, and regulatory authorities; • Organization of training and networking activities; • Collaboration on joint research projects with partner institutions and external guests; • Training and research initiatives involving the UAVR's staff; • Increasing scientific and technological awareness among stakeholders.
O2: Increase the number of scientific publications in high impact journals and highly cited ones	Scientific data related to: • The development of the SupraLife's exploratory research project.
O3: Create strong, long-lasting links between UAVR-TU/e-UBx for enabling a steadily and sustained knowledge exchange and transfer and research collaboration that will continue in the long-term	Personal, scientific and project data related to: • Organization of workshops; • Joint summer school-type events, symposia, scientific retreats, one final international conferences, short-term on-site training activities and staff exchanges; • Long-lasting mobility and training activities among the European partners; • Application for joint research projects in the scope of SupraLife's project; • Training actions for internal UAVR's researchers and training of new students to be hired in near future.
O4: New training strategies adapted to diverse scientific backgrounds	Personal data related to: • Participants of joint summer school-type events and workshops.

O5: Establishment of new partnerships with internationally leading research organizations and enterprises	Personal and project data related to: • Participation in symposia in international conferences for establishing new contacts and synergies with distinguished scientists linked to internationally leading research organization within academia and industry; • Increasing the visibility within the scientific community; • Exchange of students with distinguished leaders of research organization and enterprises; • Staff visits to internationally leading R&D organizations and industries/business players; • Recruiting new students/researchers to be part of UAVR as highly talented and qualified professionals; • The invitation of distinguished researchers and industrial leaders to the SupraLife's training activities.
O6: Establishment of new research lines and jobs in institutions in the future, namely in the Widening partner	Personal, scientific and project data related to: • Preparation of ground-breaking research projects with, but not limited to the consortium partners to stimulate the innovation capacity of UAVR; • Fomenting the exchange of staff members using funding from other mobility programs (e.g., Erasmus+).
O7: Training and maintenance of highly talented researchers at UAVR and attract new highly qualified in a sustainable manner	Personal data related to: • Training and skills development activities in the scope of the SupraLife project; • Implementation of new research projects to pursue new research work and future developments in the project field and cross-related areas.
O8: Generation of intellectual property	Scientific and personal data related to: • Collaborations and networking contacts/partnerships with academic organizations and biomedical/biotechnological companies.
O9: Regional economic and social development	Personal and scientific data related to: • The creation of new and innovative research lines and jobs in long-term for highly skilled researchers trained in SupraLife project.
O10: Setting a base for getting funds for common research activities	Personal, scientific and project data related to: • The establishment of a stable, long-lasting collaborative network with internationally leading organizations; • The training activities and actions organized in the scope of the SupraLife project; • Sustainable collaborative network within and beyond the project timeframe.

3. TYPES AND FORMATS OF DATA GENERATED WITHIN THE SUPRALIFE PROJECT

Throughout the SupraLife project, various types of data will be generated and collected, including **scientific, personal, and project data**. These data may be generated and collected in different formats, as highlighted on the Table 3.

Table 3. Types and formats of data generated in the framework of the SupraLife project.

	Type	Format
Scientific Data	Datasets from experiments	– Analysis of the chemical modification of polymeric materials and physicochemical, mechanical, morphological and biological evaluation of the developed supramolecular polymeric biomaterials by multi-nuclear NMR **(*.fid, *.1r)**; matrix-assisted laser desorption/ionization time-off-flight mass spectrometry, gas permeation chromatography-mass spectrometry, liquid chromatography-mass spectrometry – data **(*.mzxml)**; high-performance liquid chromatography – data **(*.xml)**; circular dichroism, small- and wide-angle X-ray scattering, dynamic light scattering, UV-Visible spectroscopy, attenuated total reflectance-Fourier transform infrared spectroscopy, fluorimeter, zeta-potential, quartz crystal microbalance with dissipation monitoring, X-ray diffraction, Instron, rheometer, microplate reader, flow cytometry – data **(*.csv)**; real-time polymerase chain reaction – data **(*.rdml or *.rdm)**; 3D printing/bioprinting, atomic force microscopy, transmission electron microscopy, scanning electron microscopy, fluorescence microscopy, confocal laser scanning microscopy, optical microscope – data images **(*.jpg, *.png or *.tiff)**.
	Publications	Research results and outcomes will be reported in original research articles and review papers in peer-reviewed international journals, books, book chapters, as well as in conference papers/proceedings, websites, social media networks, and other means, as text documents **(*.pdf *.doc or *.docx)**, or images **(*.jpg, *.png or *.tiff)**.
	Communication materials (posters, presentations, flyers, etc)	Data will be gathered in various formats, including text documents **(*.doc, *.docx or *.pdf)**, Power Point slides **(*.ppt or *.pptx)**, images **(*.jpg, *.png or *.tiff**). These data will be shared in events (e.g., conferences, summer school, symposia, workshops, seminars, webinars).
	Meetings with stakeholders from industry	Data will be collected as text documents **(*.doc, *.docx or *.pdf)**, or PowerPoint slides **(*.ppt or *.pptx)**.
Personal Data (will be collected in compliance with the EU General Data Protection Regulation 2016/679)	Participants attending events (conferences,	yyyyData such as name, nationality, institution, e-mail addresses, organization address, and other contacts as well as any other information deemed necessary for the specific activity. The

	summer school-type events, training activities, workshops, scientific retreats, etc.)	data will be gathered in various formats, including text documents **(*.txt, *.doc, *.docx or *.pdf)**, PowerPoint slides **(*.ppt or *.pptx)**, images. **(*.jpg, *.png or *.tiff)**, text **(*.csv)**, and excel files **(*.xlsx)**.
	Partners/Consortium	Data such as name, nationality, institution, e-mail addresses, organization address, and other contacts. The data will be gathered in various formats, including text documents **(*.doc, *.docx or *.pdf)**, PowerPoint slides **(*.ppt or *.pptx files)**, images **(*.jpg, *.png or *.tiff)**, and excel files **(*.xlsx)**.
	Staff	Most of this personal data will be stored in the form of text documents **(*.doc, *.docx or *.pdf)** or excel tables **(.xlsx)**. Individuals will be duly informed about the purpose of these initiatives, as well as the methods of data collection and processing. Only the essential personal data required will be recorded, and it will be maintained in a secure and confidential manner.
Project Data	Reports, Deliverables, Oral/poster Presentations	Continuous reporting on project implementation and work progress, deliverables, and meeting presentations (available for members of the Consortium and European Commission). Data will be stored as text documents **(*.doc, *.docx or *.pdf),** excel tables **(*.xlsx)** and PowerPoint slides **(*.ppt or *.pptx)**.

4. RE-USE OF EXISTING DATA

The SupraLife consortium is committed to making the generated scientific data available, primarily through the following channels:

Publications: Peer-reviewed scientific publications (Open Access (OA) journals) and underlying data, books, and book chapters, created by the SupraLife beneficiaries, will be made freely available on the project's website, published in OA on the Open Access Infrastructure for Research in Europe (OpenAIRE), and in the institutional repositories of the consortium partners (e.g., RIA repository – Institutional repository of UAVR, https://ria.ua.pt/; TU/e repository – Institutional repository of TU/e, https://research.tue.nl/; OSKAR Bordeaux – Institutional repository of UBx, https://oskar-bordeaux.fr/; HAL-CNRS – repository of CNRS researchers, https://cnrs.hal.science/), ensuring that the publications remain accessible beyond the project lifetime. SupraLife supports Gold OA publishing, allocating a dedicated budget to each partner to ensure that peer-reviewed scientific publications can be immediately provided in OA mode by the scientific publisher.

Joint R&D project/grant applications: The scientific data, project outcomes and exchanged knowledge and ideas will be utilized as a foundation to formulate new hypotheses and enbale the submission of new and innovative joint research project applications with the consortium partners and external world-leading academic and industrial players. However, it is important to note that,

where applicable, data will be kept confidential (information about patentable innovative biomaterials/products/devices) and can only be used within the consortium or shared with the explicit permission of the data owners in compliance with the Consortium Agreement.

5. ORIGIN AND DATA VOLUME

The collaborative efforts of the consortium members: University of Aveiro (UAVR), Eindhoven University of Technology (TU/e), University of Bordeaux (UBx) and its affiliated entities Polytechnic Institute of Bordeaux (IPB) and France National Centre for Scientific Research (CNRS), will generate new data.

The exact size of the data cannot be determined in advance due to the inherent variability in the quantity and types of data that will be generated throughout the project. The estimation of the data volume will be assessed and refined during the project lifespan as the nature and extent of data creation will become clearer.

6. UTILITY OF DATA

Besides the consortium beneficiaries (UAVR, TU/e, UBx and its affiliated entities IPB and CNRS), the data and research results with public dissemination level generated during the project activities are intended to benefit the following target groups:

- Internationally leading universities and academic research centers (e.g., in France – Institute of Supramolecular Sciences and Engineering (ISIS) and Institut Charles Sadron at the University of Strasbourg, University of Grenoble, University of Lyon; in The Netherlands – University of Leiden, University of Twente, TU/e, University of Maastricht; in Portugal – University of Porto, University of Lisbon, University of Minho, Institute for Research and Innovation in Health (i3S), international Iberian Nanotechnology Laboratory (INL); in USA – Northwestern University, Harvard University, Wyss Institute, MIT, University of Stanford, University of Colorado Boulder, Tufts University, University of Texas at Austin, CUNY Advanced Science Research Centre; among others);

- UAVR and other R&D centers, as well as large, medium, and small biotechnology industries in Europe Europe (e.g., CELLULARIS, METATISSUE, Biocant – Technology Transfer Association; Hovione, Crioestaminal; SupraPolix, SyMo-Chem, BASF, Roche, Merck KGaA);

- Health-related public and private foundations (e.g., Institut national de la santé et de la recherche médicale (Inserm), Wellcome Trust, Instituto de Salud Carlos III, La 19aixa Foundation, Oswaldo Cruz Foundation (Fiocruz), The Dreyfus Health Foundation, Fundação Calouste Gulbenkian, Levi Strauss Foundation, Project HOPE (The People-to-People Health Foundation, Inc.), The Rockefeller Foundation, Doris Duke Charitable Foundation);

- Regulatory authorities and international agencies/organization (e.g., Infarmed – National Authority of Medicines and Health Products I.P., ERS – Portuguese Health Regulatory Authority, Portuguese Ministry of Health, EMA – European Medicines Agency, EPHA – European Public Health Alliance, WHO – World Health Organisation, European Commission (DG Health and Food Safety, DG Research and Innovation);

- Media (e.g., TV's, radios, newspapers);

- Society: aiming to raise awareness and engage them in outreach activities, as well as maximizing its benefits and impact from the Centre Region to Europe.

These groups will be effectively reached through a variety of dissemination and communication tools, channels, and activities that are highlighted in the WP5 and outlined in the deliverable D5.4, entitled “Communication, Dissemination, and Exploitation Plan (Sustainability Plan)”. These efforts include leveraging the SupraLife website, social media platforms, newsletters, publications, communication materials such as brochures and flyers, as well as organizing several events like conferences, summer school-type activities, workshops, symposia, scientific retreats, and stakeholder meetings.

By employing dissemination, exploitation, and communication strategies (mentioned in Deliverable D.5.4), we will ensure that the project data reaches and resonates with the target groups, fostering engagement, knowledge sharing, and broader awareness of the SupraLife project’s objectives and outcomes.

7. FAIR DATA

7.1. MAKING DATA FINDABLE, INCLUDING PROVISIONS FOR METADATA

Each beneficiary within the SupraLife Consortium is committed to adhering to the FAIR (findable, accessible, interoperable, and reusable) data principles whenever applicable.

Metadata will be generated for datasets to enhance their discoverability and accessibility. OA data sources will be easily discoverable, identifiable, and locatable using standard identification mechanisms.

Peer-reviewed scientific publications, books, e-books, and book chapters will include Digital Object Identifiers (DOI) that are linked to the associated open data.

To maximize data accessibility, the project activity data will be organized using a numbering system based on the activity, project acronym and number, file content name, and date. OA publications will follow standard naming conventions as prescribed by the Editors of scientific journals.

To facilitate data reusability and enhance discoverability, search keywords will be utilized in research publications and information disseminated on the SupraLife project’s website and social media platforms.

Each document produced by the consortium will be clearly identified with version numbers, which will be added by each respective organization within the consortium.

The SupraLife project will develop descriptive, structural, and administrative metadata to provide comprehensive information about the data, ensuring its proper categorization, organization, and management.

7.2. MAKING DATA OPENLY ACCESSIBLE

SupraLife strongly supports early and widely open access practices and systematic sharing for any kind of document, including reports, surveys, book chapters, books, e-books, and peer-reviewed scientific publications towards raising the awareness of society as a whole about the

project results and outcomes, improving access to scientific information, creating a more equitable system of knowledge sharing open to anyone globally, and boosting the benefits of public investment in research funded under the European Union's Horizon Europe Research and Innovation Programme. The project will utilize OpenAIRE to archive publications and associated data, ensuring continued accessibility beyond the project timeframe.

All peer-reviewed scientific publications resulting from the SupraLife's project will be preferably published in OA journals.

Regarding scientific- and management-related data, the following tools will be used:

- OneDrive folders: Data owners and work package and task leaders will access these folders by logging in with their respective usernames and passwords.
- Servers: Data will be accessed from UAVR's servers, as well as those of partner institutions.
- Project website: The website will provide relevant information about project progress, activities, public deliverables, outcomes, and publications.
- Various repositories will be used for archiving publications and data:
    - Open Research Europe (ORE): https://open-research-europe.ec.europa.eu/
    - Zenodo: This repository will be used as open source and open data repository - https://zenodo.org/;
    - RIA: Institutional repository of UAVR – it is an information system that captures and preserves the research outputs of UAVR's scientific community and makes them available over the web, increasing visibility and impact. It ensures open access to the full text version of documents whenever the authors authorize it – RIA repository will feed OpenAIRE – https://ria.ua.pt/;
    - DUNAS (https://dunas.ua.pt/): Institutional research data repository of UAVR. This repository is intended to share, archive, preserve, cite, access, and explore research data produced in the university scientific research activities. The DUNAS repository will be managed by UAVR also for storing common data, focus group sessions, surveys/questionnaires, results, reports, and deliverables.
    - TU/e repository: Institutional repository of TU/e – https://research.tue.nl/;
    - TU/e Institution networked research storage and 4TU.ResearchData – https://data.4tu.nl/;
    - OSKAR Bordeaux: Institutional repository of UBx – https://oskar-bordeaux.fr/;
    - HAL-CNRS: Repository of CNRS researchers – https://cnrs.hal.science/;

Confidential information about patentable innovative biomaterials/products/devices developed during the SupraLife project and with a market projection and business potential will be restrictedly accessible to the SupraLife's consortium beneficiaries – this data will be treated as closed and confidential, and it will be stored in OneDrive folders accessible only to data owners and authorized staff using their usernames and passwords.

The project team acknowledges the importance of data protection and recognizes that the personal data collected will be treated with utmost respect and care. Personal data will be handled in accordance with the General Data Protection Regulation (GDPR) to ensure its security and compliance.

Before collecting personal data, the project will obtain consent from the individuals concerned, and inform them of how their data will be used. The personal data will only be collected and used for communication purposes between the SupraLife consortium and the individuals.

7.3. MAKING DATA INTEROPERABLE

Data generated within the SupraLife project will be made openly accessible, promoting interoperability to facilitate data exchange and reusability. The data will be provided/stored in standard formats such as **text documents (*.doc, *.docx, *.pdf), excel tables (*.xlsx), images (*.jpg, *.png or *.tiff), PowerPoint slides (*.ppt or *.pptx)**, ensuring compatibility with commonly available free software. This approach will enable seamless data exchange among researchers, institutions, organizations, countries, and other stakeholders without any restrictions.

To enhance inter-disciplinary interoperability, standardized and organized controlled vocabularies will be employed for metadata description. These controlled vocabularies will ensure consistent and structured metadata representation across different disciplines, enabling effective data integration and utilization. By adopting such standardized vocabularies, researchers from various fields will be able to understand and better interpret the metadata associated with the data, facilitating cross-disciplinary collaborations and knowledge discovery.

Overall, the SupraLife project aims to promote open access and interoperability of data, empowering researchers to exchange and reuse data effortlessly, while ensuring effective inter-disciplinary collaboration using standardized controlled vocabularies.

8. ALLOCATION OF RESOURCES

Adhering to FAIR principles involves certain costs, including expenses for OA publication, maintenance of the project website, use of repositories, and copyright licensing.

For the SupraLife project, an allocation of 7,500 EUR has already been made by each beneficiary organization specifically for OA publications. The use of Zenodo is free of charge for data sets up to 50 GB. The institutional research data repository of UAVR (DUNAS) is currently in the pilot phase and further information will be updated in the upcoming version of the DMP.

The Project Coordinator (UAVR) will be responsible for the implementation of the DMP within the SupraLife consortium. However, each beneficiary is responsible for implementing the DMP at their respective levels, including data production, quality assessment, uploading, and providing metadata, among other tasks.

9. DATA SECURITY

During the project implementation, each consortium beneficiary will assume the responsibility for the storage of the collected data. The storage and security of the data will adhere to the organizational protocols established by each beneficiary institution.

UAVR entails a specialized department dedicated to informatics, which manages the digital infrastructure of the entire university, including the hosting and housing of websites and servers.

These servers are equipped with uninterrupted power supply (UPS) systems to prevent any potential data loss.

Any personal data collected will be retained exclusively within the SupraLife consortium. Access to the data will be restricted to authorized personnel who possess an active directory with individual user logins and encrypted passwords.

10. ETHICAL ASPECTS

As reported in the Grant Agreement, all non-sensitive research outputs from the SupraLife project will be made freely accessible for verification, reuse, cross-validation and other research purposes (protection under a Creative Commons license). The project will not collect sensitive research data (e.g., clinical data from patients).

We will ensure the respect for ethical principles and the safe handling of information through different approaches such as by:

- Ensuring that scientists involved are aware of the main rules of data protection;
- Ensuring that the collected data, including personal data from participants attending the different events, is not provided to third parties.

The SupraLife project focuses on the training of PhD students, post-doc researchers, and staff/senior researchers by several means: (i) attending the various events that will be organized by the SupraLife's consortium (e.g., joint summer school-type events, "hands-on" workshops, international symposia, scientific retreats, and final international conference); (ii) short-term on-site trainings; (iii) staff exchanges/visits; and via (iv) the execution of an exploratory project focusing on the development of bioinstructive supramolecular multicomponent biomaterials to interface with living systems.

The content and topics of the SupraLife project are not expected to raise any concern from social, political, medical, and financial groups. There are also no environmental risks or harm, of any kind, being expected from the proposed tasks of the exploratory project nor from the "hands-on" workshops. This project predicts short-time on-site training activities of UAVR's PhD students, post-doc researchers and staff to the facilities of the internationally leading partners (TU/e and UBx/IPB/CNRS) to develop their work, which is related to and can benefit from the training provided by the established SupraLife network. The same flow of researchers from TU/e and UBx/IPB/CNRS is also envisioned to come to UAVR. The acquisition of know-how, knowledge and expertise, and proof-of-concept of the developed complex supramolecular multifunctional biomaterials/advanced biomedical devices during the SupraLife timeframe will be carried out using commercially available human mesenchymal stem cells, as well as other non-biological materials, namely simple chemicals and natural-origin polymers that are also fully commercially available and have already been used at UAVR, TU/e or UBx/IPB/CNRS. The SupraLife project will not involve activities or results raising security issues.

The research carried out during SupraLife timeframe does not involve or generate materials, methods, technologies, or knowledge that could be misused for unethical purposes. The facilities are managed under strict quality procedures. Moreover, the SupraLife project will not involve 'EU-classified information' as background or results. Any information of all participants obtained via the "hands-on" workshops, summer school-type events, international symposia, scientific retreats

or final international conference will be strictly processed according to the GDPR 2016/679 (EU) and no records of personal data will be maintained in any analogic or digital format.

Throughout the duration of the SupraLife project, UAVR, and all the members of the consortium, will have dedicated Data Protection Officers who will be actively consulted, involved, and will offer support.

SupraLife's beneficiaries will ensure compliance with all relevant (inter)national legislations and guidelines related with the planned activities, namely the study and handling of human-sourced biological materials (human stem cells) and disposal/recycling of chemical reagents. The project will comply with all local and national safety measures regarding the use of human and animal cells, being performed in specialized facilities that assure environment, health, and safety of directly or indirectly involved personnel. Moreover, the project is in conformity with current legislation and regulations in the countries where the research will be carried out, and the highest ethical standards and guidelines of Horizon Europe established by the European Commission, and international conventions and declarations.

11. CONCLUSIONS

The current DMP is a dynamic document that is anticipated to undergo continual refinement, improvement, and increased precision throughout the project timeframe. This evolution will be based on the project's progress, conducted/planned activities, and evolving requirements.
The SupraLife's DMP will be revised and updated on M19 (D5.3), in line with the project progress and to disclose the status of the project's reflections on data management.

12. DISCLAIMER

This project has received funding from the European Union's Horizon Europe research and innovation programme under grant agreement No. 101079482.
The content of this report reflects the views and opinions of the authors only and does not necessarily reflect those of the European Union or the European Research Executive Agency. Neither the European Union nor the European Research Executive Agency can be held responsible for them or for any use which may be made of the information contained therein.
This document is subject to updates, revisions, and extensions by the SupraLife consortium.

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