

Project Title: Sensing and predictive treatment of frailty and

associated co-morbidities using advanced personalized models and advanced interventions

Contract No: 690140

Instrument: Collaborative Project
Call identifier: H2020-PHC-2014-2015

Topic: PHC-21-2015: Advancing active and healthy ageing

with ICT: Early risk detection and intervention

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Duration: 36 months

Deliverable No: D9.2 Project Quality Plan

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CHANGE HISTORY

Ver.	Date	Status	Author (Beneficiary)	Description
1.0	1/3/2016	draft	Vasiliki Simaki (UoP)	First draft
1.1	5/3/2016	draft	Vasileios Megalooikonomou (UoP)	Several revisions
1.2	10/3/2016	draft	Yiannis Ellul (UoP)	Added Templates
1.3	12/3/2016	draft	Athanasios Benetos (INSERM)	Review and revisions
1.4	14/3/2016	draft	Vasiliki Simaki (UoP), Andreas Kanavos (UoP)	Incorporation of review comments, second draft
1.5	15/3/2016	Draft	Konstantinos Moustakas (CERTH)	Review and revisions
1.6	24/3/2016	Draft	Vasiliki Simaki, Andreas Kanavos (UoP)	Incorporation of review comments, final draft
1.7	26/3/2016	Final	Vasileios Megalooikonomou	Finalizing draft

EXECUTIVE SUMMARY

This report constitutes the Deliverable "D9.2 - Project Quality Plan" of the FrailSafe (Grant agreement No 690140), and presents the quality procedures, criteria for and areas of application.

The current deliverable is connected with "Task 9.1 Project Management" and "Task 9.2 Risk management and contingency planning", and it is expected to be utilised for the efficient quality assurance of all the work done under the FrailSafe project.

The present deliverable reports on the quality plan of the FrailSafe Project. Quality planning is a crucial component for the successful completion of the project objectives. In this direction, the Quality Manager has reviewed all requirements in order to determine the necessary activities that need to be planned. This Plan is prepared early in the project (M3) in order to demonstrate and provide the Consortium with the assurance that:

- The contract requirements and conditions have been reviewed;
- Effective quality planning has taken place;
- The quality system is appropriate.

A brief version of the quality plan of the FrailSafe project has been described also in D9.1, but it consists the main subject of the current deliverable report, and it is presented on the basis of three main pillars, namely:

- the general quality control measures and actions, such as success criteria, corrective and preventative actions, contingency planning and risk management;
- the quality control of deliverables and documentation, including document types, documents naming, and document templates;
- the quality control of the whole project, including the peer-reviewing evaluation of project's deliverables.

The proposed methodology, which is being presented here, is a revision of a similar methodology already used in previous R&D projects while the proposed quality management scheme is flexible and well-defined, thus allowing for robust Project monitoring and handling of any problems that may arise.

DOCUMENT INFORMATION

Responsible author(s)

Beneficiary

Contract Number:	H2020	90140			Acron	ym:	FRAILSAFE		
Full title	Sensing and predictive treatment of frailty and associated ousing advanced personalized models and advanced interven								
Project URL	http://fr	ailsafe-p	oroject.e	<u>u/</u>					
EU Project officer	Mr. Ra	món Sar	nmartín	Sola					
Deliverable number	er:	9.2	Title:	Project Q	uality	/ Plan			
Work package nun	nber:	9	Title:	Managem	ent &	&Ethics			
Date of delivery	Contrac	ctual	1/4/201	16 (M3)		Actual	31/3/20	16	
						10000	Actual 31/3/2016		
Status	Draft □		Final ⊠						
Nature	Report I	X	Demons	strator 🗆		Other □			
Dissemination Level	Public □ Consortium ⊠								
dissemination) required to delivery. It			ectively nes a pro	manage oject's qua	oroje ality į	ct quality f	rom pro	sary information bject planning to s, criteria for and ities.	
Keywords	Quality	procedu	res, qua	lity policy,	qual	lity plan			
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LIST OF ABBREVIATIONS AND ACRONYMS

ASB	The Advisory Stakeholder Board of the Project
DM	Dissemination Manager
DOW	Description of the Work
EAB	The Ethics Advisory Board of the Project
EM	Exploitation Manager
IPR	Intellectual Property Rights
GA	The General Assembly of the Project.
PC	Project Coordinator
PMC	The Project Management Committee of the Project.
PQB	The Project Quality Board
QM	Quality Manager
QP	Quality Plan
SC	Scientific Coordinator
STB	The Scientific and Technological Board
SM	Standarisation Manager
TL	A Task Leader
WP	Work Package
WPL	A Work Package Leader

1. INTRODUCTION

Quality planning and control is an integral part of management planning for the successful realisation of the project. The Quality Plan (QP) describes the actions and measures that will be taken by the Consortium, in order to ensure the high quality level of the project outcomes and its full conformance with its contractual requirements.

In this direction, the PQB has reviewed all requirements in order to determine the necessary activities, and has prepared the QP in order to demonstrate and provide the Consortium with the assurance that:

- a) The contract requirements and conditions have been reviewed,
- b) Effective quality planning has taken place,
- c) The quality system is appropriate.

The QP is applicable to all the project's activities, and strict compliance with it is mandatory for all participants of the FrailSafe project. In order to ensure the relevance of the quality plan, the QM should conduct quality system reviews throughout the duration of the project, and especially when contractual changes occur. The Quality Plan is a controlled document, and amendments to it shall be submitted to the PC for approval. The QM ensures that the quality plan is available to all concerned and that its requirements are met.

This section specifies the activities to be implemented, including their sequence, in order to ensure that the project and its deliverables conform to its requirements. Those responsible for ensuring that the required activities are carried out, and the resources, which are crucial for their successful completion are identified within the subsequent chapters of this document. The Quality Plan includes explanation, necessary to show how quality requirements for activities are met. A list of such activities is given below:

- Management responsibilities
- Quality Plan Review,
- Quality system,
- Document, data control and product identification and traceability
- Inspection and testing.
- Control of non-conforming prototype / deliverable,
- Corrective and preventative action,
- Internal quality audits.

2. QUALITY SYSTEM REVIEW

The Quality system is reviewed within GA meetings and will focus on the prevention of deviations during each task of the project. In such reviews, the following issues will be taken into account:

- Results from project audits;
- Results from internal audits;
- Official project Deliverables;
- Corrective action requests from all the above;
- Preventive actions on all the above;
- Project prototype deficiencies and subsystems/parts problems;
- Adequate training of the staff of the project's partners for the tasks undertaken;

- Adequate resources for the tasks undertaken.
- Complaints
- Supplier problems
- Ethics, Health and Safety Issues
- Introduction of new quality plans

The outcomes from the above shall be discussed at GA meetings, and discussion minutes shall be kept by an appointed PQB member and should include:

- Satisfaction with the audits, corrective actions and the results of complaints
- Dissatisfaction and requirements for further auditing or more corrective actions
- Satisfaction with the corrective actions taken by the relevant partner(s).

3. PROJECT QUALITY BOARD

The Project Quality Board (PQB) is also responsible for checking and evaluating the quality of reports and deliverables, which are expected to have a significant influence on the successful outcome of the project. The PQ board will review the project activities and deliverables in order to address the following issues:

- Requirements of the project;
- Organizational structure of the project;
- Coordination among the partners;
- General measures and actions taken;
- Planning and control;
- Control of the quality of the deliverables;
- Quality control of the project;
- Files and archives;
- List of quality forms to be used.

The main goals of the QP will be the following:

- Provide to all concerned a guide for the actions required by each one involved,
- Exhibit the performance of the project's quality plan in accordance to the contractual requirements,
- Assign internal members of the Project Quality Board to review specific deliverables.

The PQB is responsible for the coordination and supervision of implementing measures for the quality assurance. Also, it is responsible for the project's quality assurance matters. In accordance with the contractual agreements, the project's quality management plan will be prepared, defining organizational structure, flow of the quality system and the quality management procedures to be applied. The Quality Board is chaired by the Project Coordinator. Apart from the PC, it consists of the Scientific Coordinator (SC), the Quality Manager (QM), the Standardisation Manager (SM) and a representative of older persons (chosen by the General Assembly from names suggested by the organisation representing and working with senior citizens in the project). The PQB has an assisting role to the General Assembly. Internal members will be appointed from the PQB for the purpose of reviewing specific deliverables and reports. These are senior researchers of the project partners with extensive expertise on the subject of the specific deliverable, excluding of course its authors.

In addition to the above members other internal members will be appointed from the PQB for the purpose of reviewing specific deliverables and reports. These are senior researchers of the project partners with extensive expertise on the subject of the specific deliverable. Moreover, members of the different forums of the project will be used as reviewers especially for the public deliverables.

It should be mentioned that project members are obliged to conform to the quality plan and that the procedure for its final approval will be given by the consortium. Furthermore, at least two reviewers will be assigned for each project's deliverable. The PQB meetings will mostly take place through communication media rather than in person meetings.

4. DELIVERABLE PEER REVIEW AND EVALUATION CRITERIA

Each deliverable is linked to a Responsible Partner as defined in the projects grant agreement. The deliverable responsible partner decides on the list of contributors (authors) of the deliverable, who typically come from the partners involved in the work reported on the specific deliverable.

The peer reviewers for each deliverable are listed in Appendix 1: Table of Deliverables with Corresponding Reviewers, whereas the peer review report template is provided in Appendix 2: Template for peer review form

4.1. Review Procedures

In reviewing the Deliverables, the following procedure should be generally adhered:

- a) The Deliverable will be reviewed by the PQB as defined for the specific case (consisting of the internal experts, users' representative, the expert of Standards and the Quality Manager). The defined period for the whole review procedure is can be up to two weeks.
- b) In parallel to the PQB, the Coordinator and the Scientific Coordinator of the project will review the Deliverable to judge the degree to which the objectives are met and whether the Deliverable meets in general the standards to be expected.
- c) The responsible author(s) and the respective task leader(s) will send the consolidated peer review form including all comments of all reviewers appointed and the ones coming from the Coordinator and the Scientific Coordinator (if any) integrated, in order to be able to proceed with the relevant amendments.
- d) The responsible author(s) will revise the Deliverable upon the comments included in the consolidated peer review and will send the revised Deliverable together with their answers to the comments received justifying also the degree of their compliance to them (in the context of a specific form) to the instantiated PQB, the Project Coordinator and the Scientific Coordinator, and the responsible task leader.

4.2. Deliverable Evaluation Criteria

Each deliverable and report will be evaluated according to the following criteria:

General criteria:

- Deliverable contents thoroughness
- Innovation level
- Correspondence to project and programme objectives

Specific criteria:

- Relevance
- Response to user needs (if applicable)
- Methodological framework soundness

- · Quality of achievements
- Quality of presentation of achievements

In case of a document: deliverable layout, format, spelling, etc. Project Deliverables should fully comply with the template given in Appendix 3: Deliverable template. All submitted deliverables should abide to the following rules:

- Have a list of abbreviations used within the Deliverable
- Have a table of contents
- Have a list of Figures (including the ones provided within Appendices)
- Have a list of Tables (including the provided within Appendices)
- Start with a one-page Executive Summary
- Include a References section after the Conclusions section
- Include all detailed technical and other information in Appendices

For the convenience of partners the PQB will make available separate Word files of the Templates in question.

4.3. Timeline of Deliverable Review

The *final rating of a deliverable*, will be given by the PQB, in the following scale: *Fully accepted, Accepted with minor comments, Rejected unless modified properly, Rejected.*

Each deliverable will be evaluated according to the following schedule:

Table 1. Evaluation of FrailSafe deliverables

Deadline	Action
> 15 days before	Deliverable responsible partner and deliverable contributors prepare first draft
15 days before	Deliverable responsible partner sends first draft to the peer Reviewers
10 days before	Reviewers send comments to Deliverable responsible partner, the associated PQB members, and the Coordinator
7 days before	The deliverable responsible partner incorporates the Reviewers' comments and sends the revised version, along with a list of actions describing how he addressed the comments, to the associated PQB members and the Coordinator.
4 days before	The associated PQB members control the quality of the deliverable and if needed ask the deliverable responsible partner for further modifications
2 days before	The associated PQB members send their final rating to the Coordinator
Due Date	Deliverable submission

In case the Commission requests a revision of the submitted Deliverable, the internal review will be only repeated if the changes to the Deliverable are significant.

The Deliverable Responsible partner and the authors of a Deliverable are expected to make every possible effort to confront with the quality criteria as well as with the comments of the peer Reviewers, the PQB members, and/or the Coordinator.

4.4. Identification of prototypes and traceability

In the case of prototypes, their functionalities should be clearly presented in the corresponding deliverables according to the specifications.

4.5. Dissemination Events Scheduling and Reporting

All the dissemination activities of the partners that relate to the FrailSafe project should be supervised by the PQB in order to:

- a) Avoid publication of restricted and/or commercial in confidence data
- b) Avoid misunderstandings among the participants and publication of one's work without proper referencing
- c) Guarantee optimal use of dissemination resources of the project
- d) Guarantee proper archiving of all dissemination material

Dissemination events include but are not limited to:

- a) Publications in Scientific, Technical or Commercial Journals
- b) Presentations in Scientific Conferences
- c) Exhibition stands and demos
- d) Participation in non-project workshops, forums and/or events (EC concertation meetings, etc)

The PC and the PQB should be informed about the participation of any Participant in such an event through the completion of the appropriate form from Appendices 4-6. The PQB is responsible to approving or not the participation in such an event, after having received comments from the partners of the Consortium with experience in the related field.

For scientific publications, the following procedure will be followed:

- a) Event Participation Announcement to the whole Consortium through E-mail using the appropriate form.
- b) Written acceptance should be sent to the requesting Participant within 5 working days from receipt from both the PC and the PQB. Else, it is supposed to be positive
- c) The draft paper is then circulated to all project participants before submission for inspection. If within 15 calendar days, no objection is raised by any partner to PQB or the Coordinator, then the publication is allowed.
- d) A special provision is made in case of a submission to a conference publication: since there may be not enough time between the preparation of the final version and the required 15 days for final approval by the rest of the consortium. In this case the final version of the submitted paper should be sent to the consortium at most 1 day after the conference submission deadline. If the publication rises a major and justified objection by any partner, the article should be withdrawn from the conference.

All participants may object to the publication of confidential data or to non-inclusion of their name, if their work is also included. Comments are to be sent to the publishing Partner with

copies to the PC and the PQB. Then the author should restructure the draft paper accordingly. In case of conflict, it is the responsibility of both the PC and the PQB together to make the final decision.

The participation to dissemination activities/events requiring attendance (e.g., conferences, concentration events, workshops, seminars, etc.) is governed by the following rules:

- A partner should make his application as early as possible and not less than four weeks in advance of the event. The application shall be submitted to the PQB.
- The application should be accompanied by a copy of the event program together with a rationale describing the event and explaining the relevance of attendance to the objectives of the Project.
- The application must provide a clear breakdown of the attendance cost explaining the proposed claim for the EC contribution.
- Within two weeks after the event, the partner must provide to PQB a concise written report about the event. If possible, the report should be accompanied by the event's proceedings (or at least a suitable extract of it).

4.6. Internal Quality Audits

The QM will be responsible for the implementation of this procedure. The progress of the project will be monitored by the QM through any possible communication links (mainly by email), involving all the related partners. All day-to-day and trivial barriers of the project have to be dealt in this way. In exceptional cases, when a problem of paramount importance comes up with a certain partner, an Internal Audit Procedure will be carried out by a specific project group. This group will consist of: (a) the PC, (b) the SC and (c) the QM. All the findings of the Internal Audit will be documented by the QM in the Internal Audit Report (see Appendix 7). Then, the QM will issue corrective actions, which again will be documented in order to make all the discrepancies obsolete, within the appropriate time period. Follow up actions will be arranged, so as to ensure the effectiveness of the corrective actions. The results of the Internal Quality Audits will be distributed to all partners, related to a specific WP.

4.7. Project Reporting and Monitoring

Concerning Quality Management Scheme, Periodic Management Reports (PMRs) that have been included in the DoW will be delivered on a 6-month-basis, and in which all partners will contribute by reporting their technical and financial progress in the project. Every partner should provide the following information for the particular 6-monthly reporting period (Periodic Management Report):

- Per each Work Package and respective Tasks:
 - Objectives of the WP/Task within the reporting period
 - Status and progress towards the WP/Task Objectives
 - Deviations from the DoW and Corrective Actions
 - Plan for the next reporting period (regarding the WP/Task)
- Dissemination & Exploitation activities for the reporting and the next reporting period
- Project Meetings
- Cooperation with other partners
- Cooperation with other projects
- Deviations from planned deliverables and milestones
- Planned and actual person months per WP and Task.

The PMR will be used to depict deviations from planned milestones including delays or early finishes and their implications on the overall progress of the project. The corrective actions that

are necessary for implementation will be considered and communicated to the SC so that they are carried out in a timely manner in order to achieve the expected results. Alarms will be raised in case of significant deviations in the planned work and budget spending. A Template of the PMR is presented in Appendix 8 so as to facilitate the efficient and easy integration of the feedback from all the partners by the PC.

4.8. Corrective and preventive actions

The issues described in this section relate to general performance of a Partner and the quality of his work outcome and not to Project Deliverables, for which the previously described procedures should be followed. Any participant may raise such an issue on the work of another participant or external suppliers work.

The PC and the QM are responsible for resolving such matters within their own areas of responsibility. All complaints are to be investigated and corrective action to be agreed. Corrective and possible preventative actions are recorded and all involved are informed of the action taken (using the template forms in Appendices 7, 9, 10).

The procedure in such cases will be as follows:

- a) The PC identifies the need for corrective actions (e.g. by proposals from partners).
- b) The PC notifies the WP leader.
- c) WP leader discusses the issue with the Task leader and comes up with a proposed solution. The relevant request is documented on the appropriate form and a proposal on corrective action is being done.
- d) The solution is forwarded to the PC and the SC via the WP leader.
- e) The PC and the SC decide on the matter. The decision shall be documented according to the template of Appendix 3. The PC sends the decision to all involved parties and checks that the actions are implemented.

5. CONTROL OF REPORTING AND DOCUMENTATION

The QM is responsible for ensuring all documents are controlled effectively and that they follow the format scheme that is described in the following sections.

5.1. Document Types

The types of documents produced within FrailSafe are as indicated in the table below:

Document Document Description Type **Type** Code Deliverable D Describes the work done within a WP and/or task. Describe work done towards a specific subtask of Internal Report **IR** goal of the project for use by the consortium. Technical A scientific paper submitted for publication, or is in TR Report press, or has already published, and which is

Table 2. Document Types

		uploaded in the FrailSafe web site.
Dissemination Report DR		Describe the foreseen dissemination activities before publication in order to inform the consortium and get feedback. Dissemination activities include, (but are not limited to) magazine/journal publications, demonstration in exhibitions and conference publications, leaflets and brochures.
Review Report	RR	Filled in by a deliverable reviewer as a measure to evaluate the quality of the work done.
Meeting Program	MP	Used to communicate the schedule of a project's event or meeting. In many cases, it can be just an email to the project mailing list.
Meeting Agenda	MA	Used to communicate the purpose and items to be discussed in a physical or virtual meeting. In many cases, it can be just an e-mail to the project mailing list.
Meeting Minutes MM		Summarizes the topics dealt during the meeting as well as the actions agreed.
Conference Call Minutes	СМ	Summarizes the topics dealt during a conference call as well as the actions agreed. In many cases, it can be just an e-mail to the project mailing list.
Presentation	Р	Used to expound topics related to the project, both internally (Consortium meetings, a partner's vision/contribution, etc) and externally (conferences, dissemination events, meetings, annual review meetings, etc).
Financial Report	FR	Filled in by the partners to state their costs.
Periodic Management Report	PMR	Filled in by the partners to report on status and progress towards project objectives as well as on managerial issues, cost statements and justifications, as well as on planned and actual manpower spent within a certain reporting period.
Other	ОТН	Any other of document category that does not fall in the types described above

5.2. File Formats

The format of a file depends on its type and use, which can be either *internal* to the consortium or *external* (including the EC). The following table shows the allowable format of documents per type and use, where "doc" stands for MS-Word or equivalent compliant format, "txt" stands for plain text format, "ppt" stands for MS Power-Point presentation or equivalent compliant format, "xls" stands for MS-Excel or equivalent compliant format, and "PDF" stands for the portable document format.

Table 3. Document formats and templates

Document Type	Allowable Format for Internal Use	Allowable Format for External Use	
Deliverable	doc	PDF	
Internal Report	doc	PDF	
Technical Report	doc	PDF	
Dissemination Report	doc	PDF	
Review Report	doc	PDF	
Meeting Program	doc, txt	PDF	
Meeting Agenda	doc, txt	PDF	
Meeting minutes	doc	PDF	
Conference Call minutes	doc, txt	PDF	
Presentation	ppt	PDF	
Financial Report	doc, xls	PDF	
Periodic Management Report	doc	PDF	

5.3. Document Naming and Coding

For facilitating common browsing and storage in different platforms and OS's, no spaces should be used in the document names, and instead the dash character "-" should be used.

All project document names must start with the prefix "FrailSafe-" in order to facilitate quick identification and indexing. In particular, the following conventions are mandatory for certain types of documents. Names of deliverable documents should follow the convention:

"FrailSafe-Deliverable-Dw.n[.m]-[ShortTitle]-vX.Y.ext"				
"Dw.n[.m]"	is the deliverable number			
"W"	is the WP number			
"n"	is the numbering of the deliverable within the specific WP			
"[.m]"	(if exists) is the sub-numbering within the specific WP			
[ShortTitle]	is an indicative short title that characterises the deliverable			

"vX.Y"	is the version number
"X"	is the version
"Y"	is the sub-version
"ext"	is the file extension pertaining to the format used

For instance, the name of (the final version of) deliverable D9.2 sent to the EC can be "FrailSafe-Deliverable-D9.2-Project_ quality_plan-v1.7.pdf"

The name of remaining *FrailSafe* documents will follow the convention:

"FrailSafe-XX-ddd.pdf"

where "XX" is document type code as defined in Table 4 the "ddd" is a three-digit decimal number that will be assigned automatically to a new Technical Report by the related submission service of the FrailSafe- web site.

5.4. File Naming

There will be a unique coding system for Project documents, as indicated below:

Document Code	Document Type	Type of Forms to be used		
D	Deliverable	Deliverable Template, Peer Review Report, Summary of main Feedback and Actions taken, Non-Conforming Report, Corrective Actions Request		
TP	Technical Paper	Internal Report Template		
PR	Work Packages Plans and Progress Reports	Internal Report Template		
М	Minutes, Action Lists, Decision Lists	Internal Report Template		
ОТН	Other subjects	Internal Report Template		

Table 4. Description of Document Codes for File Naming

5.5. Document referencing and templates

There is a unique document referencing scheme which is not applicable for informal data and views exchange between Partners. The document referencing scheme is only valid for official Consortium documents, falling in one of the above categories of the previous section. Still, if a Participant selects not to classify one of his communications, he may not raise claims later, if another participant has not considered it.

Project Deliverables should fully comply with the templates given in appendices at the end of this document. This includes the guidance for the layouts of the Cover Page, headers, footers, etc. For the convenience of partners the QM will make available separate Word files of the Templates in question.

Furthermore, they should abide to the following rules:

- Have a list of abbreviations used within the Deliverable
- Have a table of contents
- Have a list of Figures (including the ones provided within Appendices)
- Have a list of Tables (including the provided within Appendices)
- Start with an one-page Executive Summary
- Include a References section after the Conclusions section
- Include all detailed technical and other information in Appendices

APPENDIX 1: TABLE OF DELIVERABLES WITH CORRESPONDING REVIEWERS

Table 5. FrailSafe Deliverables and Reviewer assignments

Del. №	Deliverable name	Lead	Туре	Due Date	Reviewer	
Mō		Beneficiary			1	2
1.1	Analysis of current practices	CERTH	R	M6	INSERM	UoP
1.2	User requirements, use cases, UCD methodology and final protocols of evaluation studies	CERTH	R	M12	INSERM	UoP
1.3	FrailSafe technical specifications and end-to-end architecture	CERTH	R	M12,M24	MATERIA	SIGLA
2.1	Clinical study methodology	INSERM	R	M6	HYPERTECH	SMARTEX
2.2	Clinical guidelines formalized	MATERIA	OTHER	M18,M27	INSERM	UoP
2.3	Completion of quantification campaign	INSERM	OTHER	M18,M25	MATERIA	UoP
2.4	Behavioural Monitoring	INSERM	OTHER	M18	UoP	MATERIA
3.1	Definition of sensor components and communication strategy	SMARTEX	R	M6	HYPERTECH	BRA
3.2	Preliminary WWBS prototype	SMARTEX	R/ DEM	M15	SIGLA	CERTH
3.3	Final WWBS prototype	SMARTEX	R/ DEM	M24	CERTH	UoP
4.1	Offline analysis of data	UoP	R/ DEM	M18,M24	CERTH	SMARTEX
4.2	Online analysis of data	UoP	R/DEM	M18,M24	CERTH	SMARTEX
4.3	Dynamic User Profiling models and Patient modelling and representation framework	CERTH	R	M6,M12	UoP	SIGLA
4.4a	Linguistic Corpus	UoP	OTHER	M18	HYPERTECH	SIGLA

Del. №	Deliverable name	Lead	Туре	Due Date	Reviewer		
Mō		Beneficiary			1	2	
4.4b	LingTester Test Results – Active (on-line) mode	UoP	R	M18,M24	CERTH	SMARTEX	
4.5a	LingTester (Prototype)	UoP	DEM	M12,M24	SIGLA	HYPERTECH	
4.5b	LingTester Test Results – Passive (off-line) mode	UoP	R	M18,M24	SIGLA	HYPERTECH	
4.6	Signal processing algorithms for extraction of frailty related indicators	UoP	R/DEM	M12,M24	SIGLA	HYPERTECH	
4.7	FrailSafe Decision Support System	UoP	R/DEM	M24, M28	CERTH	SMARTEX	
5.1	Analysis of hardware devices and software tools. Game hardware and software design.	BRA	R	M6	HYPERTECH	CERTH	
5.2	Beta version of the Synthesized AR game system.	BRA	DEM	M16	UoP	HYPERTECH	
5.3	Final Synthesized AR game system	BRA	DEM	M24	CERTH	SIGLA	
5.4	Personalised context- aware,Information Visualization	CERTH	R/DEM	M24, M28	UoP	BRA	
6.1	FrailSafe Virtual Community Platform	UoP	R/OTHER	M28,M32	BRA	SIGLA	
6.2	FrailSafe mHealth Integrated version	SIGLA	R/DEM	M18,M25, M32	HYPERTECH	BRA	
7.1	Assessment protocol	MATERIA	R	M20,M26	HYPERTECH	BRA	
7.2	Small-scale evaluation report	INSERM	R	M22	UoP	MATERIA	
7.3	Field trials report & Socio- economic guidelines	MATERIA	R	M36	INSERM	UoP	
8.1	Dissemination Plan and FrailSafe dissemination material	AGE	R	M3, M12, M24, M36	HYPERTECH	SMARTEX	
8.2	Project Web Presence	HYPERTECH	DEC	M3	SIGLA	BRA	
8.3	Exploitation Report and FrailSafe Business models	HYPERTECH	R	M24,M36	UoP	CERTH	

Del. №	Deliverable name	Lead	Туре	Due Date	Reviewer		
ΝY		Beneficiary			1	2	
8.4	IPR Protection Plan	HYPERTECH	R	M12,M24	SIGLA	BRA	
8.5	Standardisation and concertation activities report	SIGLA	R	M24,M36	HYPERTECH	SMARTEX	
8.6	Data Management Plan	HYPERTECH	R/DEC	M6,M24 M36	SIGLA	UoP	
9.1	Project reference manual and quality plan	UoP	R	M3	CERTH	INSERM	
9.2	Project Quality Plan	UoP	R	М3	INSERM	CERTH	
9.3	Periodic Management Reports	UoP	R	M6,M12, M24,M30	/ \(\) / \(\)		
9.4	Project First Report	UoP	R	M18	All	All	
9.5	Project Final Report	UoP	R	M36	All	All	
9.6	Ethics, Safety and mHealth Barriers (regulation, legislation, etc.) Manual	UoP	R	M5,M36	INSERM	MATERIA	

APPENDIX 2: TEMPLATE FOR PEER REVIEW FORM



Project:

Sensing and predictive treatment of frailty and associated co-morbidities using advanced personalized models and advanced interventions (FallBate, Grant Agreement No: 690140)



Peer Review Consolidated Report								
Deliverable	DX.X Deliverable name							
WP, No, title, activity type	Beneficiary short name							
Dissemination level	WPX - WP Title							
Lead beneficiary	Main task(s) of relevance							
Quality Manager	Type of dissemination level: PU (public) – CO (confidential)							
Review year	Month/Year							
Review result	FD (final draft) - WD (working draft)							
File name and size	"File name", File8ize kb							

FRAIL8AFE - H2020-PH0-690140 Peer Review DX.X

INTRODUCTION

The PrailBafe Consortium uses the Peer Review process for its internal quality assurance for deliverables to assure consistency and high standard for document project results.

The Peer Review is processed individually by selected reviewers. The allocated time for the review is about two weeks. The leading author of the corresponding Deliverable has the final responsibility to collect the comments and suggestions from the Peer Reviewers and decide what changes to the document and actions are to be undertaken.

Quality Manager

Surname/ Name	Beneficiary Name (short name)	Contact e-mail
Ellul loannis	UoP	ellul@upatras.gr

Peer Review List

=	Surname/ Name	Beneficiary Name (short name)	Contact e-mail
1			
2			
3			

Overall rating of the Deliverable

Excellent	Very good	Good	Poor	Very poor
		120		

FRAILSAFE - H2020-PH0-690140 Peer Review DX.X

1. 8UMMARY OF 8UGGESTED ACTIONS

The review process concluded that...

<Page break>

2. COMMENTS BY REVIEWER #1

	General comments
	Specific comments
Relevance	Reviewer's comment:
Relevance	Author's response:
Response to	Reviewer's comment:
userneeds	Author's response:
Methodological framework	Reviewer's comment:
soundness	Author's response:
Quality of	Reviewer's comment:
achievement	Author's response:
Quality of	Reviewer's comment:
presentation	Author's response:
Deliverable layout /	Reviewer's comment:
Spelling / Format	Author's response:

-3-

APPENDIX 3: DELIVERABLE TEMPLATE



Project Title: Sensing and predictive treatment of frailty and

associated co-morbidities using advanced personalized models and advanced interventions

Contract No: 690140

Instrument: Collaborative Project
Call Identifier: H2020-PHC-2014-2015

Topic: PHC-21-2015: Advancing active and healthy ageing

with ICT: Early risk detection and Intervention

Start of project: 1 January 2016
Duration: 36 months

Deliverable No: DX.X

Deliverable Title

Due date of deliverable: MXX (XX* Month 201X)

Actual submission date: XX* Month 201X

Version: XX

Date: XX* Month 201X



Horizon 2020 European Union funding for Research & Innovation

Change History

•					
	Ver.	Date	Statue	Author (Beneficiary)	Description
	x.x	XX/XXXX	draft	Author name (ORGANIZATION ACRONYM)	ToC

<PAGE BREAK>

EXECUTIVE SUMMARY

The deliverable DX,X, reports on XXXX.

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DOCUMENT INFORMATION

Contract Number	: H2020	PHO-	-690140 Aoronym: FRAILSAFE						
Full title		sensing and predictive treatment of failty and associated co-morbidities ising advanced personalized models and advanced interventions							
Project URL									
EU Project office	r Mr. Ra	smón 8	anmerti	n Sola					
Deliverable numb	er:	x.x	Title:	xxxx					
Work package nu	ımber:	x	Title:	xxxx					
Date of delivery	Contra	otual	XX/X	X/201X (MXX)	Aotual	твс			
8 tatus	Draft E				Final D				
Nature	Report	B	Demor	strator 🗆	Other 🗆				
Dissemination Public ☑ Consortium □				rtium 🗆					
Abstract (for	This de	llevera	ble repo	rts on XXX					

Keywords XXXX

Contributing authors	XXXXX (ORGANIZATION ACRONYM) XXXXX (ORGANIZATION ACRONYM)					
(beneficiaries)						
Responsible	XX	xx	Emall	xxxxxxxxx		
author(s)	Beneficiary	ACRONYM	Phone	XXXXXXXXXXX		
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2.1.1.1 XXXXXXX

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LIST OF TABLES

2 XXX

2.1 XXXX...... 2.1.1 XXXXX.....

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LIST OF FIGURES

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FRAIL 8AFE - H2020-PHC-890140 DX.X XXXXXXXXXXX

1 INTRODUCTION

XXX

2 XXX 2.1 XXXX XXXX

2.1.1 XXXXX

2.1.1.1 XXXXXX

Figure 200, 2000000

XX

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REFERENCE 8
[1] XXXXXX

[2]

-1-

APPENDIX 4: TEMPLATE FOR REPORTING NEW MAGAZINE PUBLICATIONS

				30140 Magazine Publication	MM/YY		FRAILSAÆ -H2020-PH	0-690140 Maj	jazine Publicatio	1		MM/YY
100	Horizon 2020		1. TITLE			1	6. MAGAZINE INFORM	ou				
	European Union funding for Research & Innovation]	Name of magazine	Full magazine t	(Ha			
	for Research & Innovation		2. AUTHORS LIST				Acronym	ruii magaziie i	iue			
			Z. AUTHORS LIST	Leading author (editor)		1	Publisher					
				Beneficiary Name		1	Editor	Sumame of edi				
			Sumame/ Name	(short name)	Contact e-mail		Magazine URL	Sumame or equ	tor			
Proje						1						
Sensing and predictive treatment of fr				Co-author (in alphabetic orde	ir)	1	Туре	El Solentifio	Professional	Special .	Informative	Newspap
using advanced personalized mode	els and advanced interventi	ions	Sumame/ Name	Beneficiary Name	Contact e-mail	1	Range	International	E Europe	America	National: (ple	
(FrailSafe, Grant Agree	ment No: 690140)		Sumame/ Name	(short name)	Contact e-mail		General policy	Traditional pub	ishing policy	₽ Op	en access pol	loy
						1	Author's cost					
]	Reader's cost					
Form for publication Title of Magazi Lead beneficiary! Authors Summary Related WP no, title and activity type	n in Magazine		OTHER INFO ABOUT Number of pages or words	(eg. "10 pages, single-space" or "			e. 8PECIAL REQUES' 8 peolal requestoloom Manager or Quality M	ments to be com			coordinator, 8	olentifio
Contributing task(s)	Main task(s) of relevance		Language of paper Full reference (if available)	E English	Other: (please specify)	-						
Submission date	DDMM/YY		Participance (it available)			J						
Publication date	Month/Year											
Publication status	Published, Accepted, Pended											
-11-				-2-					-3-			

APPENDIX 5: TEMPLATE FOR REPORTING NEW DEMONSTRATION EXHIBITION



FRAIL8AFE = H2020-PH0-690140 Exhibition / Event MM/YY

8. MAGAZINE INFORMATION

Name of magazine	Full magazine	title						
Acronym								
Publisher								
Editor	Sumame of ed	Sumame of editor						
Magazine URL								
Туре	₽ 8olentifio	Professional	Special .	Informative	Newspaper			
Range	International	E Europe	America	National: (ple	ase specify)			
General policy	Traditional pub	Traditional publishing policy El Open access policy						
Author's cost								
Reader's cost								

7. SPECIAL REQUESTS/COMMENTS

8pecial requests/comments	to be communicated	to the Project Coordinator,	8olentiflo
Manager or Quality Manager	prior or after the pu	blication.	

-3-

-2-

MM/YY

APPENDIX 6: TEMPLATE FOR REPORTING NEW CONFERENCE PUBLICATION



Project:

Sensing and predictive treatment of frailty and associated co-morbidities using advanced personalized models and advanced interventions (FallSafe, Grant Agreement No: 690140)



Form for presentation in Conference

Name of Conference and Abbreviation Publication			
Lead beneficiary/ Corresponding persons	Beneficiary Short name/ Author		
Summary	No more than 10 lines		
Related WP no. title and activity type	Main WP(s) of relevance		
related WF 110, one and activity type	WPX - WP Title		
Contributing task(s)	Main task(s) of relevance		
Submission date	DD/MM/YY		
Presentation date	Month/Year		
Presentation status	Presented, Accepted, Pended		

-1-

TYPE				
Oral presentation	Paper	Poster	Tutorial	Other: (please notify)
E				

FRAILSAFE - H2020-PHO-690140 Magazine Publication

Leading speaker (main author)					
Sumame/ Name	Beneficiary Name (short name)	Contact e-mail			
	Co-authors (in alphabetic order)				
Sumame/ Name	Beneficiary Name (short name)	Contact e-mail			

6. OTHER INFO ABOUT THE ARTICLE

Number of pages or words	(eg. "10 pages, single-space" or "6000 words")		
Language of paper	El English	Other: (please specify)	
Full reference (if available)			

8. CONFERENCE INFORMATION

FRAILSAFE - H2020-PHO-690140 Magazine Publication

. Voti ananya ini winin iron							
Name of conference	Full conference title						
Acronym							
Date and location							
Conference URL							
Туре	El Bolentifio Professional Special Informative New				Newspaper		
Range	International El Europe		America Nationa		National: ((please specify)	
Review policy	■ Abstract	Abstract Extended		Full Other: (p		ease notify)	
	review	abstract	рар	er			
		review	revi	ew			
Conference	■ Hardoopy	Hardcopy ar	and CI			Oth	er. (please
proceedings		CD				not	fy)
Speaker's cost		_					
Participant's cost							

MM/YY

7. SPECIAL REQUESTS/COMMENTS

Special requests/comments to be communicated to the Project Coordinator, Scientific Manager or
Quality Manager prior or after the publication.

8. POST REPORTING

Report on the participants' feedback after the conference to be communicated to the Project	
Coordinator, Scientific Manager or Quality Manager.	

4.

APPENDIX 7: TEMPLATE FOR INTERNAL AUDIT REPORT



Project:

Sensing and predictive treatment of frailty and associated co-morbidities using advanced personalized models and advanced interventions (Frail@afe, Grant Agreement No: 650140)



Internal Audit Deficiency Report No. (please notity the number)			
Title A small description of the report			
Audited Partners	Name (Partner)		
Summary	(no more than 5 lines)		
Date			

AUDITED PARTNER:	REF. No:
REFERENCE:	

PART 1: DEFICIENCY DETAILS	
Inspected by:	
Partner Representative:	
Details:	
PART 2: CAUSE	
Details:	
PART 3: CORRECTIVE ACTION	
Details:	
PART 4: FOLLOW UP AND CLOSE OUT	
Details:	

APPENDIX 8: TEMPLATE FOR PARTNER PERIODIC MANAGEMENT REPORT

Horizon 2020 European Union funding for Research & Innovation
Tor Research & Innovation

PROJECT:

SENSING AND PREDICTIVE TREATMENT OF FRAILTY AND ASSOCIATED CO-MORBIDITIES USING ADVANCED PERSONALIZED MODELS AND ADVANCED INTERVENTIONS

(FrailSafe, Grant Agreement No: 690140)



Frail 8afe Periodio Management Control (Each partner chould send the concollidate input to UoP)			
Partner			
Reporting period			
Date			

PRAILSAPE - H2020-PHC-690140	Management Report	Da

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1.1	 1.1 Objectives for the Reporting Period of all active WP and T. 	
1.1	1.2 Status and Progress towards objectives of all active WP a	nd T3
1.1	1.3 Efforts for the Next Period of all active WP and T	
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3.2	Explanation of the use of resources	

- 1 WORK PROGRESS AND ACHIEVEMENTS IN THE REPORTING PERIOD
- 1.1 Description of Work and Future Steps
- 1.1.1 Objectives for the Reporting Period of all active WP and T

WP1	The objectives of WP1 in this reporting period were to
T1.1- State-of-the-art assessment and acquisiton of methodological tods	_

1.1.2 Status and Progress towards objectives of all active WP and T

T1.1- State-of-the-art assessment and acquisiton of methodological tods	m e
<u></u>	

1.1.3 Efforts for the Next Period of all active WP and T

T1.1- State-of-the-art assessment and acquisition of methodological tods	
<u></u>	

1.1.4 Deviations from QoA and Corrective Actions

T1.1- State-of-the-art assessment and acquisition of methodological tods		

Date

6

PRAILSAPE - H2020-PHC-690240 Management Report Date 1.2 Deliverables for the Reporting Period FRAILSAFE - H2020-FHC-690240 Management Report Date Del. Date PRAILSAPE - H2020-PHC-690040 Management Report No. Level CERTH MATERIA SIGLA Foreseen HYPERTECH INSERM 1.3 Milestones for the reporting period 1.8 Cooperation with other Projects (all partners) Lead Comments Del. Date 2.3 Concentration activities and Synergies No No. Benef. 1.7 Deviations from Planned Deliverables and Milestones 2.4 Other dissemination activities 1.3 Development and Maintenance of the Frail 3afe Project web site 3 USE OF RESOURCES 1.4 Highlights / anticipated problems for next reporting period 3.1 Summary of planned and actual PMs Allocation for the reporting period 2 DISSEMINATION/EXPLOITATION INFORMATION The following table shows the person months planned and actually spent for this reporting 2.1 List of Publications period. The numbers in the "planned" row typically reflect the average distribution of the resources over the lifetime of a WP. The numbers in the "actual" rows are the actually sent PM's over the 12-month reporting period. 1.5 Cooperation within the Project All project partners have been participated in the kick-off meetings and anticipated a fruitful and Title of the Number, date Publisher Year of Pages Permanent Open Intense cooperation towards next effort foreseen in the project. periodical or the Partner WP2 WP3 WP4 WP5 WP6 WP7 series During the project kick-off meetings, several actions to be undertaken by project partners were name set and a list of actions were prepared and distributed by UoR to the Consortium. Attended Furthermore, several internal meetings, and teleconferences have been performed by WPL and the PC and the necessary tasks were assigned per each partner towards fulfilling the objectives of the reporting period. Specific cooperation among partners is as follows: 3.2 Explanation of the use of resources Foreseen Partner Cooperation within the project UoR. BRA 2.2 List of Dissemination Activities **SMARTEX** No Type of Partners Date Place Type of Size of Countries AGE activities involved audience audience addressed Attended

APPENDIX 9: TEMPLATE FOR NON-CONFORMITY REPORT

Horizon 2020
European Union funding
for Research & Innovation

PROJECT:

SENSING AND PREDICTIVE TREATMENT OF FRAILTY AND ASSOCIATED CO-MORBIDITIES USING ADVANCED PERSONALIZED MODELS AND ADVANCED INTERVENTIONS

(FrailSafe, Grant Agreement No: 690140)



Non-Conformity report			
Title Small description of the report			
Partner	Name		
Summary	No more than 5 lines		
Status D (draft)/ F (final)			

FRAIL 8AFE - H2020-PHC-890140 Non_-Conformity Report

Date

	Non-Conformity report for partner:				
Item	Item Participant			Date	
WP No	WP No 1			Date	
NCR		Issue			
NO	Deviation		Disposition		

APPENDIX 10: TEMPLATE FOR CORRECTIVE ACTION REQUEST



PROJECT:

SENSING AND PREDICTIVE TREATMENT OF FRAILTY AND ASSOCIATED CO-MORBIDITIES USING ADVANCED FERSONALIZED MODELS AND ADVANCED INTERVENTIONS

(FrailSafe, Grant Agreement No: 690140)



Crreotive Action Request No (please notify number)			
Title Small description of the report			
Requesting Participant	Name		
Summary	No more than 5 lines		
Date			

FRAIL 8AFE - H2020-PHC-880140 Corrective action request Date

Item		Participant			Date	
WP N	lo	T No		Date		
NCR		Issue				
NO	Item and De	eviation	Decision	Responsibl	e Partner	Expected date for correction

2