

Interregional Coordination for a fast and deep uptake of Personalised Health - Regions4PerMed

Report

KEY AREA 4: INNOVATION FLOW IN THE HEALTHCARE

GRANT AGREEMENT NUMBER	825812
PROJECT FULL TITLE	Interregional coordination for a fast and deep uptake of personalized health
PROJECT ACRONYM	Regions4PerMed
FUNDING SCHEME	CSA
START OF THE PROJECT	01/11/2018
DURATION	54 Months
CALL IDENTIFIER	H2020-SC1-2018-Single-Stage-RTD
PROJECT WEBSITE	http://www.regions4permed.eu/



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1. ELIXIR AS A MODEL OF RESEARCH INFRASTRUCTURE

Pro	ject Initiative title	ELIXIR
Organization name		ELIXIR
Country		United Kingdom
	gion	East-Cambridgeshire
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	bsite	https://elixir-europe.org/about-us/who-we-are/hub
	/words	Genome, bioinformatics, interoperable standards,
,	,	infrastructure for biological data, B1MG and European Joint
		program of rare diseases
Dui	ration	Not limited
	a of application	Bioinformatics research infrastructure
	Main challenges tackled	Huge amounts of data have been produced during the last years, most
	3	research centers are not able to store or transfer this data or lack the
		expertise to analyze this data. ELIXIR provides a sustainable and
		interoperable federated infrastructure for the management of sensitive
		human data at regional, national and international level. Thus,
		researchers can use ELIXIR facilities to easily access, find, store, transfer
		and analyze the datasets that match with their needs.
	Objectives	
		ELIXIR main mission is to construct and operate in a single infrastructure
		for human genomics and translational data in Europe by a federated
		system. ELIXIR consists of ELIXIR Hub that coordinates the work across
		ELIXIR and ELIXIR Nodes which coordinates the work locally.
		·
	Main concept and	ELIXIR is an intergovernmental organization that coordinates and
	methodologies involved	develops life science resources for researchers across Europe. Through
		ELIXIR, researchers can access, store, transfer and analyze data by
		"compute platform" and "tools platform". ELIXIR also provides
		researchers standards by "interoperability platform" and linkages between
		data and literature by "data platform".
	Impacts (health,	The main impacts of ELIXIR are related to different initiatives in genomic
	scientific, industrial,	data resources:
	socio-economic or others enabled by the	• ELIXIR is part of the strategic partnership Global Alliance for Genomics & Health (GA ₄ GH), to simplify searching for and request access to
	project/initiative	potentially identifiable data in regional, national and international genomic
	project/initiative	data resources.
		ELIXIR is the coordinator of Beyond One Million Genomes project
		(B1MG): Through its project, the ELSI toolkit was developed, which is a
		federated secure cross-border technical infrastructure and a B1MG
		maturity model for Personalised Medicine.
		By-COVID project is a € 12 million Horizon Europe project for the
		identification of the data challenges to overcome for an effective pandemic
		response.
		ELIXIR developed different programs to support researchers such as
		compute platform, data platform, interoperability platform, training
		platform or tools platform.
	Funding and	The ELIXIR Hub and nodes compete together for grant funding under
	Investments (please	Horizon 2020 and the Innovative Medicine Initiative (IMI). Additionally,
Z	specify the source:	the ELIXIR Hub is funded through membership fees.
Ö	public, private,	
[PT	Structural or other	
8	types of funds)	
DESCRIPTION	Key stakeholders	Researchers, computer scientists, industry
	involved	



2. FRAUNHOFER IZI AS APPLIED RESEARCH ORGANIZATION

Project Initiative title		Fraunhofer Institute for Cell Therapy and Immunology (IZI)
Organisation name		The Fraunhofer-Gesellschaft (FhG)
Country		Germany
Region		Free State of Saxony
	ct person	Thomas Tradler
	ct email	thomas.tradler@izi.fraunhofer.de
Websi		https://www.izi.fraunhofer.de/en.html
Keywo	ords:	Research infrastructure, Cell and Gene Therapies, Clinical
		trials, Applied research
Durati		Not limited
Area c	f application	Cell and Gene Therapy Research & Development Value Chain
	Main challenges tackled	To conduct Research and Development (R&D) activities in the field of biotechnology, pharmaceuticals, and medical engineering via research contracts for companies, hospitals, diagnostic laboratories and research institutes.
	Objectives	The Fraunhofer IZI investigates, develops, optimizes, and validates methods and solutions in four business fields: cell and gene therapy, drugs and vaccines for human and veterinary medicine, molecular diagnostics and immunodiagnostics focuses on the discovery and clinical validation of DNA and RNA as well as extracorporeal therapies.
	Main concept and methodologies involved	The Fraunhofer IZI is a professional service provider in the field of R&D. It performs research and capabilities as well as conducting and pursuing internal and joint (R&D) project in close collaboration with industry and academia. The cell and gene therapy strategy involves the entire value chain from early development through Good Manufacturing Practices (development and manufacturing) to clinical trials. The strategy of facilitating the innovation flow starts with applied research activities jointly with industry and academic partners to promote knowledge and innovation. The second step is the excellence research partnerships with international partners to develop innovative technologies and products and the part three of the strategy is the creation of spin-off companies and the support for establishing subsidiaries of international companies in Germany.
	Impacts (health, scientific, industrial, socio-economic or others enabled by the project/initiative (max 200 words)	Thanks to the strategy of fostering the innovation flow, joint research and contract research projects (with companies such as Novartis, Daiichi-Sankyo); international research partnerships (with, i.e., McMaster University, Monash University); the creation of spin-off companies (Ribonomix GmbH, Bioville, or Epitopic); gain expertise and knowledge by supporting the establishment of international companies in Saxony (Bellaseno or Apocell) or formation of cluster for regional development (SaxoCell®) were possible.
DESCRIPTION	Funding and Investments (please specify the source: public, private, Structural or other types of funds) Key stakeholders	In 2020, the budget of Fraunhofer IZI was € 37,9 million (48,3% from industry, 28,6% German national and regional government, 22,2% from other and 0,9% from EU). IZI partners were 179 industries and 172 from Academia in 2020. Industry, researchers, innovative centers
DES	involved	muusu y, researchers, iimovauve centers



3. GALICIAN IMMUNOTHERAPY MANUFACTURING CENTER

Project Initiative title	Galician Immunotherapy Manufacturing Center
Organisation name	Galician Health Service (SERGAS-GALARIA)
Country	Spain
Region	Galicia
Contact person	Mariona Baliu-Piqué
Contact email	Maria.Baliu.Pique@sergas.es
Website:	Not available: https://www.youtube.com/watch?v=2P2qNyAMGFg
Keywords:	Immunotherapies, clinical trials
Duration:	Not limited
Area of application	Cellular Immunotherapy Research Unit
Main challenges tackled	The first steps for the creation of a centralized manufacturing center for cellular immunotherapies for Galicia.
Objectives	To facilitate the access for Galician population to new therapies produced in this center.
Main concept and methodologies involved	The Galician immunotherapy manufacturing center, the first center based in Galicia, which focus on clinical and research activities. It is based on the production of Advanced Therapy Medicinal Products (ATMP), promotion and development of new clinical trials. Its research activity is based on the mobilization of research projects, union of research groups and lines in competitive calls to boost the execution, testing and validation of new drugs at research level.
Impacts (health, scientific, industrial, socio-economic or others enabled by the project/initiative	Establishment of a partnership among SERGAS, Hospital Clínic de Barcelona and IDIBAPS to produce new Chimeric Antigen Receptor T-cell (CAR-T) drugs for hematologic cancer treatment. Building-up the Cellular Immunotherapy Research Unit in Galicia.
Funding and Investments (please specify the source: public, private, Structural or other types of funds) Key stakeholders involved	The Galician immunotherapy manufacturing center was initially funded with more than € 3 Million (100% public nature).
Key stakeholders involved	Clinicians, researchers, hospitals, universities, research foundations and institutes



4. MEDEA PROJECT

Projec	t Initiative title	Clinical Implementation of an e-Health based Pharmacogenetics and Personalised Prescription System
Organ	isation name	Clinical Research Center of the Health Area of Badajoz of the Extremadura Health Service (SES)
Count	ry	Spain
Region		Extremadura
	ct person	Adrián Llerena
	ct email	allerena@unex.es
Websi		https://www.proyectomedea.es/en/home/
Keywo	ords:	Genetic and genomic information, clinical data collection, e- health Drug prescription, Prevention and Intervention, clinical trials
Durati	on:	2017-2022
	f application	Public procurement of innovation in pharmacogenetics and pharmacogenomics project
	Main challenges tackled	A health innovation program that looks forward the promotion of research, development and innovation in private companies for the development of personalised prescription system through the use of Public Procurement of Innovation (PPI). The first challenge for the companies, during the competitive call, is to develop a clinical decision support system with a clinical visualization supporting tool, named Personalised Oriented Prescription System (PoPS). The second challenge is the genotyping and the third challenge is the evaluation of the adverse drug reaction. The last challenge for the companies is to develop a tool for clinical trials.
	Objectives	In addition to encourage the private sector to develop and validate a personalised prescription program integrated in the electronic medical data record, MedeA also aims to generate an individualized selection strategy avoiding unnecessary risks and increase the effectiveness of the studies.
DESCRIPTION	Main concept and methodologies involved	Personalised Oriented Prescription System (PoPS) is based on three databases: genetic biomarkers, the drug-drug interactions and the electronic medical record data. Altogether will offer a screening and prevention model for adverse drug reactions (ADR)
DE	Impacts (health, scientific, industrial, socio-economic or others enabled by the project/initiative	At the end of MedeA project, there will be a personalised prescription system validated under real clinical conditions in the Extremadura Health System linked to the electronic prescription system. This will include the genetic information together with other relevant information for its use for patients in polytherapy and for prediction and prevention of Adverse Drug Reactions (ADR).
	Funding and Investments (please specify the source: public, private, Structural or other types of funds)	The budget of MedeA project is € 5,3 Million for integration of personalised prescription program in the electronic medical records.
	Key stakeholders involved	Patients, clinicians, pharmacists, medical personnel



5. EUROPEAN ASSISTANCE FOR INNOVATION PROCUREMENT (EAFIP)

Project Initiative title	EAFIP Initiative
Organisation name	DG Connect of European Commission
Country	Belgium
Region	Brussels
Contact person	-
Contact email	HTTPS://EAFIP.EU/CONTACT/
Website:	https://eafip.eu/
Keywords:	Innovation, procurement, unmet needs, value-based healthcare, technology readiness level (TRL)
Duration:	Not limited
Area of application	Innovation procurement in healthcare sector
Main challenges tackled	To promote good practices and reinforce the use of innovation procurements in different fields by following the same key strategic steps.
Objectives	The objective of the AEFIP-initiative is to promote and support other public procurers to use Pre-commercial Procurement (PCP) and Public Procurement of innovation (PPI) and boost the digital-green economy recovery through Innovation Procurement tools (PCP & PPI).
Main concept and methodologies involved	The whole cycle of innovation procurement starts with the curiosity driven research until the actual deployment of the solutions, sectors that are involved are procurers, healthcare professionals, researchers and industry. The full-blown application of EAFIP methodology encompasses 10 steps, being critical for the success of the operation, the first five preparatory steps including business case methodology before launching a procurement.
Impacts (health, scientific, industrial, socio-economic or others enabled by the project/initiative	Thanks to this methodology, a knowledge-packed toolkit with 3 modules was created, EAFIP-methodology provided free of charge technical and legal assistance to public procurers in the development and implementation of innovation procurement tools. EAFIP has also organized a series of webinars and workshops to explain the EAFIP methodology for innovation procurement.
Funding and Investments (please specify the source: public, private, Structural or other types of funds) Key stakeholders	EAFIP initiative started in 2015 financed by EU Commission (DG Connect).
Key stakeholders involved	Legal advisors, procurers, policymakers



6. CÓDIGO 100

Proje	ect Initiative title	Código100
Orga	nisation name	Galician Health Service (SERGAS), Galician Health Knowledge Agency (ACIS)
Cour	itry	Spain
Regio		Galicia
Cont	act person	José María Romero Fidalgo
Cont	act email	codigo100@sergas.es
Webs	site:	https://codigo100.sergas.es/
Keyv	vords:	Innovation, Public Procurement of Innovation Technology, Prototype, healthcare
Dura	tion:	2016-2021
	of application	Personalised Medicine, Public Health Services
	Main challenges	Código100 is an innovation plan focused on ageing, one of the main
	tackled	priorities of the Autonomous Community of Galicia.
	Objectives	Código100 aimed to obtain ideas from patients, professionals and
		companies in the sector. The ideas were grouped around the three main lines:
		- Patient empowerment: to strengthen the rights and capacities of citizens.
		- Therapies: to increase the Personalised Medicine in SERGAS by developments in biotechnology and the biomedical area.
		- Professional solutions: to increase and gain skills of clinical personnel,
		update the system and promote an innovative culture among
		professionals.
		Código 100 promoted solutions that brought clear healthcare benefits within SERGAS as well as acted as a catalyst for innovation in the healthcare sector.
	Main concept and	The first task of Código100 involved the implementation of a market
	methodologies	survey stage to detect the needs required for each project. The
	involved	information exchanged during this consultation stage was essential to
		address real healthcare challenges through technological solutions that
		had not currently existed.
	Impacts (health, scientific, industrial,	As a result of Código100, different products were developed and purchased by SERGAS, such as:
	socio-economic or	- the SIPAD: a mobile APP for diabetes patients.
	others enabled by the	•
	project/initiative	- SHARE, which is a large platform containing all kind of multimedia
		content to provide patients clinical information. It analyses the patient
		profile and detects the main areas that patient is interested in obtaining
		information, creating personalised content for each patient.
		- e4Quant: a quantitative test of apolipoprotein E4 detection in human
		plasma designed by Biocross for prediction of Alzheimer disease
		progression.
		- PharmaHIC-ReTER Test by Healthincode: a Next Generation Sequencing
		for genetic biomarkers related to the response to biological drugs for
		rheumatology PQreader+Promonitor Test by Progenika Biopharma: an integrated point
7		of care solution for quantitative monitoring of biological drugs and
O		qualitative monitoring of anti-drug antibodies in human blood for
PT.		rheumatology uses.
KI		- OMTX100 by Oncomatrix: a non-invasive diagnosis of solid tumor (colon
DESCRIPTION		and lung cancer) in blood exosomes using a novel combination of genes
		expressed in tumor epithelial cells and in tumor microenvironment cells.
		The state of the s



Funding and Investments (please specify the source: public, private, Structural or other types of funds)	Código100 was funded by € 13 million (80% from ERDF funding by collaboration with the Ministry of Science and Innovation).
Key stakeholders	General practitioners, clinicians, researchers
involved	



7. LOMBARDY CLUSTER TECHNOLOGIES FOR LIVING ENVIRONMENTS

Proje	ect Initiative title	Lombardy Cluster Technologies for Living Environments
Orga	nisation name	TAV - TechForLife Cluster
Cour	ntry	Italy
Regi	on	Lombardy
	act person	Cristina De Capitani
	act email	<u>cristina.decapitani@polimi.it</u>
Web		https://duster.techforlife.it/
	vords:	Technology, Healthcare, Innovation
	of application	From 2013 to now
Alea	of application Main challenges tackled	Clinical developments This network focuses on the management of chronicity and development of new innovative technologies to tackle the chronicity.
	Objectives	The Cluster was created with the aim of facilitating and supporting research, innovation and training in the Lombardy region by acting as a catalyst of multidisciplinary and multiprofessional skills and expertise. It also supports and facilitates the commercial development of the Cluster. In terms of personalised medicine, the Cluster of Technologies for Living Environment (Cluster TAV) focuses on the patient lifestyle, including nutrition and rehabilitation that affects the management of chronicity.
	Main concept and methodologies involved	The Cluster TAV supports and promotes research, innovation and training in the development of technologies for the living environment, by a synergistic relationship between industry and research.
		To support the innovation process, the Cluster promotes collaboration and dialogue between partners and establishes a regulatory and institutional ecosystem that enables to start and participate in common projects on personalised medicine.
DESCRIPTION	Impacts (health, scientific, industrial, socio-economic or others enabled by the project/initiative	projects were developed, such as:



Funding and	The TAV - TechForLife Cluster is sustained by annual fees from
Investments (please	TAV partners, participation as partners in different projects.
specify the source:	
public, private,	
Structural or other	
types of funds)	
Key stakeholders	General practitioners, clinicians, researchers, patients, industry
involved	



8. GALICIAN NETWORK OF HEALTH LIVING LABS (LABSAÚDE)

Project Initiative titl	Galician Network of Health Living Labs (LABSAUDE)
Organisation name	University Hospital Complex of Ourense (CHUO), Galician Health Service (SERGAS), Galician Health Knowledge Agency (ACIS)
Country	Spain
Region	Galicia
Contact person	José María Romero Fidalgo
Contact email	labsaude@sergas.es
Website:	https://labsaude.sergas.es/
Keywords:	Technology, Personalised Medicine
Duration:	From 2015 to now
Area of application	Personalised Medicine, personalised technologies, Public Health Services, open innovation, Technological Readiness Level (TRL) 7 or higher
Main cha tackled	Living Lab of Ourense is the first Living Lab from Galician Network of Health Living Labs (LABSAUDE) to provide an environment for patients, medical and health personnel, and biomedical companies to share knowledge and design, test and evaluate together innovative solutions with healthcare real needs.
Objectives	The Galician Network of Health Living Labs - LABSAÚDE, is an initiative that will convert Galician Hospitals and health centers into authentic testing laboratories for new solutions, products or services related to health and thus create an ecosystem of multidisciplinary and multisectoral co-creation networks (health professionals, administration, patients and industry) focused on end users.
	The Living Lab of Ourense is the pilot project from LABSAÚDE. The aim of the network is to spread the creation of this type of end-user-tailored settings to the rest of the Galician hospitals.
Main concept methodologies involved	Thanks to the Living lab of Ourense, SERGAS is testing a real future situation in where patients suffer with two or more chronic diseases and need more socio-health care services, that situation is forecast for 2050 throughout Europe. Within this initiative, the solutions for healthy ageing that issue in Ourense will become an international benchmark.
	Different innovation lines are expected for Living Lab scenarios in the context of LABSAÚDE: elderly and/or chronic patients care, Patient empowerment; ICT/ Telemedicine and teleservice; Robotics and virtual reality; Impact of the environment in hospital stays; or Other (biosafety, new materials, nutrition)
Impacts (scientific, ind socio-economic others enabled project/initiative)	or the patients with severe stroke. Wards are also available for testing and evaluating solutions based on the use of new technologies, for example dynamic posturographer for rehabilitation after hip fractures. In total, there are 15 wards that enable to continuously test the experience of patients and check their response to innovative formulas to address ageing-diseases (i.e., respiratory rehabilitation in patients with exacerbated COPD).
Ц	It also allows to develop and scale solutions as a spearhead towards the



		market by public call for project selection.
	Funding and Investments (please specify the source: public, private, Structural or other types of funds)	The facilities of the Living Lab of Ourense were built by PPI-based project Hospital2050.
	Key stakeholders involved	Medical and healthcare professionals, biomedical companies, patients