



Maternal, Neonatal Health and Developmental Research Network SAMID (RD16/0021/0001)

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(Valencia; Spain)

UENPS VALENCIA 2016





Aims & Objectives

- Promote research throughout the perinatal period (pregnancy & fetal development), infancy and childhood.
- Evaluate endogenous (genetic) and exogenous (environmental) factors that influence growth, development and health of the individual from fetal stage to the end of growth and development and influence upon adult age health status.





Primary objective

To study conditions that occur in the perinatal period, infancy and adolescence and how they may influence adult health status from different perspectives:

- Etiopathogenic (mechanisms)
- Preventive
- Therapeutic
- Evolving (long-term consequences).





Strategic objectives

- To generate new high-quality and relevant social and health knowledge in a specific time frame of human life span.
- 2. To optimize human and technical research resources.
 - ✓ Incorporating young researchers
 - ✓ Stimulating multidisciplinary approach to problems.
 - ✓ Enhancing financial support for multinational resources
- 3. To promote incorporation of new groups and research centers to the RETIC RED SAMID enhancing the quality and quantity of biomedical research in Spain.
- 4. To promote the incorporation of RETIC RED SAMID to European Obstetrics and Research Infrastructures to contribute to the establishment of an European Maternal and Pediatric Research Network.





Tactical objectives

- To promote the figure of the Pre-and-Post Doctoral Research Fellows in the field of experimental and clinical Perinatology
- To further involve Staff neonatologists and obstetricians in clinical and basic research (protected time in referral centers)
- To establish clinical and methodological synergies
- To develop animal models
- To establish a network of European centers of excellence to launch multicenter clinical trials.



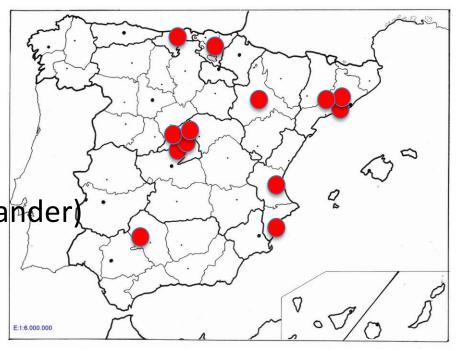


- 13 acknowledged Research Groups
- 120 clinical and basic researchers
- 12 young investigators with full research contract
- 1 network manager
- Budget allocated to promote the network, meetings, scientist stages, publications, webpage, etc.





- HU La Fe (Valencia)
- HU La Paz (Madrid; 2)
- HU 12 de Octubre (Madrid)
- HU Gregorio Marañón (Madrid)
- HU Cruces (Barakaldo, Bilbao)
- Universidad de Granada
- Universidad de Murcia
- HU Marqués de Valdecilla (Santander)
- H del Mar (Barcelona)
- H Vall d'Hebrón (Barcelona)
- H S Joan de Deu (Barcelona)
- Universidad de Zaragoza







- Cooperation with related Health Research Institutes:
 - Ancillary Methodological Platforms (epidemiology, statistics, economy, legal, ethics, etc.)
 - Analytical Platform ("omics"; basic laboratory needs)
- Basic Scientists:
 - Biologists, Geneticists, Analytical Chemists
 Pharmacologists, Microbiologist, Pathologist, etc.
- Clinical staff
 - Obstetrics, Surgery, Radiology, Neonatology, Nutrition, Endocrinology, etc.





Associated Clinical Groups

- A total of 23 hospitals
- Devoted to actively participate in clinical trial.
- Adequate infrastructure to recruit eligible patients
- Infrastructure to obtain, process, store and ship biological samples
- Are endowed with special funding
- Can constitute in the future an Acknowledge Research Group



INFRASTRUCTURE OF SAMID NETWORK





Acknowledged Groups

Committees

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EXPERIMENTAL

CLINICAL

RESEARCH

RESEARCH

TRAINING TEACHING

COMMUNICATION

INTRANETWORK COORDINATION

10





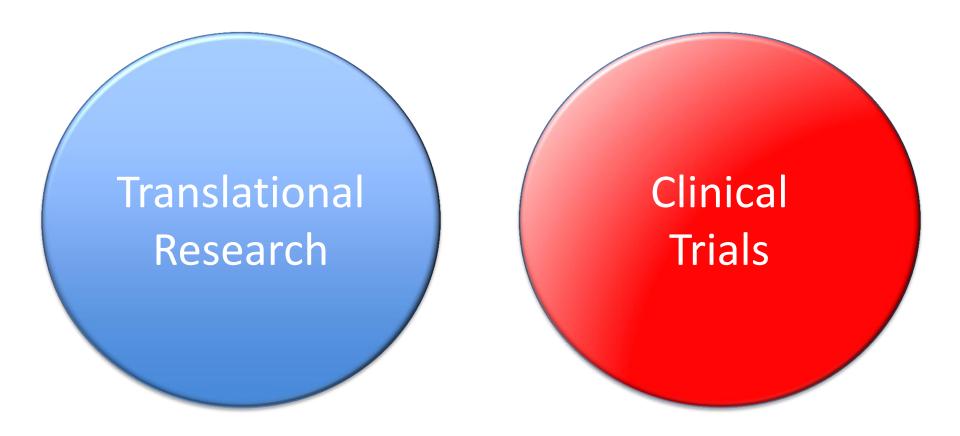
Web SAMID network: www.redsamid.net







Research involvement of SAMID







Program #1 accepted by the ISCIII

(Ministry of Economy, Industry, Competitiveness; Kingdom of Spain)

STATEMENT OF THE PROGRAMME RETICS COORDINATOR

RETIC Code RD16/0022/0001

Network (Thematic area): MATERNAL, INFANT AND DEVELOPMENTAL HEALTH RESEARCH NETWORK						
Programme Title:	Interventions during pregnancy, neonatal period, and pediatric age for prevention and treatment of conditions influencing adult health.					
Programme Leader:	MAXIMO VENTO TORRES					
Applicant institution:	FUNDACION PARA LA INVESTIGACIÓN HOSPITAL LA FE					
Work institution:	HOSPITAL UNIVERSITARIO Y POLITÉCNICO LA FE					
Requested budget:	2.473.100,00 €					



Applicant institution:

Work institution:

Requested budget:



RETIC Code

Program #2 accepted by the ISCIII

(Ministry of Economy, Industry, Competitiveness; Kingdom of Spain)

STATEMENT OF THE PROGRAMME RETICS COORDINATOR

Before submitting your application please read carefully the instructions provided on the last page,

you will not be able to correctly unload this document

2.070.750 Euros

otherwise you will not be as	RD16/0022/0015						
Network (Thematic area): MATERNAL, INFANT AND DEVELOPMENTAL HEALTH RESEARCH NETWORK							
Programme Title:	CARDIOVASCULAR RISK-RELATED PRENATAL FACTORS						
Programme Leader:	ELISA LLURBA OLIVÉ						

FUNDACIÓ INSTITUT DE RECERCA DE LA VALL D'HEBRON (VHIR)

HOSPITAL UNIVERSITARIO DE LA VALL D'HEBRON





Program #1 accepted by the ISCIII

(Ministry of Economy, Industry, Competitiveness; Kingdom of Spain)

OBJECTIVE 1

To study biomarkers of Intrauterine Growth Retardation, preeclampsia and chorioamnionitis capable of predicting postnatal developmental disorders related to neurocognitive development and nutritional and metabolic status.

OBJECTIVE 2

To study the efficacy of clinical indicators and biomarkers to early predicting neurocognitive and developmental outcomes and identify risk factors for neurologic sequelae in children after acute neurologic injuries, and to analyze the effectiveness of diagnostic methods and prevention strategies.

OBJECTIVE 3

To study neonatal risk factors for adverse nutritional and metabolic outcomes and possible preventive nutritional strategies.

OBJECTIVE 4

To study pre-and-postnatal environmental factors associated with neurologic, nutritional, and metabolic conditions in the perinatal and childhood periods.





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DESCRIPTION OF EACH WORKPACKAGE

WP N°	1	Start Date/Start Event		01/01/2017		End Date/End Event		31/12/2021		
WP TITLE	PRENATAL RISK FACTORS FOR ABNORMAL NEONATAL AND LONGTERM OUTCOME									
PI Family Name	LLURBA	GOMEZ	VENTO	CABAÑAS	PALLAS	LOPEZ DE HEREDIA	GARCIA ALGAR	RODRIGUEZ		
Group N°	5	4	1	2	3	6	7	10		
N° of Resarchers	7	11	17	16	18	10	10	9		

OBJECTIVES Max. 1,200 characters

^{1.1} To establish a protocol for the diagnosis and obstetrical management of conditions prompting preterm delivery including preeclampsia, IUGR, chorioamnionitis.

^{1.2} To study the interrelation between epidemiological, toxic, environmental, socio-economic variables and maternal risk factors leading to impairment of neurodevelopment.

^{1.3} To study the relationship between Doppler hemodynamic compromise of umbilical blood flow and alteration of neurodevelopment.

^{1.4} To study the interrelation between perinatal complications, postnatal adaptation and neurodevelopment.

^{1.5} To develop a fetal model of IUGR, to study factor affective fetal growth and test possible preventive and therapeutic strategies.





DRUG COMPANIES

PUBLIC & PRIVATE RESEARCH ENTITIES

HOSPITALS
NETWORK
(SAMID)

Enpr-EMA **IDENTIFICATION OF RESEARCH PRIORITIES**



STUDY DESIGN



SCReN

SELECTION OF PERTINENT INSTITUTIONS (DATA BASE; INFRASTRUCTURE)

FEED BACK





However the niche of opportunity for the promotion of clinical trials funded by International Institutions requires changes in logistic approach!





Spanish Clinical Research Network (SCReN) Platform for Clinical Research and Trials













Enpr-EMA

Objective:

To facilitate studies in order to increase availability of medicinal products authorised for use in the paediatric population.

Initiatives:

- ✓ fostering high-quality, ethical research on the quality, safety and efficacy of medicines for use in children
- ✓ helping with the recruitment of patients for clinical trials
- ✓ enabling collaboration between networks and stakeholders
- ✓ avoiding unnecessary duplication of studies
- ✓ building up scientific and administrative competence at a European level
- ✓ promoting **European Commission framework** programme applications.









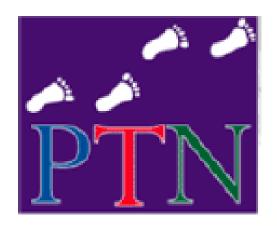


Innovative medicines initiative

- The Innovative Medicines Initiative (IMI) is Europe's largest public-private initiative aiming to speed up the development of better and safer medicines for patients.
- IMI supports collaborative research projects and builds networks of industrial and academic experts in order to boost pharmaceutical innovation in Europe.
- IMI is a joint undertaking between the European Union and the pharmaceutical industry association EFPIA.
- IMI 2 Call 6.







Pediatric Trials Network





Paediatric Trials Network

- The Paediatric Trials Network (PTN), sponsored by the National Institute of Child Health and Human Development (NICHD), is an alliance of clinical research sites cooperating in the design and conduct of paediatric clinical trials.
- PTN research provides evidence for optimal dosing of commonly used medications in infants and children, improving healthcare for these patients.





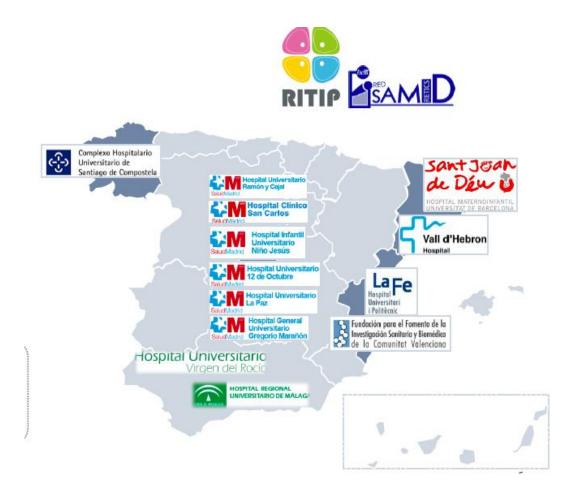
Spanish Paediatric Clinical Trials Network (RECLIP)

RECLIP Presentation Report





Who we are a) national thematic networks







Who we are: b) Current partners

CURRENT PARTNERS OF RECLIP







RECLIP SCIENTIFIC STRUCTURE







Objectives of RECLIP

The main objective of RECLIP is to provide a fully developed and professionalized network of reference national clinical trial units, ensuring an appropriate infrastructure for conducting safe and effective paediatric clinical trials.

HOW?

Through a coordinated and super-specialized network and a professionalized, multidisciplinary and organized stable structure, integrating the most important paediatric clinical research units across the country, located in the most important hospitals in Spain.

WITH WHOM?

RECLIP develops its activity through the **cooperation with key stakeholders** inside and outside the network. The network joins strong and experienced national paediatric clinical and academic institutions with a long history of cross-collaboration with each other and the pharmaceutical industry.

WHY?

The collaboration between reference entities in RECLIP allows to **reduce the uncertainty** and the risk inherent to the development of paediatric trials and **helps to accelerate** the process. All partners offer their expertise and best practices for the achievement of the aim of RECLIP.

Thanks for your attention!