





TEDDY is an independent multidisciplinary and multinational Network including partners from 18 countries and is aimed at facilitating the performance of good quality paediatric studies and research. TEDDY is a member of Enpr-EMA, the European Network of Paediatric Research at the European Medicines Agency.

MISSION

-  To implement good practices and tools to develop studies and research on medicinal products in children in compliance with EU legislation and guidelines
-  To promote, participate in and provide methodological expertise to set up, develop and conduct paediatric national or multinational clinical trials and other paediatric studies
-  To collaborate with existing paediatric networks and research organisations
-  To develop communication and educational tools tailored for children in order to facilitate their participation in all the relevant experimental procedures including assent and consent forms preparation



TEDDY NETWORK is coordinated by

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ACTIVITIES



PROJECTS

TEDDY started as a FP6 funded project and promoted, together with its members, several FP7 projects including research-driven paediatric clinical trials: CloSed, DEEP, GAPP, NeoMero, NeoVanc. In the new Horizon 2020 framework, SMART - Small Medicines Advanced Research Training, a Twinning project started in 2016, is gathering 4 TEDDY members, and PedCRIN - Paediatric Clinical Research Infrastructure Network, an Infradev3 project coordinated by ECRIN and involving CVBF-TEDDY as a partner, will soon be kicked off.



INFORMATIVE TOOLS

- ★ TEDDY is now performing an online survey to collect general information on investigational centres in EU and non-EU countries participating in paediatric research. Results will be made available to EnprEMA (the European Network for Paediatric Research at the EMA) and will be used in the implementation phase of EPCT-RI, the proposed European Paediatric Clinical Trials - Research Infrastructure.
- ★ A database, the European Paediatric Medicines Database (EPMD), containing information on paediatric drugs authorised by the European Medicines Agency (EMA) under the centralised procedure, has been created and is available online.
- ★ An Inventory of the Ethical and Regulatory procedures for paediatric clinical trial (CT) approval and conduction in EU and non-EU countries has been performed. It includes information on the relevant local and national requirements and procedures to apply Ethics Committees and concerned Authorities' authorisation and to submit periodic information to the same Bodies. The inventory is going to be implemented with pharmacovigilance requirements.



PRODUCTS

- ★ Study Specific Procedures for paediatric clinical trials.
- ★ Videos and age-appropriate informative material to duly inform children and families.
- ★ E-learning courses to train Investigators and patients for an effective participation in clinical trials.



Join the TEDDY NETWORK!

For more information please contact
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or visit our website

www.teddynetwork.net