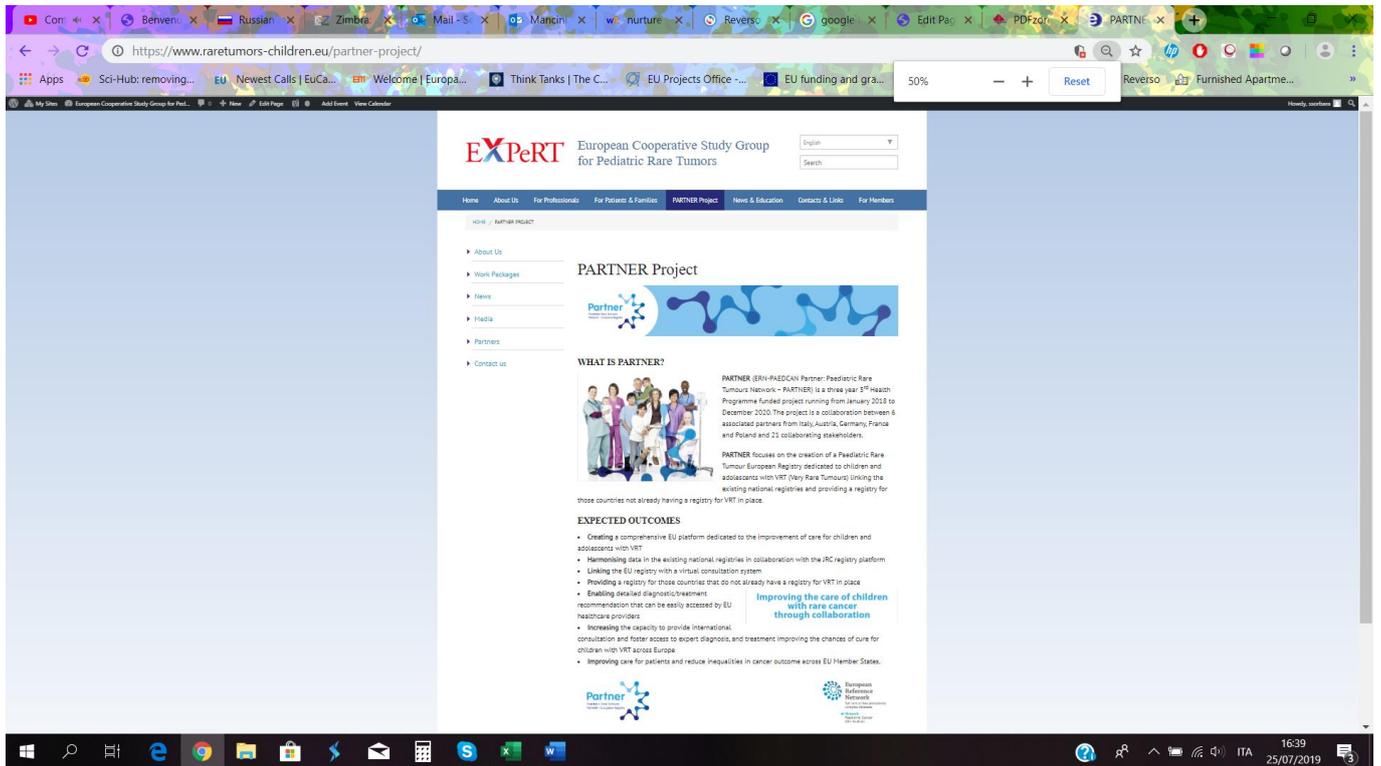




**ERN-PAEDCAN Partner - Paediatric Rare Tumours Network - European Registry
(N°777336)**

D2.3 – Project website launched



EXPeRT European Cooperative Study Group for Pediatric Rare Tumors

English Search

Home About Us For Professionals For Patients & Families **PARTNER Project** News & Education Contacts & Links For Members

Home / PARTNER PROJECT

- About Us
- Work Packages
- News
- Media
- Partners
- Contact us

PARTNER Project



WHAT IS PARTNER?

PARTNER (ERIN-PAEDI-CAN Partner: Paediatric Rare Tumours Network - PARTNER) is a three year ST Health Programme funded project running from January 2018 to December 2020. The project is a collaboration between 6 associated partners from Italy, Austria, Germany, France and Poland and 21 collaborating stakeholders.

PARTNER focuses on the creation of a Paediatric Rare Tumour European Registry dedicated to children and adolescents with VRT (Very Rare Tumours) linking the existing national registries and providing a registry for those countries not already having a registry for VRT in place.

EXPECTED OUTCOMES

- Creating a comprehensive EU platform dedicated to the improvement of care for children and adolescents with VRT
- Harmonising data in the existing national registries in collaboration with the JRC registry platform
- Linking the EU registry with a virtual consultation system
- Providing a registry for those countries that do not already have a registry for VRT in place
- Enabling detailed diagnostic/treatment recommendations that can be easily accessed by EU healthcare providers
- Increasing the capacity to provide international consultation and foster access to expert diagnosis, and treatment improving the chances of cure for children with VRT across Europe
- Improving care for patients and reduce inequalities in cancer outcome across EU Member States.

Improving the care of children with rare cancers through collaboration