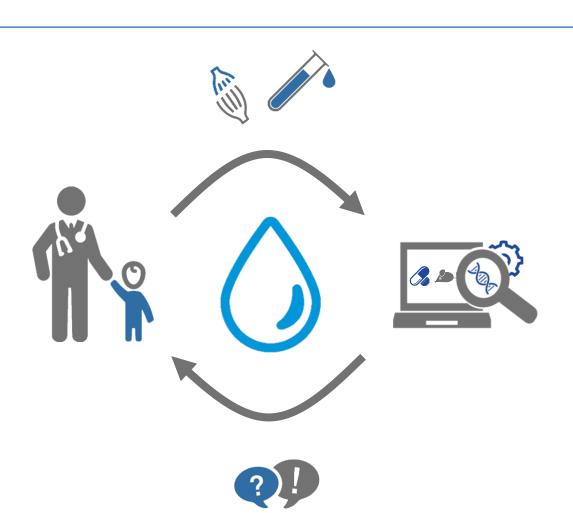
TEARDROP: Building a Belgian Pediatric Oncology specific Bioportal

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Introduction

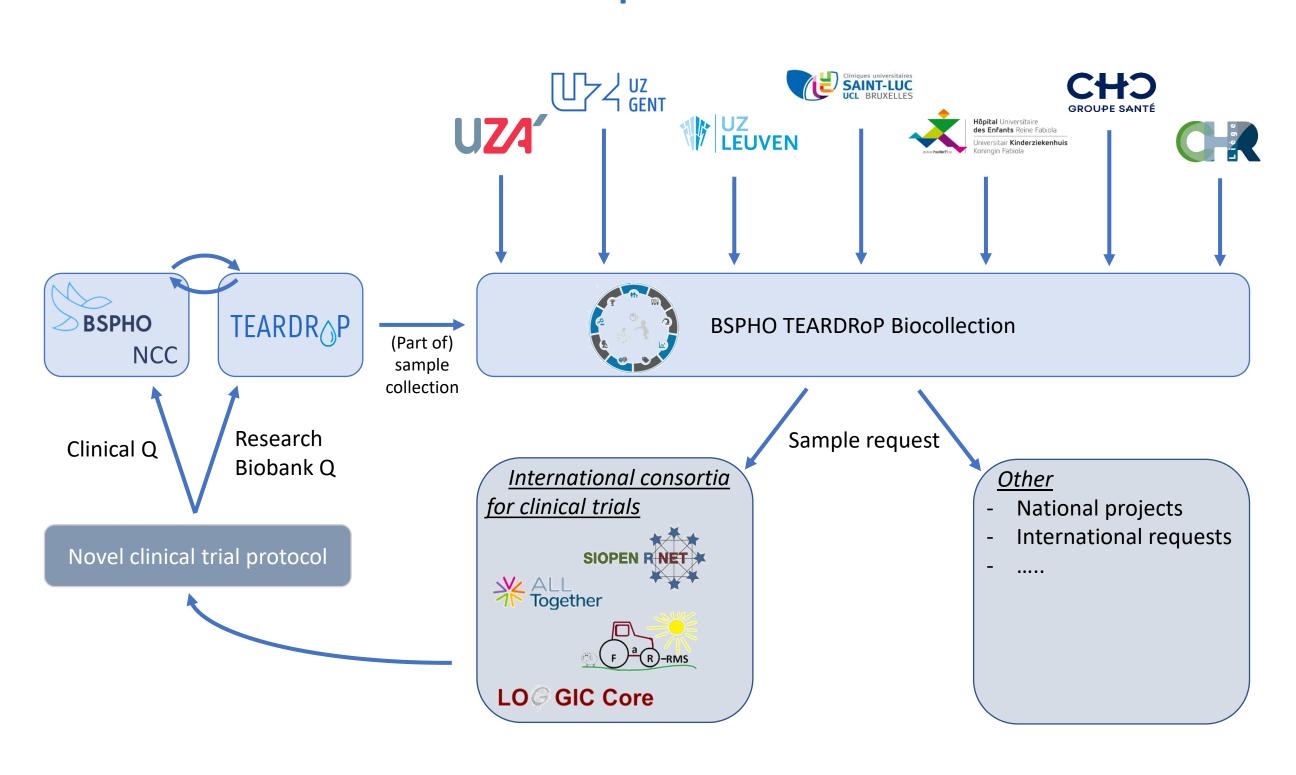
- Pediatric cancer is rare and requires specialized care for developing children
- Survival rates have stagnated in recent decades
- Need for international clinical trials and novel treatment options
- Increasing trial complexity leads to high sampling demands
- State-of-the-art research requires high quality, well annotated, longitudinal, patient sampling



TEARDROP Oncology consortium aims to:

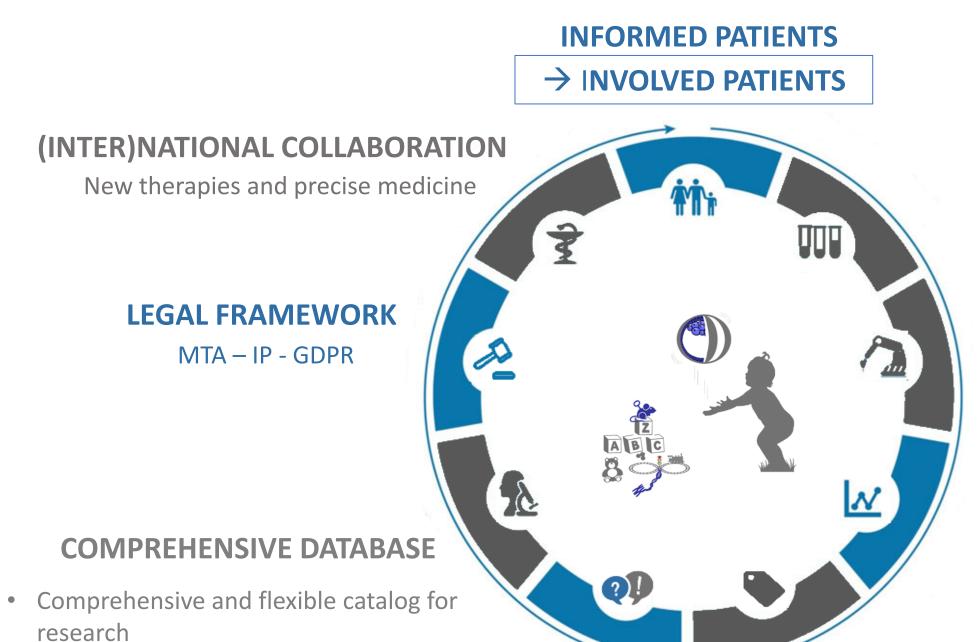
- Facilitate access to patient material for researchers and provide clinicians preclinical data on new treatment concepts
- Establish a national pediatric-oncology specific biocollection, integrating patient and sample data for (inter)national research
- Provide uniform procedures for high quality sample collection implemented in routine clinical care resulting in more complete sample collections, fit for use in the next generation of fundamental research efforts

Power of centralization



Collaboration with the **BSPHO NCC** enables Belgian pediatric oncology centers to **participate in international** research efforts. Basic clinical annotations and storage information on collected samples is not always easily accessible outside the study objectives of the international clinical trials. Moreover, off-study patients are often underrepresented. Setting-up a national bioportal, a registry linked to a virtual biobank collection for pediatric oncological patients as an intermediate vehicle ensures better data accessibility and overview, strengthening the Belgian position in international collaborations.

TEARDRoP Hallmarks



DIALOGUE

Clinicians, researchers, patient advisory board

TUMORBANK LOCAL

- Harmonized sample collection for
- excellent quality Sample processing & storage

TUMORBANK

- VIRTUAL = REGISTRY Collection of core clinical and
- epidemiological information • Additional site -specific activities and suppl. samples/data

CLINICAL TRIALS vs FUTURE RESEARCH

The goal of TEARDROP is to be a vehicle to assimilate the different (inter)national clinical trials, projects and studies into a unified infrastructure over time. TEARDROP will result in a national biobanking portal encompassing fully clinically annotated sample sets that can be easily explored and retrieved thanks to a preset national legal framework. It aims to break existing barriers, and lower thresholds for research into pediatric oncology.

Rise of the infrastructure

TEARDRAP Pediatric Oncology specific Bioportal



Site-specific flows

Less labour-intensive

Efficiency

GDPR-compliant

Harmonized Sample Collection

Sample manual 'translation'

Prospective collection

Common denominator

Quality criteria:

Standard set of samples

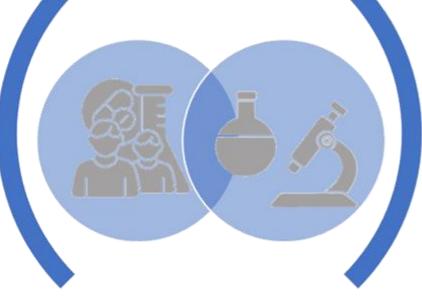
Bottom-up approach for sample

Basic samples: high quality SOPs

SOPs to adhere to clinical routine

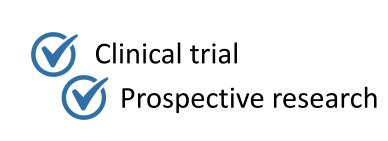
Advanced samples: BSPHO

reference labs





National Bioportal



National guidelines

Overview at sample level Strengthen position @ International consortia

Sample and clinical annotation resource

- Define a minimal dataset
- Both on sample and clinical annotation
- Incorporating:
- Legal obligation
- Cancer registry requirements
- International clinical trial eCRF standard

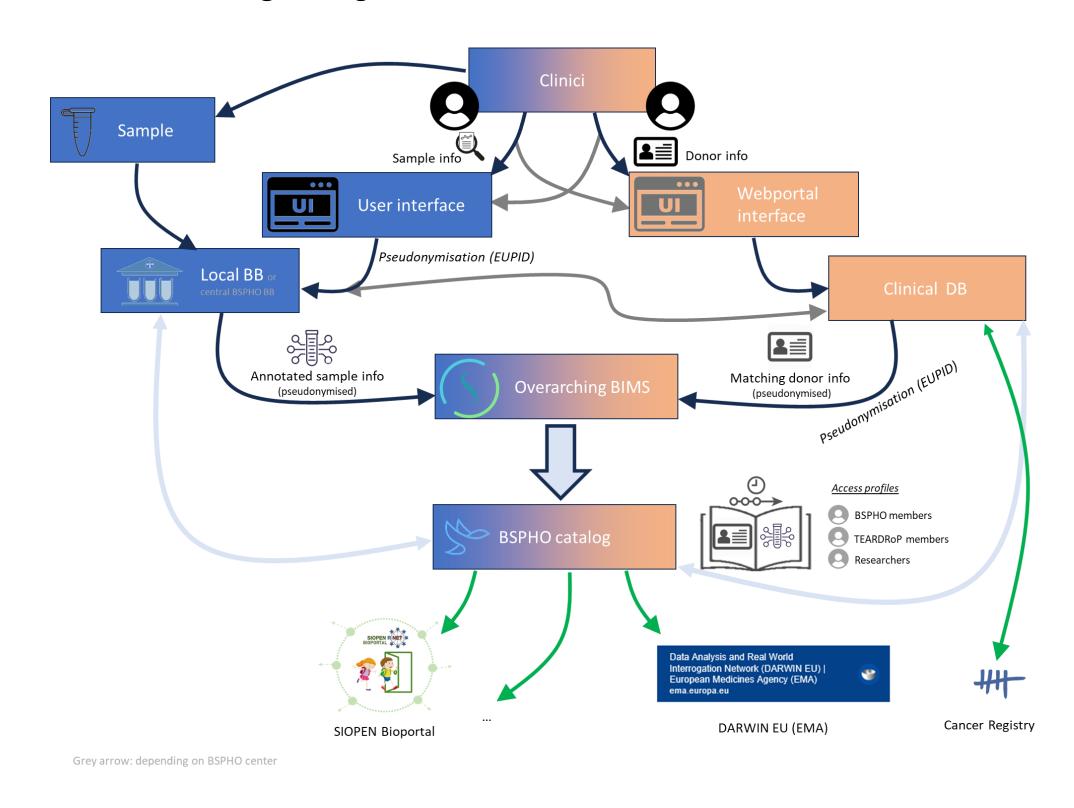
Bioportal management structure

- TEARDRoP board
- Consortium partners
- Project coordinators
- Disease specific working groups

Roadmap to success

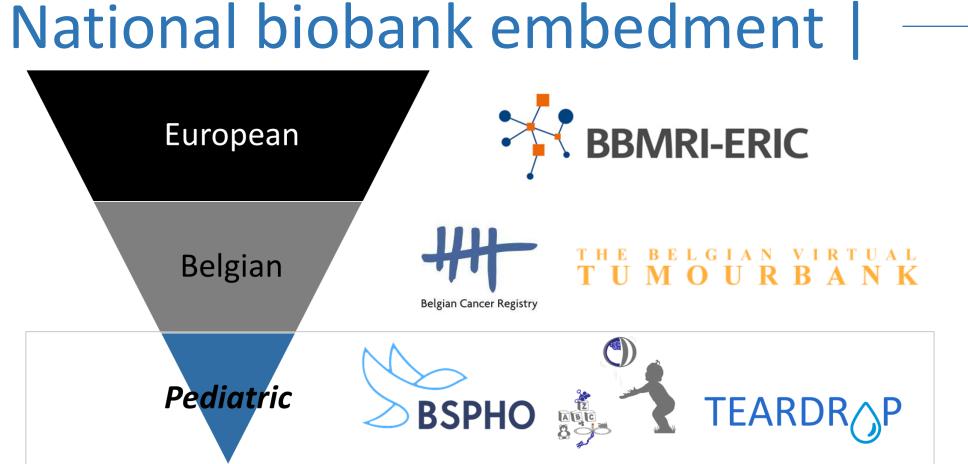
Overview & traceability

The value of any biospecimen collection is largely determined by the quality of the accompanying sample annotation. The data will be merged by means of a Datawarehouse in an overarching BIMS to be able to offer a comprehensive searchable catalog through the BSPHO website.



Sample routing is kept as close to **clinical routine** as possible, to improve efficiency and relieve the burden on both the clinical teams and teams involved in routine patient care. Sample types are defined:

- Basic sample types will be collected and processed in local biobanks.
- Advanced sample type processing will be organised by BSPHO reference labs



There are general Belgian registries while a pediatric oncology-specific sample collection is a growing need. Many neighbouring countries have successfully implemented a similar structure. Therefore, we collaborate with existing initiatives to ensure (inter)national biobank embedment.



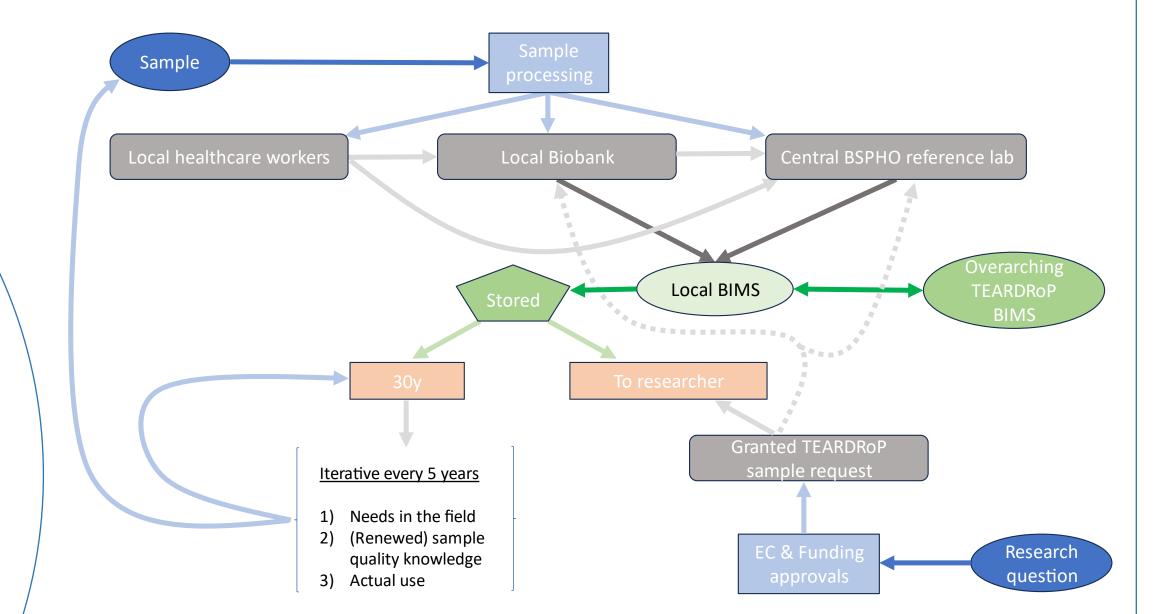
TEARDROP as a vehicle to set-up a high-quality, harmonized national PedOnco sample collection

Key takeaways



TEARDROP in a research-inspired nation-wide supportive role through clinical embedment

TEARDROP infrastructure to tackle patient sampling challenges to allow elevated BSPHO (inter)national clinical trial support.



The procedure for **sample requests** within the BSPHO community is structured to reflect collective decision-making. A brief project application template will allow users to provide a scientific rationale which will be reviewed by the TEARDROP disease specific working group on adherence to the overarching goals of the BSPHO and the embedment into the (inter)national pediatric oncology research community.



























