Privanova

RESEARCH & CONSULTING

Privanova is part of the FACILITATE: a multidisciplinary R&D project ran by an international consortium and funded by the EU within the Horizon2020 Innovative Medicines Initiative.

FACILITATE reimagines the dynamics of medical data processing by giving the data subjects transparency and control over the utilisation of their data while granting reliable material for medical innovation. Its strategy represents a unique, innovative opportunity for medicines drug development and regulation to better understand the clinics of diseases, and to evaluate the effectiveness of products in the healthcare system.

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FACILITATE

Framework for clinical trial participants data reutilisation for a fully transparent and ethical ecosystem

Budget

€ 6 886 711
From Jan. 2022 to Dec. 2025

Consortium

27 Partners from including SMEs, Pharmaceutical Industry, Academia, Hospitals, and Patient Organisations

Privanova's role

Privanova leads the WP7 on Business exploitation and sustainability. We guide the project partners and help them define the FACILITATE business plan in terms of use cases, potential, benefits, strategy requirements and associated options ensuring patients' empowerment.

In addition, we take part in WP2 where we identify major legal issues and propose solutions to address them in order to return coded/ indirectly identifiable data from sponsors conducting clinical trials to patients and to allow research with such data focusing on the interplay between the Clinical Trials Regulation (CTR) and the General Data Protection Regulation (GDPR).

Impact

To avoid the current situation where clinical data are siloed in separate medical systems without possibility to be used beyond their original, limited single-sided purpose, FACILITATE provides clear rules in a trusted ethical, legal and regulatory ecosystem to strongly engage patients as data generators.

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