

A Post-Authorization Safety Study (PASS) to Assess Long-term Safety in Patients with Pompe Disease Treated with Avalglucosidase alfa in the Real-World Setting: the SAVANT (Safety assessment of AVAlglucosidase Alfa as a Novel Treatment) study

First published: 15/11/2023

Last updated: 15/07/2025

Study

Ongoing

Administrative details

EU PAS number

EUPAS107472

Study ID

107473

DARWIN EU® study

No

Study countries

-  Belgium
 -  Canada
 -  France
 -  Germany
 -  Italy
 -  Netherlands
 -  Spain
 -  United Kingdom
 -  United States
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Study description

This PASS is a non-interventional, multicenter, longitudinal study of safety with prospectively collected data in participants with Pompe disease who are receiving ERT with avalglucosidase alfa as per SoC determined by their treating physician and will be followed for safety reporting using the PASS Recommended Schedule of Assessments provided in the protocol as a guidance.

The site selection will be worldwide, with participating sites from multiple geographical regions, particularly sites participating in the Pompe Registry and in countries where home infusion is allowed per local regulation.

The Pompe Registry (www.registrynxt.com) is a global, multi-center, multi-national, longitudinal, observational, and voluntary program that is currently conducted to track the disease history and outcomes in patients with Pompe disease

Study status

Ongoing

Research institutions and networks

Institutions

Sanofi

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Institution

Contact details

Study institution contact

Trial Transparency Team Trial Transparency Team contact-us@sanofi.com

Study contact

contact-us@sanofi.com

Primary lead investigator

Trial Transparency Team Trial Transparency Team

Primary lead investigator

Study timelines

Date when funding contract was signed

Planned: 06/10/2023

Actual: 06/10/2023

Study start date

Planned: 30/06/2024

Actual: 15/07/2024

Date of final study report

Planned: 31/12/2029

Sources of funding

- Pharmaceutical company and other private sector

More details on funding

Sanofi

Regulatory

Was the study required by a regulatory body?

No

Is the study required by a Risk Management Plan (RMP)?

EU RMP category 3 (required)

Methodological aspects

Study type

Study type list

Study topic:

Disease /health condition

Study type:

Non-interventional study

Main study objective:

Evaluate the important identified risk of IARs including hypersensitivity and anaphylactic reactions

Study drug and medical condition

Medicinal product name

NEXVIADYME

Medicinal product name, other

Nexviadyme

Study drug International non-proprietary name (INN) or common name

AVALGLUCOSIDASE ALFA

Population studied

Age groups

- Term newborn infants (0 - 27 days)
- Infants and toddlers (28 days - 23 months)
- Children (2 to < 12 years)

- Adolescents (12 to < 18 years)
 - Adults (18 to < 46 years)
 - Adults (46 to < 65 years)
 - Adults (65 to < 75 years)
 - Adults (75 to < 85 years)
 - Adults (85 years and over)
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Estimated number of subjects

100

Study design details

Outcomes

Incidence rate and event rate of participants with IARs
Incidence rate and event rate of IARs

Cumulative incidence of IARs leading to treatment discontinuation

Incidence rate and event rate of medication errors

Cumulative incidence of medication errors leading to AE/SAE/treatment discontinuation,

Incidence of all AEs during study participation, including but not limited to AESIs and AEs leading to permanent treatment discontinuations,

Incidence of severe hypersensitivity reactions, anaphylactic reactions, and severe immune mediated reactions, Incidence of AEs with permanent treatment discontinuation as outcome,

Incidence of ADA/NAb over time

Data analysis plan

The primary analysis approach is descriptive, and the same approach will apply to both the primary and secondary endpoints. No hypothesis testing is planned.

Stratification by prior exposure category (ie, avagliflozin use prior to enrollment, prior use of other ERTs), region (Europe versus rest of the world) and infusion setting (clinic/hospital versus home) will also be performed, given adequate representations in each stratum.

Data management

ENCePP Seal

The use of the ENCePP Seal has been discontinued since February 2025. The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.

Data sources

Data sources (types)

[Other](#)

Use of a Common Data Model (CDM)

CDM mapping

No

Data quality specifications

Check conformance

Unknown

Check completeness

Unknown

Check stability

Unknown

Check logical consistency

Unknown

Data characterisation

Data characterisation conducted

No