

# An observational, prospective study to assess the use, effectiveness, and safety of voxelotor in patients with sickle cell disease (SCD) in a real-life setting (VoxEval)

**First published:** 05/06/2024

**Last updated:** 24/02/2025

Study

Cancelled

## Administrative details

### EU PAS number

EUPAS1000000186

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### Study ID

1000000186

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### DARWIN EU® study

No

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### Study countries

 France

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### Study status

Cancelled

## Contact details

### Study institution contact

Delphine BERZIN delphine.berzin@pfizer.com

Study contact

[delphine.berzin@pfizer.com](mailto:delphine.berzin@pfizer.com)

### Primary lead investigator

Jean-Benoit ARLET

Primary lead investigator

## Study timelines

### Date when funding contract was signed

Actual: 31/07/2023

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### Study start date

Planned: 15/10/2024

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### Date of final study report

Planned: 31/12/2030

## Regulatory

### Was the study required by a regulatory body?

Yes

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

Not applicable

## Methodological aspects

**Study type:**

Non-interventional study

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**Scope of the study:**

Drug utilisation

Effectiveness study (incl. comparative)

Safety study (incl. comparative)

## Study drug and medical condition

**Medicinal product name**

OXBRYTA

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

VOXELOTOR

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**Anatomical Therapeutic Chemical (ATC) code**

(B06AX03) voxelotor

voxelotor

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**Medical condition to be studied**

Sickle cell disease

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 The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.

## Use of a Common Data Model (CDM)

### **CDM mapping**

No

## Data quality specifications

### **Check conformance**

Yes

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### **Check completeness**

Yes

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### **Check stability**

Yes

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### **Check logical consistency**

Yes

## Data characterisation

### **Data characterisation conducted**

Not applicable