

# Post-Authorization Safety Study to Assess the Effectiveness of the Newly Implemented Risk Minimization Measures for Topiramate: HCP and Patient Knowledge and Behavior Survey

**First published:** 06/11/2025

**Last updated:** 06/11/2025

Study

Planned

## Administrative details

### EU PAS number

EUPAS1000000806

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### Study ID

1000000806

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### DARWIN EU® study

No

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### Study countries

 France

 Germany

-  Poland
  -  Spain
  -  Sweden
  -  United Kingdom
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## Study description

Cross-sectional, multinational, and non-interventional on-line survey to assess the impact of the newly implemented Risk Minimization Measures (RMMs) on the knowledge and self-reported behaviour of HCPs involved in prescription, dispensing, and/or consultation of topiramate mono-component products and of Women of Childbearing Potential (WOCBP) treated with topiramate mono-component products for migraine and/or epilepsy in the EU and UK.

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## Study status

Planned

# Research institutions and networks

## Institutions

### Syneos Health

 United Kingdom

**First published:** 23/04/2015

**Last updated:** 06/03/2024

**Institution**

**Non-Pharmaceutical company**

**ENCePP partner**

## Contact details

### **Study institution contact**

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**Study contact**

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### **Primary lead investigator**

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**Primary lead investigator**

### **ORCID number:**

0000-0003-0026-9360

## Study timelines

### **Date when funding contract was signed**

Planned: 20/09/2024

Actual: 20/09/2024

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### **Study start date**

Planned: 18/03/2026

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### **Data analysis start date**

Planned: 17/09/2026

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### **Date of final study report**

Planned: 09/12/2026

## Sources of funding

- Pharmaceutical company and other private sector

## More details on funding

Topiramate Safety Consortium of MAHs, led by Janssen

## Study protocol

[Protocol\\_abstract\\_Topiramate PASS\\_7054980\\_SURVEY.pdf](#) (238.2 KB)

## Regulatory

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

Yes

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

EU RMP category 1 (imposed as condition of marketing authorisation)

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### **Regulatory procedure number**

EMA/H/A-31/1520

## Other study registration identification numbers and links

PCSESPA0008

## Methodological aspects

### Study type

### Study type list

**Study topic:**

Human medicinal product

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**Study type:**

Non-interventional study

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**Scope of the study:**

Assessment of risk minimisation measure implementation or effectiveness

**Data collection methods:**

Primary data collection

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**Study design:**

Cross-sectional, multinational, non-interventional survey among HCPs involved in prescription, dispensing, and/or consultation of topiramate mono-component products and WOCBP ( $\geq 13$  to  $\leq 49$  years of age) exposed to it for migraine and/or epilepsy in 6 European countries (5 EU member states and the UK)

**Main study objective:**

To assess the impact of the newly implemented RMMs on the knowledge and self-reported behavior of HCPs who prescribe topiramate mono-component products and of patients treated with topiramate mono-component products in the EU and UK.

To assess the awareness of:

WOCBP treated with topiramate mono-component products with respect to the receipt of the Educational Materials (EMs).

HCPs with respect to the receipt of/access to Direct Healthcare Professional Communications (DHPC) and EMs for topiramate mono-component products.

To assess the knowledge of:

WOCBP treated with topiramate mono-components products, with respect to risks associated with use of topiramate during pregnancy and measures to prevent exposed pregnancies.

HCPs with respect to risks associated with use of topiramate during pregnancy and measures to prevent pregnancies among WOCBP exposed to topiramate mono-component products.

To assess the self-reported behavior of:

WOCBP treated with topiramate mono-component products, with respect to measures to prevent exposed pregnancies.

HCPs with respect to measures to prevent pregnancies among WOCBP exposed to topiramate mono-component products.

## Study Design

### **Non-interventional study design**

Cross-sectional

## Study drug and medical condition

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

TOPIRAMATE

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### **Anatomical Therapeutic Chemical (ATC) code**

(N03AX11) topiramate

topiramate

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### **Medical condition to be studied**

Epilepsy

Migraine

## Population studied

### **Short description of the study population**

The target population of the survey will be composed of women of childbearing potential (WOCBP) who are treated with topiramate and HCPs grouped into three categories: 1) those who prescribe topiramate, 2) those who dispense topiramate, and 3) those who consult with or see patients receiving topiramate, in the targeted countries.

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### **Age groups**

- **Paediatric Population (< 18 years)**
    - Adolescents (12 to < 18 years)
  - **Adult and elderly population (≥18 years)**
    - Adults (18 to < 46 years)
    - Adults (46 to < 65 years)
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### **Special population of interest**

Women of childbearing potential not using contraception

Women of childbearing potential using contraception

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### **Estimated number of subjects**

500

## Study design details

## **Outcomes**

The HCP and patient surveys will begin with screening questions, followed by items assessing awareness, knowledge, and behavior. For each dimension, the proportion of participants who correctly respond to at least 80% of the relevant questions will be calculated.

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## **Data analysis plan**

Data from patients and HCPs will be analyzed separately, with all analyses being descriptive. The patient dataset will be analyzed by indication, while the HCP will be analyzed by (1) HCP category and (2) medical specialty. The main analysis will be conducted for all countries combined, for all EU countries (excluding the UK), and for each individual country.

Within the HCP and patient datasets, a derived variable in the form of a score summarizing all responses at individual participant level will be created for each of the main dimensions of the survey (awareness, knowledge, and behavior). The individual score will be calculated as the proportion of correct responses to all questions within each domain. Success for each dimension is defined as a score of at least 80%.

For each dimension, the number and proportion of participants who have a score of at least 80% will be calculated. Success for each dimension is defined as having at least 80% of successful participants.

At a study level (i.e., all dimensions combined), the newly implemented RMMs will be considered satisfactory if the behavior outcome and at least one of the two other outcomes (awareness and/or knowledge) are equal to or above the defined threshold of 80% of successful participants.

## **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 source(s), other**

The data source will be on-line questionnaires completed by HCPs and patients (or parents/caregivers) who have passed the screening and who have given consent.

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### **Data sources (types)**

[Other](#)

[Patient surveys](#)

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### **Data sources (types), other**

Surveys from HCPs including general practitioners (GPs), specialists in internal medicine, other specialists managing epilepsy and/or migraine (including neurologists, general pediatricians, neuro-pediatricians), obstetricians/gynecologists (OB/GYN), midwives, and pharmacists

## Use of a Common Data Model (CDM)

### **CDM mapping**

No

## Data quality specifications

**Check conformance**

Unknown

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**Check completeness**

Unknown

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**Check stability**

Unknown

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**Check logical consistency**

Unknown

## Data characterisation

**Data characterisation conducted**

Unknown