# Cariprazine in patients with schizophrenia – real-life data analysis based on the NEAK database (HU-NIS-CARIPRAZINE-01/2024)

First published: 21/12/2024

**Last updated:** 06/03/2025





# Administrative details

EU PAS number	
EUPAS1000000422	
Study ID	
1000000422	
DARWIN EU® study	
No	
Study countries	
Hungary	

## **Study status**

Ongoing

Research institutions and networks

# **Institutions**

# Gedeon Richter

First published: 01/02/2024

Last updated: 01/02/2024

Institution

# Contact details

### **Study institution contact**

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Study contact

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### **Primary lead investigator**

Darko Djuric

**Primary lead investigator** 

# Study timelines

Date when funding contract was signed

Planned: 17/12/2024

Actual: 17/12/2024

### Study start date

Planned: 01/01/2025

Actual: 05/02/2025

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

Planned: 15/05/2025

# Sources of funding

• Pharmaceutical company and other private sector

# Regulatory

Was the study required by a regulatory body?

No

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

Not applicable

# Methodological aspects

Study type

Study type list

**Study topic:** 

Human medicinal product

### Study type:

Non-interventional study

### Scope of the study:

Drug utilisation

### **Data collection methods:**

Secondary use of data

# Study Design

### Non-interventional study design

Cohort

# Study drug and medical condition

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

CARIPRAZINE HYDROCHLORIDE

**OLANZAPINE** 

**ARIPIPRAZOLE** 

**RISPERIDONE** 

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

(N05AX15) cariprazine

cariprazine

(N05AX12) aripiprazole

aripiprazole

(N05AH03) olanzapine

olanzapine (N05AX08) risperidone risperidone

### Medical condition to be studied

Schizophrenia

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The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.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

National Health Insurance Fund (Hungarian acronym: NEAK) database

# **Data sources (types)**

Drug utilisation data

# Use of a Common Data Model (CDM)

# **CDM** mapping

No

# Data quality specifications

# Unknown Check completeness Unknown

# **Check stability**

**Check conformance** 

Unknown

# **Check logical consistency**

Unknown

# Data characterisation

### **Data characterisation conducted**

No