# Cross-Sectional Survey of Rheumatologists and Dermatologists to Assess the Effectiveness of the Baricitinib Additional Risk Minimization Measures in Canada

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# Administrative details

EU PAS number EUPAS1000000438	
<b>Study ID</b> 1000000438	
DARWIN EU® study	
Study countries  Canada	

#### Study description

Survey study of HCPs to assess effectiveness of aRMM.

#### **Study status**

**Planned** 

# Research institutions and networks

# **Institutions**



# Contact details

#### **Study institution contact**

Kristin Meyers meyers\_kristin\_joy@lilly.com

Study contact

meyers\_kristin\_joy@lilly.com

### **Primary lead investigator**

Philippe Assad

Primary lead investigator

# Study timelines

#### Date when funding contract was signed

Planned: 24/09/2024 Actual: 23/07/2024

#### Study start date

Planned: 03/02/2025

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

Planned: 01/04/2026

# Sources of funding

• Pharmaceutical company and other private sector

# More details on funding

Eli Lilly and Company

# Study protocol

LY3009104 B042 NI Canada Protocol Version 1.pdf(754.47 KB)

# Regulatory

Was the study required by a regulatory body?

Yes

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

Non-EU RMP only

# Other study registration identification numbers and links

# Methodological aspects

# Study type

# Study type list

#### Study type:

Non-interventional study

#### Scope of the study:

Assessment of risk minimisation measure implementation or effectiveness

#### **Data collection methods:**

Primary data collection

# Study drug and medical condition

#### Name of medicine

**OLUMIANT** 

#### Name of medicine, other

baricitinib

# Data management

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