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

First published: 14/01/2025

Last updated: 13/02/2025

Study Planned

Administrative details

EU PAS number

EUPAS1000000438

Study ID

1000000438

DARWIN EU® study

No

Study countries

Canada

Study description

Survey study of HCPs to assess effectiveness of aRMM.

Study status

Planned

Research institutions and networks

Institutions

YOLARX Consultants

France

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Institution | **Non-Pharmaceutical company** | **ENCePP partner**

Contact details

Study institution contact

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Study contact

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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

I4V-CA-B042

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

Medicinal product name

OLUMIANT

Medicinal product name, other

baricitinib

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 The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.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

Unknown

Check completeness

Unknown

Check stability

Unknown

Check logical consistency

Unknown

Data characterisation

Data characterisation conducted

No