Real-World Impact of Bimekizumab on Disease Activity in axial spondyloarthritis patients including the EARLY subpopulation (EXPEDITE)

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Administrative details

EU PAS number	
EUPAS1000000519	
Study ID	
1000000519	
DARWIN EU® study	
No	
Study countries	
Austria	
Belgium	
European Union	

France
Germany
Greece
Italy
☐ Netherlands
Poland
Spain
Switzerland
United Kingdom
Study status
Ongoing
Contact details
Study institution contact
UCB Cares UCBCares.global@ucb.com
Study contact
UCBCares.global@ucb.com
Primary lead investigator
UCB Cares
Primary lead investigator

Study timelines

Date when funding contract was signed

Planned: 01/04/2025

Actual: 09/04/2025

Study start date

Planned: 30/05/2025 Actual: 21/05/2025

Date of final study report

Planned: 25/10/2027

Sources of funding

Pharmaceutical company and other private sector

More details on funding

UCB Biopharma SRL

Regulatory

Was the study required by a regulatory body?

No

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

Not applicable

Other study registration identification numbers and links

Internal study ID: SPA002

Methodological aspects

Study type

Human medicinal product	
Study type:	
Non-interventional study	
Study drug and medical condition	
Name of medicine	
BIMZELX	
Study drug International non-proprietary name (INN) or common no BIMEKIZUMAB	ame
Anatomical Therapeutic Chemical (ATC) code	
(L04AC21) bimekizumab	
bimekizumab	
Medical condition to be studied	
Axial spondyloarthritis	
Data management	
ENCePP Seal	
2.133.1 333.	

Study topic:

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 The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.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

Check completeness

Yes

Check stability

Yes

Check logical consistency

Yes

Data characterisation

Data characterisation conducted

No