

Juvenile Idiopathic Arthritis Registry

OM1®

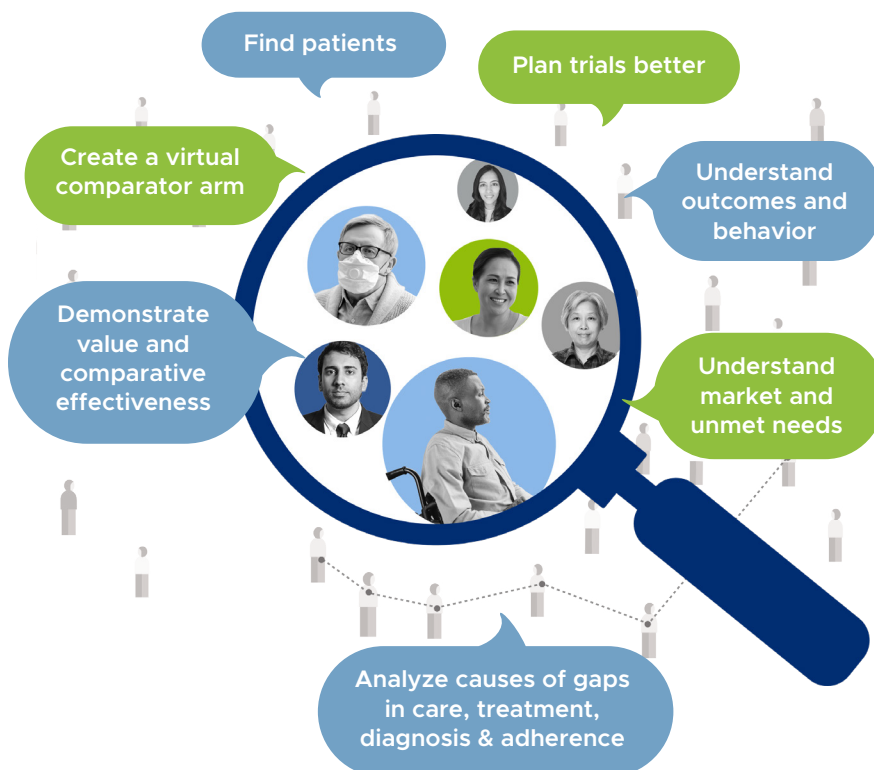
Better understand, compare and predict outcomes

The OM1 Juvenile Idiopathic Arthritis (JIA) registry is a continually updating linked database of more than **4,000** patients prospectively followed with deep clinical, claims, patient reported, laboratory and other data, such as measures of disease activity and treatment response.

The registry provides unique insights into patient journeys, comparative effectiveness, provider behavior and competitive intelligence that is not possible with existing smaller and less complete registries and datasets. The broader OM1 Real-World Data Cloud includes more than **58,500** total JIA patients that can be used to apply models and other advanced techniques aimed at bringing the right treatment to the right patient more quickly.

What do you need to do?

Whether you're focused on improving clinical development, seeking approval or market access, comparing treatment benefit, or trying to determine which prescribers are prescribing your treatment and why, the OM1 JIA Registry can help you rapidly meet your goals:



How we can help

Through our JIA registry, we're delivering faster, more cost-effective access to:

- Clinically Deep Data
- Research Grade Data
- Analytics Ready Data
- Mean follow-up of 6 years

Our JIA Registry products

- Data Licenses
- Data Analytics
- Registry-Based Studies



Learn more and get started

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