

Rheumatoid Arthritis

**Tocilizumab Real-Life Human Factors (RLHFs) Validation Study**

**Trial Status**  
Completed

**Trial Runs In**  
1 Countries

**Trial Identifier**  
NCT02682823 WA29917

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The source of the below information is the publicly available website [ClinicalTrials.gov](https://clinicaltrials.gov). It has been summarised and edited into simpler language.

***Trial Summary:***

This study is designed to evaluate RLHFs concerning administration of the tocilizumab autoinjector AI-1000 G2 in adults with rheumatoid arthritis (RA) who have been receiving subcutaneous (SC) tocilizumab using the commercially available prefilled syringe and needle safety device (PFS-NSD). The study will enroll participants with RA, a subset of whom will be assigned to perform self-injection with the AI-1000 G2. Enrolled caregivers (CGs) and healthcare professionals (HCPs) will administer the AI-1000 G2 injection to the remaining study participants.

**Hoffmann-La Roche**  
Sponsor

**Phase 4**  
Phase

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**NCT02682823 WA29917**  
Trial Identifiers

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***Eligibility Criteria:***

**Gender**  
All

**Age**  
≥ 18 Years

**Healthy Volunteers**  
No

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