

Wollenhaupt J, Lee EB, Curtis JR, et al. Safety and efficacy of tofacitinib for up to 9.5 years in the treatment of rheumatoid arthritis: final results of a global, open-label, long-term extension study. Arthritis Res Ther. 2019;21(1):89.



7,061
PATIENTS
with moderate to severe RA³

>22,800
PY of drug exposure (all doses; as of March 2017)¹

>260,000

PATIENTS
have been prescribed Xeljanz worldwide

(all indications as of June 2019)²

P-XEL-ISR-033

XELJANZ 5 mg BID is the only approved dosage for the treatment of RA. (Xeljanz (tofacitinib) latest approved Israeli prescribing information)
BID=twice daily; LTE=long-term extension; PY=patient-years; QD=once daily; RA=rheumatoid arthritis. 1. Cohen S, et al. Presented at the American College of Rheumatology Annual Scientific Meeting, October 19–24, 2018; Chicago, IL, USA. Abstract 963. 2. Data on file as of June 2019 Pfizer Inc., New York, NY. 3. SB Cohen et al, Long-term Safety of Tofacitinib up to 9.5 Years: A Comprehensive Integrated Analysis of the Rheumatoid Arthritis Clinical Development Program, ACR oct 18, Presentation No. 963.

* Wollenhaupt J, Lee EB, Curtis JR, et al. Safety and efficacy of tofacitinib for up to 9.5 years in the treatment of rheumatoid arthritis: final results of a global, open-label, long-term extension study. Arthritis Res Ther. 2019;21(1):89.

Xeljanz has three indications¹

RA, PsA and UC and has been prescribed for more than 260,000 patients² worldwide



- 1. Xeljanz (tofacitinib) latest approved Israeli prescribing information
 - 2. Pfizer data on file as of June 2019. Pfizer Inc. New York, NY