

## **FACT SHEET**

# Humira

#### **DRUG INFORMATION**

• FDA Approval: 11 total indications. Major disease indications include rheumatoid arthritis (2003), psoriatic arthritis (2005), and Crohn's disease (2007)

Company: AbbVieDrug Type: Biologic

• On the U.S. Market since: 2002

 Generic/Biosimilar approved by FDA: 2016 (Not on the market)

#### **PRICING**

• Annual List Price: \$44,0001

• Price Trend: 100% increase from 2012 to 2018<sup>2</sup>

• Annual Medicare Spending: \$5,339,052,157<sup>3</sup>

• Increase in Average Spend per Medicare Beneficiary (2014-2018): 94%

Annual Medicaid Spending: \$1,642,701,843

• Medicare Spending Rank: 54

#### PATENT LANDSCAPE

• First Patent Filed: 1994

• Total Patent Applications: 257

• Patents Issued: 130

• Filed After FDA Approval: 90%

• Duration of Patent Protection: 39.0 years (Nov 1994 - Nov 2033)

# DELAYED COMPETITION IN THE U.S.

- Humira biosimilars launched in Europe in Oct 2018. Within 6 months, prices dropped by 70%.
- In the U.S., AbbVie negotiated deals with 8 drugmakers to delay the launch of biosimilars until 2023.
- Between 2018 and 2023, an estimated \$77 billion will be spent on Humira in the U.S.

#### **DID YOU KNOW?**

AbbVie has been granted just 6 patents for Humira in Europe. Lower-cost biosimilar versions of the drug have been available in that region since 2018. By contrast, AbbVie's 130 granted U.S. patents will prohibit biosimilars from entering the market until 2023.

### **VISIT I-MAK.ORG/HUMIRA**