

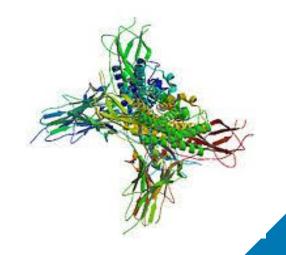


Biosimilars: The Next Frontier PBGH & IBI Webinar Series Event

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Kaiser Permanente (KP) Facts

12.4 million members

8 States and the District of Columbia

714 Medical Offices

39 Hospitals

23,271 Physicians

63,306 Nurses

\$84.5B 2019 Operating Revenue

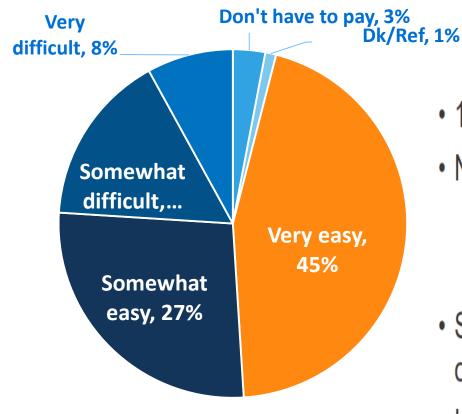
\$14 B 2019 Pharmacy Expenses





Visit kp.org/ataglance

What problem are we trying to solve? 1 in 4 people in the US have a difficult time affording their medicine



- 1 in 4 people have a difficult time affording their medications¹
- Newer, innovative therapies cost ~\$500K-\$2M per patient
 - CAR-T therapy
 - Gene therapy
- Specialty drugs now account for more than half of drug budgets in many organizations
- How can we afford to provide these therapies?

Source: Kaiser Family Foundation Health Tracking Poll (conducted Aug 6-11, 2015)



Kaiser Permanente Process for Biosimilars



Endorsements

 Evidence presented to key stakeholders for assessment & approval

Contracts

 Contracting negotiates and secures contract

Formulary

 Biosimilar is vetted through P&T

Education

Drug Use
 Management
 communicates and
 educates field

Operations

- eHR, ordering systems point to biosimilar as preferred
- Implement inventory strategy

Surveillance

- Track utilization
- Re-educate where needed
- Monitor for safety/efficacy

Note: Kaiser Permanente is an integrated health system, so some processes may be unique

What do patients & clinicians want to know about biosimilars?

Is it just as effective?



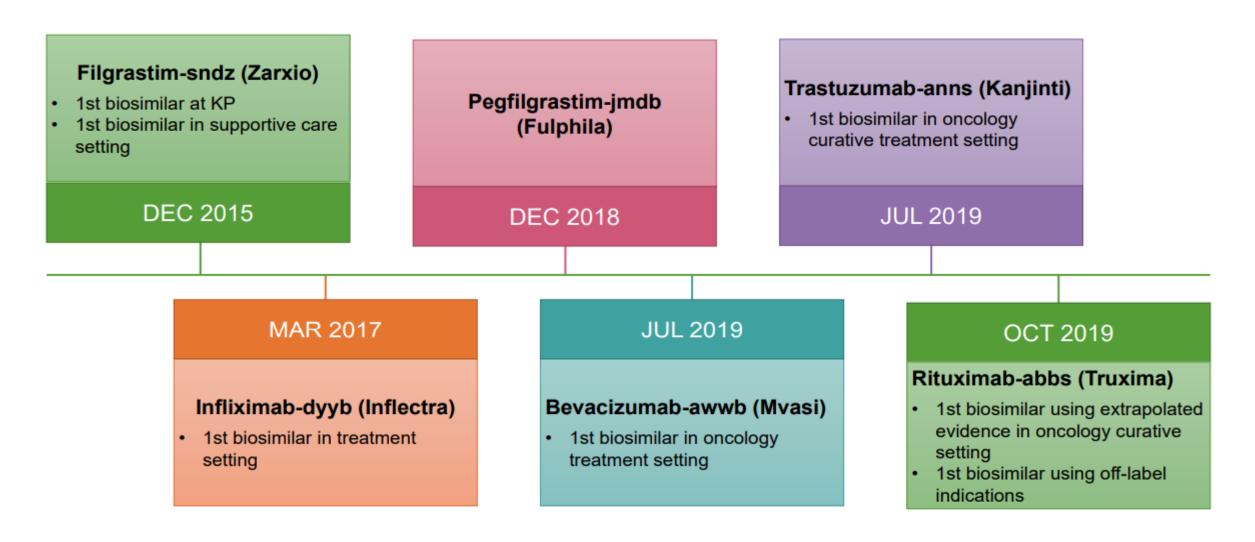
Is it just as safe?

Are the presentations the same?

Are there differences in patient-convenience?

It is more cost-effective?

Kaiser Permanente Journey with Biosimilars



Currently there are 28 biosimilars approved by the FDA, with only 12 on the market

Kaiser Permanente Success With Biosimilars

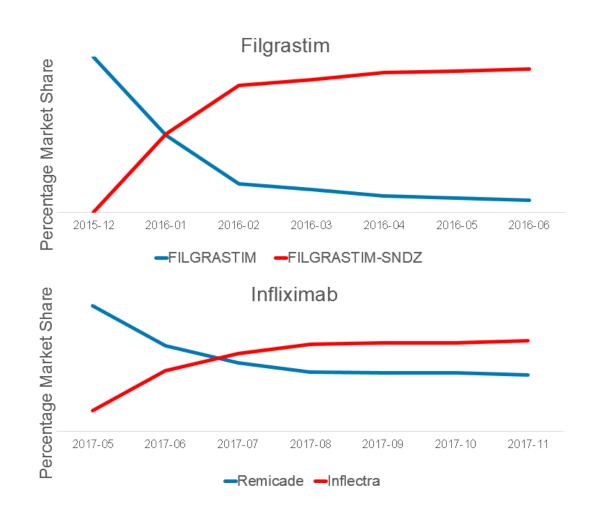
Currently using 6 biosimilars

80-95% Biosimilar market share

Estimated savings: ~\$200M since inception

Higher/faster success rates with:

- Supportive care vs treatment
- Palliative vs curative
- Increasing familiarity with biosimilar concept





The Permanente Federation

PERMANENTE MEDICINE. The Impact Of Biosimilar Competition In Europe

	Price per TD 2016 / Year before Biosimilar entrance
EPO	Total market
Portugal	-66%
Slovakia	-53%
Norway	-51%
HGH	
Finland	-52%
Poland	-42%
Norway	-37%
Fertility	
Denmark	-24%
Spain	-14%
Sweden	-10%

	Price per TD 2016 / Year before Biosimilar entrance
G-CSF	Total market
Romania	-62%
Slovakia	-61%
Slovenia	-57%
Anti-TNF	
Sweden	-39%
Norway	-32%
Denmark	-24%
Insulins	
Finland	-18%
France	-5%
Ireland	-3%

Anti-TNF	Volume TD 2016/ Year before Biosimilar entrance
Bulgaria	190%
Slovakia	93%
Sweden	74%
Portugal	63%
Czech	59%
EPO	
Poland	237%
Greece	196%
Italy	39%
Czech	36%
Bulgaria	36%

The entrance of biosimilars increases price competition

Competition drives down price

Lower prices increase patient access

The Impact of Biosimilar Competition in Europe - European Commission

Conclusions

- The US Biosimilars Market is at a critical juncture as uptake is improving but slower compared to Europe due to patent litigation amongst pharmaceutical companies, pay for delay, restricted access by PBMs/insurers, reimbursement mechanisms, and a lack of physician and patient education regarding availability and efficacy, and due to the FDA's interchangeability requirement
- This is despite the fact that there are substantial financial savings (\$54 B from 2017-2026) per the Rand Corporation) to be gained through the use of biosimilars and improved access for patients to these medications
- Biosimilar adoption provides a solution to the drug pricing and spending debate like generics did for brand name drugs
- Additional policies need to be considered to help uptake and viability of the biosimilars market in the U.S.



Questions?

