## **Operationalizing Biosimilars Best Practices: 3 Practical Steps Toward Implementation**

Biosimilars have layered additional complexities into the traditional evaluation process used by Pharmacy and Therapeutic (P&T) committees in considering a new product for formulary inclusion. To ensure efficient implementation, a thoughtful assessment of the strategies and tactics related to an institution's operational considerations is required.<sup>1,2</sup>

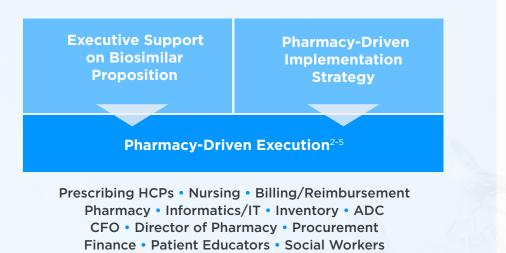
## Strategic Considerations for P&T Evaluation<sup>1,2</sup>



#### **Pharmacy-Driven Implementation Strategy**

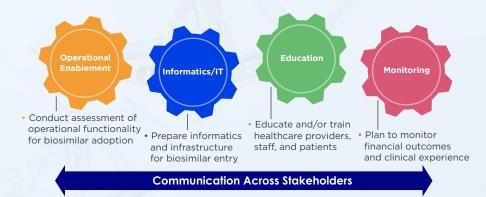
Supported by defined timelines, roles and responsibilities, and necessary resources for execution at a given institution

# Multidisciplinary Collaboration Will Be Essential to Execute the P&T Implementation Strategy<sup>3,4</sup>



**Registration & Scheduling** 

### Four Critical Elements That May Be Included in a Biosimilar Implementation Strategy<sup>2-4,6-9</sup>



The considerations found on the next page may help you develop the implementation strategy that will be most effective at your institution.

## **Example Considerations for Biosimilar Implementation Strategy**

### **1** PREPARATION

#### **Operational Enablement**

- Integrate benefits verification process with electronic health record (EHR) systems to facilitate a quick order process<sup>7</sup>
- Set up biosimilar prior authorizations in the EHR to ensure coverage<sup>7</sup>
- Develop a patient transition plan for how biosimilar transitions will be sequenced (e.g., all at once, phased in by clinic or therapeutic area, existing vs treatment-naive patients)<sup>1,4</sup>
- Identify where the product will be stored and how much should be stocked
- Evaluate what inventory management system modifications need to be made (including automated dispensing devices and software) to integrate the biosimilar and enable accurate identification<sup>2,4</sup>

### **2** IMPLEMENTATION

#### Informatics/IT

- Identify what modifications need to be made to existing order sets and protocols to include biosimilar products<sup>4</sup>
- Plan for converting patient orders (e.g., new vs established patients, single order vs treatment plan, route for co-signature)<sup>1,2,4</sup>
- Determine which EHR functionality/interface can be leveraged to support the organization's biosimilar conversion goals<sup>2,4</sup>

#### Education

- Identify the most critical stakeholders to target for biosimilar education<sup>7,9</sup>
- Determine what education stakeholders will need to prepare for biosimilar adoption<sup>79</sup>
- Select who will be responsible for patient education<sup>5,8</sup>
- Identify which patient education materials will be needed to support biosimilar use<sup>5,8</sup>

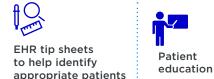
## **3** MONITORING

#### Monitoring

- Determine how the institution will ensure that biosimilar protocol is followed<sup>2,7</sup>
- Identify how financial and drug use evaluation outcomes will be monitored in order to determine if value of the biosimilar was achieved<sup>2,7</sup>

Clear communication of the biosimilar initiative and implementation strategy should be maintained across stakeholders throughout the medication management continuum.<sup>3</sup>

## Pfizer Provides Resources to Help With Seamless Biosimilar Implementation for Your Practice, Hospital, or Health System





EHR biosimilar implementation support tools



Peer-to-peer speaker programs



Emerging potential best practices for implementation and considerations for P&T

#### PfizerBiosimilars.com downloadable resources and educational videos

## For more information, please speak with your Pfizer representative and visit PfizerBiosimilars.com

References: 1. Soefje S. Considerations for adding biosimilars to formulary. https://www.pppmag.com/article/2368. Accessed February 24, 2021. 2. Griffith N, McBride A, Stevenson JG, Green L. Formulary selection criteria for biosimilars: considerations for US health-system pharmacists. *Hosp Pharm.* 2014;49(9):813-825. 3. ASHP Expert Panel on Formulary Management. ASHP guidelines on the pharmacy and therapeutices committee and the formulary system. *Am J Health Syst Pharm.* 2008;65(13):1272-1283. 4. Ventola CL. Evaluation of biosimilars for formulary inclusion: factors for consideration by P&T committees. *P T.* 2015;40(10):680-689. 5. Lucio SD, Stevenson JG, Hoffman JM. Biosimilars: Implications for health-system pharmacists. *Am J Health Syst Pharm.* 2013;70(22):2004-2017. 6. Rumore MM, Vogenberg FR. Biosimilars: still not quite ready for prime time. *P T.* 2016;41(6):366-375. 7. AMCP Partnership Forum: biosimilars: eurrent perspectives in immune-mediated inflammatory diseases. *Expert Opin Biol Ther.* 2019;19(10):1001-1014. 9. Oskouei ST. Following the biosimilar breadcrumbs: when health systems and manufacturers approach forks in the road. *J Manag Care Spec Pharm.* 2017;23(12):1245-1248.

