

FDA approvals and expedited pathways used -- New Molecular Entities (NMEs)

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Anthem Blue Cross and Blue Shield (Anthem) reviews the activities of the Food and Drug Administration (FDA)'s approval of drugs and biologics on a regular basis to understand the potential effects for both our providers and members.

The FDA approves new drugs/biologics using various pathways of approval. Recent studies on the effectiveness of drugs/biologics going through these different FDA pathways illustrates the importance of clinicians being aware of the clinical data behind a drug or biologic approval in making informed decisions.

Here is a list of the approval pathways the FDA uses for drugs/biologics:

- **Standard Review** – The Standard review process follows well-established paths to make sure drugs/biologics are safe and effective when they reach the public. From concept to approval and beyond, FDA performs these steps: reviews research data and information about drugs and biologics before they become available to the public; watches for problems once drugs and biologics are available to the public; monitors drug/biologic information and advertising; and protects drug/biologic quality. *To learn more about the Standard Review process, [click here](#).*
- **Fast Track** – Fast Track is a process designed to facilitate the development, and expedite the review of drugs/biologics to treat serious conditions and fill an unmet medical need. *To learn more about the Fast Track process, [click here](#).*
- **Priority Review** – A Priority Review designation means FDA's goal is to take action on an application within 6 months. *To learn more about the Priority Review process, [click here](#).*
- **Breakthrough Therapy** – A process designed to expedite the development and review of drugs/biologics which may demonstrate substantial improvement over available therapy. *To learn more about the Breakthrough Therapy process, [click here](#).*

- **Orphan Review** – Orphan Review is the evaluation and development of drugs/biologics that demonstrate promise for the diagnosis and/or treatment of rare diseases or conditions. [To learn more about the Orphan Review process, click here.](#)
- **Accelerated Approval** – These regulations allowed drugs/biologics for serious conditions that filled an unmet medical need to be approved based on a surrogate endpoint. [To learn more about the Accelerated Approval process, click here.](#)

New Molecular Entities Approvals: January - August 2020

Certain drugs/biologics are classified as new molecular entities (“NMEs”) for purposes of FDA review. Many of these products contain active ingredients that have not been approved by FDA previously, either as a single ingredient drug or as part of a combination product; these products frequently provide important new therapies for patients.

Anthem reviews the FDA-approved NMEs on a regular basis. To facilitate the decision-making process, we are providing a list of NMEs approved from January to August 2020 along with the FDA approval pathway utilized.

Generic Name	Trade Name	Standard Review	Fast Track	Priority Review	Break-through Therapy	Orphan Review	Accelerated Approval
Abametapir	Xeglyze	X					
Amisulpride	Barhemys	X					
Avapritinib	Ayvakit		X	X	X	X	
Belantamab mafodotin	Blenrep			X	X	X	X
Bempedoic acid	Nexletol	X					
Brexucabtagene autoleucel	Tecartus			X	X	X	X
Capmatinib	Tabrecta			X	X	X	X
Decitabine/ cedazuridine	Inqovi			X		X	
Eptinezumab-jjmr	Vyepti	X					
Fostemsavir	Rukobia		X	X	X		
Inebilizumab	Uplizna	X			X	X	
Isatuximab	Sarclisa	X				X	
Lurbinectedin	Zepzelca			X		X	X
Nifurtimox	Lampit			X		X	X
Oliceridine	Olinvyk	X	X				
Opicapone	Ongentys	X					
Osilodrostat	Isturisa	X				X	
Ozanimod	Zeposia	X					
Peanut (Arachis hypogaea) allergen powder-dnfp	Palforzia	X	X		X		

Generic Name	Trade Name	Standard Review	Fast Track	Priority Review	Break-through Therapy	Orphan Review	Accelerated Approval
Pemigatinib	Pemazyre			X	X	X	X
Remimazolam	Byfavo	X					
Rimegepant	Nurtec ODT			X			
Risdiplam	Evrysdi		X	X	X	X	
Ripretinib	Qinlock		X	X	X	X	
Sacituzumab-hziy	Trodelvy		X	X	X	X	X
Selpercatinib	Retevmo			X	X	X	X
Selumetinib	Koselugo		X	X	X	X	
Tafasitamab	Monjuvi	X	X		X	X	X
Tazemetostat	Tazverik			X		X	X
Teprotumumab-trbw	Tepezza		X	X	X	X	
Triheptanoin	Dojolvi	X	X			X	
Tucatinib	Tukysa		X	X	X	X	
Viltolarsen	Viltepso		X	X		X	X

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