

Other Products

PRODUCT	INDICATION OR PROPOSED INDICATION	APPROVED/FILED*		
		U.S.	EU	JAPAN
Myfembree (relugolix, estradiol, and norethindrone acetate) ^(a)	Moderate to severe pain associated with endometriosis	Approved August 2022		
Ngenla (somatrogen) ^(b)	Pediatric growth hormone deficiency	Approved June 2023	Approved February 2022	Approved January 2022
Pevnar 20/Apexxnar (Vaccine)	Active immunization to prevent pneumonia, invasive disease and otitis media caused by Streptococcus pneumoniae (adults)	Approved June 2021	Approved February 2022	
	Active immunization to prevent pneumonia, invasive disease and otitis media caused by Streptococcus pneumoniae (pediatric)	Approved April 2023		Filed March 2023
TicoVac (Vaccine)	Active immunization to prevent tick-borne encephalitis disease	Approved August 2021		Filed March 2023
Paxlovid^(c) (nirmatrelvir and ritonavir)	COVID-19 in high-risk adults	Approved May 2023	Approved February 2023	Approved February 2022
Nurtec ODT/Vydura (rimegepant)	Acute treatment of migraine with or without aura (adults)	Approved February 2020	Approved April 2022	
	Prevention of episodic migraine (adults)	Approved May 2021	Approved April 2022	
Litfulo/Ritfulo (ritlecitinib)	Alopecia areata	Approved June 2023	Filed September 2022	Approved June 2023
Zavzpret (zavegepant) (intranasal)	Acute treatment of migraine with or without aura (adults)	Approved March 2023		
PF-06886992 (Vaccine)	Active immunization to prevent serogroups ABCWY meningococcal infections (adolescent and young adults)	Filed December 2022	Filed June 2023	
Abrysvo (Vaccine)	Active immunization to prevent respiratory syncytial virus infection (maternal)	Filed February 2023	Filed January 2023	Filed February 2023
	Active immunization to prevent respiratory syncytial virus infection (older adults)	Approved May 2023	Filed January 2023	Filed May 2023
etrasimod	Ulcerative colitis (moderately to severely active)	Filed December 2022	Filed November 2022	
Braftovi (encorafenib) and Mektovi (binimetinib)	BRAF ^{V600E} -mutant metastatic non-small cell lung cancer	Filed April 2023		
elranatamab (PF-06863135)	Multiple myeloma triple-class relapsed/refractory	Filed February 2023	Filed February 2023	Filed June 2023
Talzenna (talazoparib)	Combination with Xtandi (enzalutamide) for adult patients with homologous recombination repair (HRR) gene-mutated mCRPC ^(d)	Approved June 2023	Filed February 2023	
fidanacogene elaparovvec (PF-06838435) ^(e)	Hemophilia B	Filed June 2023	Filed June 2023	

* For the U.S., the filing date is the date on which the FDA accepted our submission. For the EU, the filing date is the date on which the EMA validated our submission.

^(a) Being developed in collaboration with Sumitomo Pharma America, Inc. (formerly known as Myovant Sciences Ltd.)

^(b) Being developed in collaboration with OPKO.

^(c) Previously authorized under EUA in the U.S. (December 2021) and approved by the FDA in high-risk adults (May 2023). Remains under EUA for children (12-18 years of age; >88lbs) in the U.S.

^(d) Listed patient population applies to U.S. only. Patient population in the filed application in the EU is an all-comers population in men with mCRPC.

^(e) Being developed in collaboration with Spark Therapeutics, Inc.

In China, the following products received regulatory approvals in the last twelve months: Xeljanz for the treatment of adult patients with active psoriatic arthritis in October 2022; and Prevenar 13 in infants and children aged 6 weeks to 15 months, in April 2023.