

**SELECTED PHARMACEUTICALS IN LATE STAGE U.S. AND E.U. DEVELOPMENT OR REGISTRATION as of 7/16/20**

Therapeutic Area	Product Name	Indication Sought	U.S. Development Stage	E.U. Development Stage	
Cardiovascular and Metabolism	XARELTO® (rivaroxaban)	Infrainguinal revascularized PAD (2) Pediatric VTE (2)	Phase III Phase III		
	INVOKANA® (canagliflozin) aprocitentan	Treatment of Diabetic Kidney Disease in adults with T2DM (2) Difficult to treat hypertension (2)	Approved 9/19 Phase III	Approved 6/20 Phase III	
Immunology	TREMFYA® (guselkumab)	Psoriatic arthritis Crohn's Disease Pediatric Psoriasis UC Monotherapy	Approved 7/20 Phase IIB/III Phase III Phase IIB/III	Filed 10/19 Phase IIB/III Phase III Phase IIB/III	
	SIMPONI® SIMPONI ARIA®	Pediatric Ulcerative Colitis Polyarticular Juvenile Idiopathic Arthritis (pJIA) Juvenile Psoriatic Arthritis (jPsA)	Phase III Filed 3/20 Filed 3/20	Phase III Phase III	
	STELARA® (ustekinumab)	Pediatric Psoriasis (6-11 year-olds)	Filed 9/19	Approved 1/20	
Infectious Diseases and Vaccines	Rilpivirine Long Acting nanosuspension for injection	HIV Infection	Complete Response 12/19	Filed 7/19	
	pimodivir (JNJ-3872) VAC52150 VAC89220	Influenza A (2) Monovalent Ebola Virus Vaccine HIV Px Vaccine	Phase III Phase III Phase III	Phase III Approved 7/20 Phase III	
	SPRAVATO™ (esketamine) ponesimod paliperidone palmitate 6 month long-acting injectable	Major Depressive Disorder with Active Suicidal Ideation and Intent Monotherapy for multiple sclerosis Treatment of Schizophrenia	Filed 10/19 Filed 3/20 Phase III	Filed 1/20 Filed 3/20 Phase III	
Oncology	DARZALEX®	Relapsed Refractory MM 1+ prior lines w/PomDex (2) Relapsed Refractory multiple myeloma w/carfilzomib/dex (Amgen sponsored) (2) Amyloidosis (AMY 3001) (2) Smoldering multiple myeloma (SMM3001) (2) Frontline multiple myeloma transplant ineligible in combination w/ bortezomib, lenalidomide and dexamethasone (MMY3019, CEPHEUS) (2) Frontline multiple myeloma transplant eligible in combination w/ bortezomib, lenalidomide and dexamethasone (MMY3014, PERSEUS) (2)	Phase III Filed 2/20 Phase III Phase III Phase III Phase III	Phase III Phase III Phase III Phase III	
	IMBRUVICA® (ibrutinib)	Subcutaneous Formulation in patients with relapsed or refractory multiple myeloma (MMY3012, COLUMBA) (2) Treatment naive patients with Mantle Cell Lymphoma in combination with Bendamustine and Rituximab (randomized study MCL-3002) (2) Relapsed/refractory patients with Indolent Non-Hodgkins Lymphoma in combination with Bendamustine and Rituximab or R-CHOP; (randomized study FLR-3001) (2) Frontline Chronic Lymphocytic Leukemia in combination with Rituximab (Young and Fit) (ECOG 1912) (2)	Approved 5/20 Phase III Phase III Approved 4/20	Approved 6/20 Phase III Phase III Filed 1/20	
		Frontline Chronic Lymphocytic Leukemia in combination with venetoclax (fixed duration) (GLOW) (2) Relapsed/refractory patients with Mantle Cell Lymphoma in combination with venetoclax (PCYC-1143) (2)	Phase III Phase III	Phase III Phase III	
		ERLEADA® (apalutamide)	Localized prostate cancer High risk prostate cancer (PROTEUS)	Phase III Phase III	Phase III Phase III
		apalutamide/ abiraterone acetate BALVERSA™ (erdafitinib)	Prostate Cancer metastatic castration resistant chemotherapy naïve Urothelial cancer (2) Tumor Agnostic	Phase III Phase III Approved 4/19 Phase IIB	Phase III Phase III Phase III Phase IIB
	niraparib / abiraterone acetate	L1 Prostate cancer metastatic castration-resistant in combination with abiraterone acetate and Prednisone	Phase III	Phase III	
	Ciltacabtagene Autoleucl (BCMA CAR-T) Amivantamab (EGFR/cMET)	Relapsed refractory multiple myeloma (2) Non Small Cell Lung Cancer	Phase II Phase II	Phase II Phase II	
	Pulmonary Hypertension	OPSUMIT®	Fontan-palliated in adolescent (>12 years old) and adult patients Pediatric pulmonary arterial hypertension	Phase III Phase III	Phase III Phase III
		UPTRAVI®	Chronic thromboembolic pulmonary hypertension Pulmonary arterial hypertension IV	Complete Response 1/19 Phase III	Submission Withdrawn 11/19
			Chronic thromboembolic pulmonary hypertension Pediatric pulmonary arterial hypertension	Phase III Phase III	Phase III
Macitentan w/tadalafil FDC	Pulmonary arterial hypertension	Phase III	Phase III		

\* This information is accurate as of the date hereof to the best of the Company's knowledge. Johnson & Johnson assumes no obligation to update this information.

(2) INVOKANA licensed from Mitsubishi Tanabe Pharma Corporation; Long acting HIV injectable treatment regimen of rilpivirine and cabotegravir developed in collaboration with ViiV Healthcare Ltd; aprocitentan developed in collaboration with Idorsia; XARELTO co-developed with Bayer HealthCare; paliperidone palmitate includes technology licensed from Alkermes, Inc.; DARZALEX licensed from Genmab A/S; IMBRUVICA developed in collaboration with Pharmacyclics, LLC, an AbbVie company Pimodivir (JNJ-3872) licensed from Vertex Pharmaceuticals, Inc.; BALVERSA discovered in collaboration with Astex Pharmaceuticals, Inc.; Niraparib licensed from Tesaro; BCMA CAR-T licensed from Legend Biotech; Monovalent Ebola Vaccine developed in collaboration with Bavarian Nordic A/S, and MVA-BN-Filo® is licensed-in from Bavarian Nordic A/S.



**Janssen Pharmaceutical Companies of Johnson & Johnson**  
**Selected NME Pharmaceutical Pipeline - Recent Approvals/Potential Filings\***  
 Selective Highlights as of July 16, 2020

NMEs APPROVED 2019 & 2020	NMEs IN REGISTRATION	NME PLANNED FILINGS 2020-2023*	
Oncology	Neuroscience	Oncology	Neuroscience
<p><b>ERLEADA® (apalutamide) (EU)</b> Non-metastatic prostate cancer</p> <p><b>BALVERSA™ (erdafitinib) (US)</b> Urothelial cancer</p>	<p><b>Ponesimod</b> Monotherapy for multiple sclerosis</p>	<p><b>Niraparib</b> Metastatic castration-resistant prostate cancer (mCRPC)</p> <p><b>Ciltacabtagene Autoleucel (BCMA CAR-T)</b> Multiple Myeloma</p> <p><b>BALVERSA™ (erdafitinib) (EU)</b> Urothelial cancer</p> <p><b>Cusatuzumab</b> Acute myeloid leukemia</p> <p><b>Amivantamab (EGFR/cMET)</b> Non Small Cell Lung Cancer</p> <p><b>JNJ-7957 (BCMA/CD3), JNJ-7564 (GPC5D/CD3)</b> Multiple myeloma</p>	<p><b>Seltorexant</b> Adjunctive treatment for major depressive disorder with insomnia symptoms</p>
			Infectious Diseases & Vaccines
			<p><b>Pimodivir</b> Influenza A</p> <p><b>RSV Sr vaccine</b> RSV vaccine</p>
			Cardiovascular & Metabolism
			<p><b>RPGR AAV Gene Therapy</b> X-linked retinitis pigmentosa</p>
			<p><b>Aprocitanan</b> Difficult to treat hypertension</p>
Neuroscience		Immunology	
<p><b>SPRAVATO™ (Esketamine)</b> Treatment resistant depression</p>		<p><b>JNJ-4500 Anti-NKG2D</b> Crohn's disease</p>	
Infectious Diseases & Vaccines			
<p><b>Monovalent Ebola Vaccine (EU)</b> Ebola virus Disease</p>			

\*Filings/approvals assumed to be in the US and EU unless otherwise noted. This information is accurate as of the date hereof to the best of the Company's knowledge. Johnson & Johnson assumes no obligation to update this information. Pimodivir (JNJ-3872) licensed from Vertex Pharmaceuticals, Inc.; Niraparib licensed from Tesaro; BALVERSA discovered in collaboration with Astex Pharmaceuticals, Inc.; BCMA CAR-T licensed from Legend Biotech; Cusatuzumab licensed from Argenx; Lazertinib licensed from Yuhan Corporation; JNJ-4500 Anti-NKG2D licensed from Novo Nordisk; Retinal assets (Achromatopsia: AAV-CNGA3, AAV-CNGB3) and (X-Linked Retinitis Pigmentosa: AAV-RPGR) licensed from MeiraGTx; Seltorexant being developed in collaboration with Minerva Neurosciences; DUOBODY platform licensed from Genmab relates to several bispecific antibody programs (JNJ-6372, JNJ-7957, JNJ-7564); Monovalent Ebola Vaccine developed in collaboration with Bavarian Nordic A/S, and MVA-BN-Filo® is licensed-in from Bavarian Nordic A/S.