

**SELECTED PHARMACEUTICALS IN LATE STAGE U.S. AND E.U. DEVELOPMENT OR REGISTRATION as of 10/15/19**

Therapeutic Area	Product Name	Indication Sought	U.S. Development Stage	E.U. Development Stage
Cardiovascular and Metabolism	XARELTO® (rivaroxaban)	Prevention of Symptomatic VTE and VTE-related death in high-risk, medically ill pts (2) VTE prophylaxis in ambulatory cancer patients receiving chemo at high-risk for VTE (2) Infringuinal revascularized PAD (2) Pediatric VTE (2)	Approved 10/19 Phase III Phase III Phase III	
	INVOKANA® (canagliflozin) apocritentan	Treatment of Diabetic Nephropathy in adults with T2DM (2) Difficult to treat hypertension (2)	Approved 9/19 Phase III	Filed 7/19 Phase III
Immunology	TREMFYA® (guselkumab)	Psoriatic arthritis Crohn's Disease Pediatric Psoriasis	Filed 9/19 Phase IIB/III Phase III	Phase III Phase IIB/III Phase III
	SIMPONI® STELARA® (ustekinumab)	Pediatric Ulcerative Colitis Pediatric Psoriasis (6-11 year olds) Ulcerative Colitis Lupus	Phase III Filed 9/19 Filed 12/18 Phase III	Phase III Filed 6/19 Approved 9/19 Phase III
Infectious Diseases and Vaccines	Rilpivirine Long Acting nanosuspension for injection pimodivir (JNJ-3872) VAC52150	HIV Infection  Influenza A (2) Monovalent Ebola Virus Vaccine	Filed 4/19  Phase III Phase III	Filed 7/19  Phase III Phase III
	SPRAVATO™ (esketamine)  ponesimod paliperidone palmitate 6 month long-acting injectable	Treatment-resistant depression Major depressive disorder who have active suicidal ideation with intent Monotherapy for multiple sclerosis Treatment of Schizophrenia	Approved 3/19 Filed 10/19 Phase III Phase III	Filed 10/18 Phase III Phase III Phase III
Oncology	DARZALEX®	Frontline multiple myeloma transplant eligible in combination with bortezomib, thalidomide and dexamethasone (MMY 3006, CASSIOPEIA) (2)	Approved 9/19	Filed 3/19
		Frontline multiple myeloma transplant ineligible patients in combination with lenalidomide and low dose dexamethasone (MMY3008, MAIA) (2)	Approved 6/19	Filed 3/19
		Relapsed Refractory MM 1+ prior lines w/PomDex (2)	Phase III	Phase III
		Relapsed Refractory multiple myeloma w/carfilzomib/dex (Amgen sponsored) (2)	Phase III	
		Amyloidosis (AMY 3001) (2)	Phase III	Phase III
		Subcutaneous Formulation in patients with relapsed or refractory multiple myeloma (MMY3012, COLUMBA) (2)	Filed 7/19	Filed 7/19
		Smoldering multiple myeloma (SMM3001) (2)	Phase III	Phase III
	IMBRUVICA® (ibrutinib)	Frontline multiple myeloma transplant ineligible in combination w/ bortezomib, lenalidomide and dexamethasone (MMY3019, CEPHEUS) (2)	Phase III	Phase III
		Frontline multiple myeloma transplant eligible in combination w/ bortezomib, lenalidomide and dexamethasone (MMY3014, PERSEUS) (2)	Phase III	Phase III
		Treatment naïve patients with Mantle Cell Lymphoma in combination with Bendamustine and Rituximab (randomized study MCL-3002) (2)	Phase III	Phase III
		Relapsed/refractory patients with Indolant Non-Hodgkins Lymphoma in combination with Bendamustine and Rituximab or R-CHOP; (randomized study FLR-3001) (2)	Phase III	Phase III
		Previously untreated and relapsed/refractory patients with Waldenstrom's Macroglobulinemia in combination with Rituximab (PCYC-1127) (2)	Approved 8/18	Approved 8/19
ERLEADA® (apalutamide)	Frontline Chronic Lymphocytic Leukemia in combination with Rituximab (Young and Fit) (ECOG 1912) (2)	Phase III	Phase III	
	Frontline Chronic Lymphocytic Leukemia in combination with obinutuzumab (PCYC-1130) (2)	Approved 1/19	Approved 8/19	
	Frontline Chronic Lymphocytic Leukemia in combination with venetoclax (fixed duration) (GLOW) (2)	Phase III	Phase III	
	Frontline Chronic graft-versus-host-disease (cGVHD) (2)	Phase III	Phase III	
	Relapsed/refractory patients with Mantle Cell Lymphoma in combination with venetoclax (PCYC-1143) (2)	Phase III	Phase III	
apalutamide/ abiraterone acetate BALVERSA™ (erdafitinib) niraparib (PARP inhibitor) niraparib / abiraterone acetate BCMA CAR-T	Metastatic castrate sensitive prostate cancer (TITAN)	Approved 9/19	Filed 06/19	
	Localized prostate cancer	Phase III	Phase III	
	High risk prostate cancer (PROTEUS)	Phase III	Phase III	
	Prostate Cancer metastatic castration resistant chemotherapy naïve	Phase III	Phase III	
	Urothelial cancer (2)	Approved 4/19	Phase III	
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 Chronic thromboembolic pulmonary hypertension	Complete Response 1/19 Phase III Phase III	Filed 8/18 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; apocritentan 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 Pharmacocyclics, 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;

**Janssen Pharmaceutical Companies of Johnson & Johnson**  
**Selected NME Pharmaceutical Pipeline - Recent Approvals/Potential Filings\***  
 Selective Highlights as of October 15, 2019

NMEs APPROVED 2019	NMEs IN REGISTRATION	NME PLANNED FILINGS 2019-2023*	
Oncology	Neuroscience	Oncology	Neuroscience
<b>ERLEADA® (apalutamide) (EU)</b> Non-metastatic prostate cancer  <b>BALVERSA™ (erdafitinib) (US)</b> Urothelial cancer	<b>SPRAVATO™ (Esketamine) (EU)</b> Treatment resistant depression	<b>Niraparib</b> Metastatic castration-resistant prostate cancer (mCRPC)  <b>BCMA CAR-T</b> Multiple Myeloma  <b>BALVERSA™ (erdafitinib) (EU)</b> Urothelial cancer  <b>Lazertinib</b> Non Small Cell Lung Cancer  <b>Cusatuzumab</b> Acute myeloid leukemia  <b>JNJ-6372 (EGFR/cMET)</b> Solid Tumor  <b>JNJ-7957 (BCMA/CD3), JNJ-7564 (GPRC5D/CD3)</b> Multiple myeloma	<b>Ponesimod</b> Monotherapy for multiple sclerosis  <b>Seltorexant</b> Adjunctive treatment for major depressive disorder
			Infectious Diseases & Vaccines
			<b>Pimodivir</b> Influenza A  <b>RSV Sr vaccine</b> RSV vaccine
			Cardiovascular & Metabolism
			<b>CNGB3 and CNGA3 AAV Gene Therapy</b> Achromatopsia  <b>RPGR AAV Gene Therapy</b> X-linked retinitis pigmentosa  <b>Aprocitentan</b> Difficult to treat hypertension
		Immunology	
		<b>JNJ-4500 Anti-NKG2D</b> Crohn's disease	

\*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 Argencx; 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)