

# OUR PIPELINE IS POISED TO DELIVER NOW AND IN THE FUTURE

	WAVE 1 <sup>1</sup>				CLINICAL-STAGE NMEs	WAVE 2 <sup>2</sup>				Research engine is targeting 10-12 IND filings in FY21
TARGET APPROVAL	FY21	FY22	FY23	FY24	FY25 and Beyond					
<b>ONCOLOGY</b>	 <b>mobocertinib</b> 2L NSCLC with EGFR exon 20 insertion mutation <sup>3</sup>	 <b>pevonedistat</b> HR-MDS	 <b>mobocertinib</b> 1L NSCLC with EGFR exon 20 insertion mutation <sup>3</sup>	 <b>pevonedistat</b> Unfit AML <sup>5</sup>	 <b>TAK-981</b> Multiple cancers	 <b>TAK-676</b> Solid tumors	 <b>TAK-252</b> Solid tumors	Multiple cancers	 <b>TAK-102</b> Multiple cancers	
<b>RARE GENETICS &amp; HEMATOLOGY</b>	 <b>maribavir</b> R/R CMV infect. in transplant	 <b>TAK-609</b> Hunter CNS (IT) <sup>6</sup>	 <b>maribavir</b> 1L CMV infect. in HSCT	 <b>TAK-611</b> MLD (IT)	 <b>TAK-573</b> R/R MM	 <b>TAK-605</b> Multiple cancers	 <b>TAK-186</b> EGFR Solid Tumor	 <b>TAK-940</b> CD19+ hematologic malignancies		
<b>NEUROSCIENCE</b>			 <b>soticlestat</b> DS	 <b>Orexin 2R-ag</b> (TAK-994/TAK-925) Narcolepsy T1	 <b>Orexin 2R-ag</b> (TAK-994/TAK-861/TAK-925) NT2, IH, Additional Indications	 <b>TAK-831</b> <sup>7</sup> CIAS NS	 <b>TAK-653</b> <sup>7</sup> TRD			
<b>GASTRO-ENTEROLOGY</b>	 <b>Eohilia</b> <sup>4</sup> EoE Approval date TBD		 <b>soticlestat</b> LGS		 <b>TAK-071</b> Parkinson's Disease	 <b>TAK-341</b> Parkinson's Disease	 <b>TAK-041</b> <sup>7</sup> Anhedonia in MDD			
<b>VACCINES</b>	 <b>TAK-003</b> Dengue Vaccine	 <b>TAK-019</b> Novavax COVID-19 Vaccine (JP)			 <b>TAK-999</b> AAT Liver Disease	 <b>TAK-671</b> Acute Pancreatitis	 <b>TAK-062</b> Celiac Disease	 <b>TAK-101</b> Celiac Disease	 <b>sibofimloc</b> Crohn's Disease (post-op and ileitis)	 <b>TAK-510</b> Nausea & vomiting
		 <b>TAK-919</b> Moderna COVID-19 Vaccine (JP)			 <b>TAK-906</b> Gastroparesis	 <b>TAK-954</b> POGD	 <b>TAK-951</b> Nausea & vomiting	 <b>TAK-039</b> Hepatic encephalopathy		
					 <b>TAK-426</b> Zika Vaccine	 <b>TAK-214</b> Norovirus Vaccine				

● Breakthrough and/or Fast Track Designations

● China Breakthrough and/or Japan SAKIGAKE Designation

Orphan potential in at least one indication  
 Orexin franchise

Additions in Q4: TAK-861, TAK-186, TAK-510  
 COVID-19 Vaccines

1. Projected approval dates depend on data read-outs; some WAVE 1 target approval dates assume accelerated approval  
 2. Certain WAVE 2 programs may be accelerated into WAVE 1 depending on future data read outs  
 3. Approval date assumes filing on Phase 2 data  
 4. In active discussions with the FDA. Projected approval subject to outcome of discussions  
 5. COVID-19 related shift in enrollment now suggests regulatory filing in FY24 and potential approval FY25

6. Filing of TAK-609 is subject to feedback from FDA on the ongoing extension trial and may change  
 7. Partnership with Neurocrine Biosciences  
 8. Timeline changes: Eohilia (FY21), TAK-609 (FY22), maribavir 1L (FY23), TAK-611 (FY24)  
 Takeda's Fiscal Year ends March 31 of the following year; e.g., "FY20" refers to the twelve-month period ending March 31, 2021.  
 All timelines are approximate estimates of May 11, 2021. For glossary of disease abbreviations please refer to appendix.

# MAXIMIZING THE VALUE OF OUR GLOBAL AND REGIONAL BRANDS

	PHASE 1 & 2	PHASE 3			FILED			
ONCOLOGY		<p><b>NINLARO</b><sup>®</sup> Proteasome inhibitor Maint. ND MM post-SCT (US, EU)</p> <p><b>NINLARO</b><sup>®</sup> Proteasome inhibitor Maint. ND MM no SCT (US, EU, CN)</p>	<p><b>ICLUSIG</b><sup>®</sup> BCR-ABL inhibitor FL Ph+ ALL (US)</p> <p><b>ALUNBRIG</b><sup>®</sup> ALK inhibitor 2L ALK+NSCLC H2H with alectinib (GL)</p>	<p><b>Cabozantinib</b> <i>Exelixis</i> VEGFR/RTK inhibitor mCRPC combo w/atezolizumab (JP)</p> <p><b>Cabozantinib</b> <i>Exelixis</i> VEGFR/RTK inhibitor 2L mNSCLC combo w/atezolizumab (JP)</p>	<p><b>ALUNBRIG</b><sup>®</sup> ALK inhibitor 1L &amp; 2L ALK+NSCLC (JP, CN)</p> <p><b>NINLARO</b><sup>®</sup> Proteasome inhibitor Maint. ND MM no SCT (JP)</p>	<p><b>ADCETRIS</b><sup>®</sup> <i>Seattle Genetics</i> CD30 ADC CTCL (CN)</p> <p><b>Cabozantinib</b> <i>Exelixis</i> VEGFR/RTK inhibitor 1L RCC combo w/nivolumab (JP)</p>		
RARE GENETICS & HEMATOLOGY	<p><b>NATPARA</b><sup>®</sup> PTH replacement Hypothyroidism (JP)</p>	<p><b>TAKHZYRO</b><sup>®</sup> Anti-kallikrein mAb HAE pediatric (GL)</p> <p><b>TAKHZYRO</b><sup>®</sup> Anti-kallikrein mAb BMA (GL)</p>	<p><b>ADYNOVATE</b><sup>®</sup> Pediatric HemA (EU)</p> <p><b>VONVENDI</b><sup>®</sup> vWF replacement vWD Pediatric on-demand &amp; surgery (GL)</p>	<p><b>TAKHZYRO</b><sup>®</sup> Anti-kallikrein mAb HAE (JP)</p>	<p><b>VONVENDI</b><sup>®</sup> vWF replacement vWD Adult Prophylaxis (GL)</p>			
NEUROSCIENCE								
GASTRO-ENTEROLOGY	<p><b>ENTYVIO</b><sup>®</sup> α4β7 mAb Pediatric UC/CD (GL)</p>	<p><b>ENTYVIO</b><sup>®</sup> α4β7 mAb SubQ CD (US, JP)</p> <p><b>ENTYVIO</b><sup>®</sup> α4β7 mAb GvHD Prophylaxis (EU, JP)</p>	<p><b>Vonoprazan</b> PCAB H. Pylori (CN)</p> <p><b>ALOFISEL</b><sup>®</sup> mesenchymal stem cells Perianal Fistulas in CD (US)</p>	<p><b>ALOFISEL</b><sup>®</sup> mesenchymal stem cells Perianal Fistulas in CD (JP)</p>	<p><b>ENTYVIO</b><sup>®</sup> α4β7 mAb SubQ UC (US, JP)</p> <p><b>Vonoprazan</b> PCAB Oral disintegrated tablet formulation (JP)</p>	<p><b>Vonoprazan</b> PCAB Reflex Esophagitis Maintenance (CN)</p> <p><b>Vonoprazan</b> PCAB Duodenal ulcer (CN)</p>	<p><b>GATTEX</b><sup>®</sup> GLP-2R agonist Pediatric-SBS (JP)</p> <p><b>GATTEX</b><sup>®</sup> GLP-2R agonist Adult-SBS (JP)</p>	
VACCINES								
PDT		<p><b>CUVITRU</b><sup>®</sup> IgG 20% (human) subcutaneous PID (JP)</p>	<p><b>HYQVIA</b><sup>®</sup> <i>Halozyme</i> IgG 10% + Recombinant Human Hyaluronidase CIDP (US, EU)</p>	<p><b>HYQVIA</b><sup>®</sup> <i>Halozyme</i> IgG 10% + Recombinant Human Hyaluronidase Pediatric PID (US, EU<sup>1</sup>)</p>				

● Orphan Drug Designation (in any region / indication for a given asset) ● Potential for registration enabling Ph-2 study  
 ✓ Approved since Q3 FY20 ✗ Discontinued/deprioritized

Status as of May 11, 2021; region abbreviations: GL = global (USA, Europe, Japan, China). Pipeline not all inclusive; programs also ongoing in other Therapeutic Areas

1. Ph4 study completed Q4 FY20

