


















# WE HAVE A RICH PIPELINE OF INNOVATIVE NMEs IN WAVE 1 WITH TARGET LAUNCH BY END OF FY24

TARGET APPROVAL <sup>1</sup> →	FY20	FY21	FY22	FY23	FY24	FY19 STAGE-UPS AND PIVOTAL DATA READOUTS	
 <b>ONCOLOGY</b>		<b>TAK-788<sup>2</sup></b> 2L NSCLC with EGFR exon 20 insertion mutation 		<b>TAK-007</b> Hematologic malignancies 	<b>TAK-924</b> Unfit AML 	<ul style="list-style-type: none"> <li>✓ TAK-007 Ph1/2 addition</li> <li>✓ TAK-924 Ph3 start in Unfit AML</li> <li>✓ TAK-788 Ph3 start in 1L NSCLC</li> <li>✓ TAK-924 Ph2 readout in HR-MDS</li> </ul>	
 <b>RARE DISEASES</b> <i>Immunology Hematology Metabolic</i>		<b>TAK-620</b> CMV infect. in transplant 		<b>TAK-611</b> MLD (IT) 	<b>TAK-607<sup>3</sup></b> Complications of prematurity 	<ul style="list-style-type: none"> <li>✓ TAK-607 Ph2 start in comp. of prematurity</li> <li>✓ TAK-611 Ph2 start in MLD (IT)</li> <li>✓ TAK-755 Ph3 start in cTTP</li> </ul>	
 <b>NEUROSCIENCE</b>		<b>TAK-609</b> Hunter CNS (IT) 		<b>TAK-755</b> cTTP 	<b>TAK-935</b> DEE 	<b>Orexin2R-ag</b> (TAK-925/994) Narcolepsy T1 	<ul style="list-style-type: none"> <li>✓ TAK-994 Ph1 start</li> </ul>
 <b>GASTRO-ENTEROLOGY</b>	<b>TAK-721</b> EoE 					<ul style="list-style-type: none"> <li>✓ TAK-721 Ph3 EoE induction readout</li> <li>✓ TAK-721 Ph3 EoE maintenance readout</li> </ul>	
 <b>VACCINES</b>		<b>TAK-003</b> Dengue Vaccine				<ul style="list-style-type: none"> <li>✓ TAK-003 Ph3 part 2 readout</li> </ul>	






1. Projected timing of approvals depending on data read-outs  
 2. Projected approval date assumes filing on Phase 2 data  
 3. Currently non-pivotal Phase 2 study; assumes interim stage gates would allow for consideration of filing on Phase 2 data

✓ Stage-ups/additions since April 1, 2019  
 ✓ Stage-ups/additions since earnings announcement October 31, 2019

 Orphan potential in at least one indication  
 Estimated dates as of February 4, 2020  
 For glossary of disease abbreviations please refer to appendix.




# OUR EARLY STAGE NMEs AND NEXT-GENERATION PLATFORMS IN WAVE 2 PROVIDE SUSTAINED GROWTH BEYOND FY25






	TARGET APPROVAL <sup>1</sup> →	FY25 AND BEYOND				FY19 STAGE-UPS AND PIVOTAL DATA READOUTS	
 <b>ONCOLOGY</b>	<p><b>TAK-164</b> <i>GI malignancies</i></p> <p><b>TAK-573</b> <i>R/R MM</i></p>	<p><b>TAK-252</b> <i>Solid tumors</i></p> <p><b>TAK-981</b> <i>Multiple cancers</i></p>			<p>CELL THERAPY AND IMMUNE ENGAGERS</p> <p>TARGETED INNATE IMMUNE MODULATION</p> <p>NEXT-GEN CHECKPOINT MODULATORS</p>	<p>✓ TAK-252 Ph1 start in solid tumors</p>	
 <b>RARE DISEASES</b> <i>Immunology Hematology Metabolic</i>	<p><b>TAK-079<sup>2</sup></b> <i>MG, ITP</i></p> <p><b>TAK-755</b> <i>iTTP, SCD</i></p>	<p><b>TAK-754</b> <i>HemA</i></p>			<p>GENE THERAPY</p>	<p>✓ TAK-755 Ph2 start in iTTP</p> <p>✓ TAK-755 Ph1/2 start in SCD</p>	
 <b>NEUROSCIENCE</b>	<p><b>TAK-341</b> <i>Parkinson's Disease</i></p> <p><b>TAK-418</b> <i>Kabuki Syndrome</i></p> <p><b>WVE-120101</b> <i>Huntington's Disease</i></p>	<p><b>Orexin2R-ag</b> <i>Sleep Disorders</i></p> <p><b>TAK-653</b> <i>TRD</i></p> <p><b>WVE-120102</b> <i>Huntington's Disease</i></p>	<p><b>TAK-041</b> <i>CIAS NS</i></p> <p><b>TAK-831</b> <i>CIAS NS</i></p> <p><b>TAK-935</b> <i>CRPS</i></p>		<p>GENE THERAPY</p> <p>OTHER PLATFORMS <i>RNA Modulation Antibody Transport Vehicle</i></p>	<p>✓ TAK-925 Ph1 start in OSA</p> <p>✓ TAK-935 Ph2 start in CRPS</p>	
 <b>GASTRO-ENTEROLOGY</b>	<p><b>Kuma062</b> <i>Celiac Disease</i></p> <p><b>TAK-954</b> <i>POGD</i></p>	<p><b>TAK-101</b> <i>Celiac Disease</i></p> <p><b>TAK-906</b> <i>Gastroparesis</i></p>	<p><b>TAK-018</b> <i>Crohn's Disease (post-op and ileitis)</i></p> <p><b>TAK-951</b> <i>Nausea &amp; vomiting</i></p>	<p><b>TAK-671</b> <i>Acute Pancreatitis</i></p>	<p>GENE THERAPY</p> <p>MICROBIOME</p> <p>CELL THERAPY</p>		
 <b>VACCINES</b>	<p><b>TAK-214</b> <i>Norovirus Vaccine</i></p>	<p><b>TAK-426</b> <i>Zika Vaccine</i></p>	<p><b>TAK-021</b> <i>EV71 vaccine</i></p>				

1. Projected timing of approvals depending on data read-outs
2. TAK-079 to be developed in Rare Diseases indications myasthenia gravis (MG) and immune thrombocytopenic purpura (ITP) – FPI expected Q4 FY19

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# MAXIMIZING THE VALUE OF OUR APPROVED PROGRAMS

	PHASE 1 & 2	PHASE 3	FILED	FY19 STAGE-UPS
 <b>ONCOLOGY</b>	<p><b>ALUNBRIG®</b> ALK inhibitor ALK+NSCLC (JP)</p> <p><b>NINLARO®</b> Proteasome inhibitor R/R MM triplet Tx (GL)</p> <p><b>ICLUSIG®</b> BCR-ABL inhibitor TKI res. chronic phase CML (US)</p> <p><b>NINLARO®</b> Proteasome inhibitor R/R MM doublet Tx (US, EU, JP)</p> <p><b>ALUNBRIG®</b> ALK inhibitor 2L ALK+NSCLC 2<sup>nd</sup> gen TKI (GL)</p>	<p><b>NINLARO®</b> Proteasome inhibitor ND MM (GL)</p> <p><b>ICLUSIG®</b> BCR-ABL inhibitor FL Ph+ ALL (US)</p> <p><b>ALUNBRIG®</b> ALK inhibitor 1L ALK+NSCLC (JP, US, CN)</p> <p><b>NINLARO®</b> Proteasome inhibitor Maint. ND MM no SCT (GL)</p> <p><b>NINLARO®</b> Proteasome inhibitor Maint. ND MM post-SCT (US, EU, CN)</p> <p><b>ALUNBRIG®</b> ALK inhibitor 2L ALK+NSCLC H2H with alectinib (GL)</p> <p><b>Cabozantinib</b> <i>Exelixis</i> VEGFR/RTK inhibitor 1L RCC (JP)</p>	<p><b>Niraparib</b> <i>GlaxoSmithKline</i> PARP 1/2 inhibitor Ovarian cancer – maint. (JP)</p> <p><b>ADCETRIS®</b> <i>Seattle Genetics</i> CD30 ADC R/R sALCL (CN)</p> <p><b>ADCETRIS®</b> <i>Seattle Genetics</i> CD30 ADC R/R HL (CN)</p> <p><b>Niraparib</b> <i>GlaxoSmithKline</i> PARP 1/2 inhibitor Ovarian cancer – salvage (JP)</p> <p><b>ADCETRIS®</b> <i>Seattle Genetics</i> CD30 ADC 1L PTCL (EU)</p> <p><b>Cabozantinib</b> <i>Exelixis</i> VEGFR/RTK inhibitor 2L HCC (JP)</p> <p><b>ALUNBRIG®</b> ALK inhibitor 1L ALK+NSCLC (EU)</p> <p><b>NINLARO®</b> Proteasome inhibitor Maint. ND MM post-SCT (JP)</p>	<ul style="list-style-type: none"> <li>✓ ADCETRIS 1L PTCL filed (EU)</li> <li>✓ NINLARO NDMM transplant filed (JP)</li> <li>✓ Cabozantinib 2L RCC filed (JP)</li> <li>✓ ALUNBRIG Ph3 start H2H alectinib</li> <li>✓ ALUNBRIG Ph2 start 2<sup>nd</sup> gen TKI</li> <li>✓ Niraparib ovarian salvage filed (JP)</li> <li>✓ Niraparib ovarian maint. filed (JP)</li> <li>✓ Cabozantinib 2L HCC filed (JP)</li> </ul>
 <b>GASTRO-ENTEROLOGY</b>	<p><b>ENTYVIO®</b> α4β7 mAb Pediatric UC/CD (US)</p>	<p><b>GATTEX</b> GLP-2R agonist Adult-SBS (JP)</p> <p><b>ALOFISEL®</b> mesenchymal stem cells Perianal Fistulas in CD (US, JP)</p> <p><b>GATTEX</b> GLP-2R agonist Pediatric-SBS (JP)</p> <p><b>ENTYVIO®</b> α4β7 mAb GvHD Prophylaxis (EU, JP)</p> <p><b>ENTYVIO®</b> α4β7 mAb SubQ CD (US, JP)</p>	<p><b>ENTYVIO®</b> α4β7 mAb Crohn's Disease (CN)</p> <p><b>ENTYVIO®</b> α4β7 mAb SubQ UC (US, EU, JP)</p> <p><b>ENTYVIO®</b> α4β7 mAb SubQ CD (EU)</p> <p><b>ENTYVIO®</b> α4β7 mAb Ulcerative Colitis (CN)</p> <p><b>Vonoprazan</b> PCAB Prev. of L-ASA ulcers (JP)</p>	<ul style="list-style-type: none"> <li>✓ ENTYVIO sc UC filed (received CRL)</li> <li>✓ ENTYVIO CD filed (CN)</li> <li>✓ ENTYVIO UC filed (CN)</li> <li>✓ ALOFISEL Ph3 start in CPF</li> <li>✓ Vonoprazan L-ASA ulcer prevention filed (JP)</li> </ul>
 <b>RARE DISEASES</b>	<p><b>NATPARA</b> PTH replacement Hypothyroidism (JP)</p>	<p><b>TAKHZYRO</b> Anti-kallikrein mAb HAE pediatric (GL)</p> <p><b>OBIZUR</b> <i>Ipsen</i> FVIII replacement CHAWI (US, EU)</p> <p><b>VONVENDI</b> vWF replacement vWD Prophylaxis</p> <p><b>TAKHZYRO</b> Anti-kallikrein mAb HAE (JP)</p> <p><b>VONVENDI</b> vWF replacement vWD Pediatric</p> <p><b>ADYNOVATE</b> Pediatric HemA (EU)</p>	<p><b>TAKHZYRO</b> Anti-kallikrein mAb HAE prophylaxis (CN)</p> <p><b>VONVENDI</b> vWF replacement vWD (JP)</p>	<ul style="list-style-type: none"> <li>✓ TAKHZYRO Ph3 start in HAE pediatric</li> <li>✓ TAKHZYRO Ph3 start in HAE (JP)</li> </ul>
 <b>NEUROSCIENCE</b>		<p><b>BUCCOLAM</b> GABA Allosteric Modulator Status Epilepticus (JP)</p>		
 <b>PLASMA-DERIVED THERAPIES</b>		<p><b>CINRYZE</b> PD C1 Esterase inhibitor HAE prophylaxis (JP)</p> <p><b>HYQVIA</b> <i>Halozyme</i> IgG 10% + Recombinant Human Hyaluronidase CIDP</p> <p><b>HYQVIA</b> <i>Halozyme</i> IgG 10% + Recombinant Human Hyaluronidase Pediatric PID (US)</p>		

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- ✓ Stage-ups since earnings announcement October 31, 2019
- Orphan Drug Designation (in any region / indication for a given asset)
- Potential for registration enabling Ph-2 study