Takeda Oncology Pipeline

Beginning with patients' needs, we build our pipeline by identifying the most promising science and collaborate with leading teams around the globe to accelerate innovation. Our oncology R&D activities are focused on translating science into curative or transformative potential treatments by targeting tumor vulnerabilities and through novel strategies that leverage the power of innate immunity.

These therapies in development are investigational and have not been approved by any regulatory bodies for use in the investigational indications listed below.

INVESTIGATIONAL THERAPY	PLATFORM	INVESTIGATIONAL INDICATION	PHASE	ADDITIONAL INFORMATION
Cabozantinib*	Targeted therapy	Metastatic castration- resistant prostate cancer (JP)	Phase 3	Tyrosine kinase inhibitor including MET/AXL/VEGFR Cabozantinib is being developed in Japan in collaboration with Exelixis.
		2L metastatic non-small cell lung cancer (JP)	Phase 3	
lxazomib*	Proteasome inhibitor	Maint. newly diagnosed multiple myeloma post- stem cell transplant (US, EU)	Phase 3	- Proteasome inhibitor
		Maint. newly diagnosed multiple myeloma without stem cell transplant (US, EU, CN)	Phase 3	
Mobocertinib*	Targeted therapy	2L EGFR Exon20 insertion non-small cell lung cancer (CN)	Filed	Oral EGFR Exon20 tyrosine kinase inhibitor
		2L EGFR Exon20 insertion non-small cell lung cancer (JP)	Phase 3	
		1L EGFR Exon20 insertion non-small cell lung cancer	Phase 3	
Ponatinib*	Targeted therapy	FL Ph+ acute lymphocytic leukemia (US)	Phase 3	BCR-ABL inhibitor Takeda shares development rights with Incyte Corp. (Europe, Turkey and Israel) and Otsuka Pharm. (Asia Pacific territories).
		Pediatric Ph+ acute lymphoblastic leukemia	Phase 1	
Relugolix* (TAK-385)	Small molecule	Prostate cancer (JP, Asian countries)	Phase 3	GnRH antagonist
				Relugolix is being developed in Japan and Asian countries in collaboration with Myovant Sciences, Inc.
Niraparib*	Targeted therapy	Triple-negative breast cancer		PARP 1/2 inhibitor
			Phase 3	Niraparib is being developed in collaboration with GSK. Takeda has development and commercialization rights for the treatment of all tumor types in Japan, and all tumor types excluding prostate cancer in South Korea and Taiwan.

INVESTIGATIONAL THERAPY	PLATFORM	INVESTIGATIONAL INDICATION	PHASE	ADDITIONAL INFORMATION
Modakafusp Alfa (TAK-573)	I-O cold-to-hot	Multiple myeloma	Phase 1/2	Immune targeting attenuated cytokine
		Solid tumors	Phase 1/2	Modakafusp alfa (TAK-573) was licensed from Teva Pharmaceuticals.
Subasumstat (TAK-981)	I-O cold-to-hot	Solid tumors	Phase 1/2	SUMOylation inhibitor
		Non-Hodgkin lymphoma	Phase 1/2	
		Multiple myeloma	Phase 1/2	
TAK-007	I-O redirected immunity	B-cell malignancies	Phase 2	CD19 CAR NK TAK-007 is being developed in collaboration with The University of Texas MD Anderson Cancer Center.
TAK-186	I-O redirected immunity	Solid tumors	Phase 1/2	EGFR x CD3 targeting COBRA (COnditional Bispecific Redirected Activation) T cell engager immunotherapy
TAK-102	I-O redirected immunity	Solid tumors	Phase 1	GPC3 targeted CAR-T TAK-102 was licensed from Noile-Immune Biotech.
TAK-103	I-O redirected immunity	Solid tumors	Phase 1	Mesothelin targeted CAR-T TAK-103 was licensed from Noile-Immune Biotech.
TAK-500	I-O cold-to-hot	Solid tumors	Phase 1	Targeted STING agonist
TAK-676	I-O cold-to-hot	Solid tumors	Phase 1	STING agonist
TAK-940	I-O redirected immunity	B-cell malignancies	Phase 1	CD19-1XX CAR-T TAK-940 is being developed in collaboration with Memorial Sloan Kettering Cancer Center.
TAK-280	I-O redirected immunity	Solid tumors	Phase 1/2	B7-H3 x CD3 targeting COBRA (COnditional Bispecific Redirected Activation) T cell engager immunotherapy

^{*}Marketed products have received approval in one or more jurisdictions.

All programs have global development rights unless otherwise noted.

GL = global (USA, Europe, Japan, China) / EU = Europe / JP = Japan / CN = China / GEM = global emerging markets



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