Takeda Oncology Pipeline

Our research and development efforts focus on advancing medicines for thoracic, gastrointestinal and hematologic cancers.

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	MODALITY	INVESTIGATIONAL INDICATION	PHASE	ADDITIONAL INFORMATION
Mirvetuximab soravtansine-gynx (TAK-853) **(Japan rights only)**	ADC	*Folate receptor alpha (FRα)-positive platinum-resistant ovarian cancer*	Filed	Takeda has development and commercialization rights for the treatment in Japan.
		Folate receptor alpha (FRα)-positive platinum-sensitive ovarian cancer	Phase 3	
Rusfertide (TAK-121)	Small molecule	*Polycythemia vera*	Filed	Collaboration with Protagonist Therapeutics. Protagonist is responsible for development in the U.S. through the completion of the Phase 3 VERIFY trial. Takeda has rights for ex-U.S. development and is responsible for leading global regulatory and commercialization activities. Hepcidin mimetic peptide (injection)
Elritercept (TAK-226)	Complex biologic	*2L anemia-associated myelodysplastic syndromes*	Phase 3	Takeda has development and commercialization rights for the treatment worldwide outside of mainland China, Hong Kong and Macau. Activin A/B ligand trap
		1L anemia-associated myelodysplastic syndromes	Phase 3	
		Anemia-associated myelofibrosis	Phase 2	
TAK-928	Complex biologic	*2L squamous non-small cell lung cancer*	Phase 3	Collaboration with Innovent Biologics. Takeda will lead global co-development and U.S. co-commercialization and has exclusive commercial rights outside the U.S. and mainland China, Hong Kong, Macau and Taiwan.
		Solid tumors	Phase 2	
TAK-921	ADC	*3L+ gastric cancer*	Phase 3	Collaboration with Innovent Biologics. Takeda has development and commercialization rights for the treatment worldwide outside of mainland China, Hong Kong, Macau and Taiwan. Ongoing Phase 3 study in 3L+ gastric cancer is being conducted in Japan and China.
		Solid tumors	Phase 1	
TAK-505	Complex biologic	*Solid tumors*	Phase 1	Interventional Phase 1 trial being conducted in the U.S. Bispecific PD-L1 Vd1 gamma delta T-cell engager (injection)
TAK-188	ADC	*Solid tumors*	Phase 1/2	Takeda has global development and commercialization rights to TAK-188, which utilizes licensed technology from Heidelberg Pharma and Memorial Sloan Kettering Cancer Center.
TAK-168	Small molecule	*Solid tumors*	Phase 1	Collaboration with Kumquat Biosciences. Kumquat is responsible for development (KQB168) through the completion of the Phase 1 trial (NCT06994806).

Learn more at **TakedaOncology.com**

Takeda | ONCOLOGY