

R&D Pipelines Covering Various Therapeutic Areas

(As of 1 Sep 2023)



Pre Clinical		Phase I/II		Phase III	Approval for marketing/ Emergency use authorization	
JS011 Undisclosed	JS013 CD93	JS006 TIGIT	JS007 CTLA-4	JS010 CGRP	Bevacizumab VEGF	Toripalimab PD-1
JS018 IL-2	JS104 Pan-CDK	JS009 CD112R	JS014 IL-21	UBP1213sc BLYS	Tifcemalimab BTLA	Adalimumab TNF-α
JS114 Nectin4 ADC	JS115 BCMA ADC	JS015 DKK1	JS105 PI3K-α	JS103 Uricase	Ongericimab PCSK9	Deuremidevir Hydrobromide Tablets RdRp
JS120 IDH1	JS121 SHP2	JS107 Claudin18.2 ADC	JS110 XPO1	JS401 ANGPTL3	JS005 IL-17A	Etesevimab ^{1*} S protein
JS122 FGFR2	JS123 ATR	JS111 EGFR exon 20	JS112 Aurora A	JS026 S protein		
JS205 EGFR x cMet	JS206 IL-2 x PD-1	JS113 EGFR 4th Gen	JS116 KRAS			
JS208 Undisclosed	JS211 PD-L1 x Undisclosed	JS203 CD3 x CD20	JS001sc PD-1			
JS209 CD112R x TIGIT	VV993 3CL protease	JS207 PD-1 x VEGF	JS019 CD39			
JS008 Undisclosed	JT109 Vaccine for Zika virus	JS003 PD-L1	JS012 Claudin18.2			
		JS101 Pan-CDK	JS108 Trop2 ADC			
			JS201 PD-1 x TGF-β			

● Oncology
 ● Metabolism
 ● Immunology
 ● Neurologic
 ● Infectious disease

*Received Emergency Use Authorization from the FDA

- Etesevimab is expected to no longer generate revenue.
- In August 2023, the Company conducted friendly negotiations with IMPACT Therapeutics, Inc. ("IMPACT Therapeutics"). Based on the Company's commercial considerations, both parties have agreed to terminate their cooperation on Shanghai Junpai Yingshi Bio Pharmaceutical Co., Ltd. (the "JV Company") and the PARP inhibitor senaparib (code: JS109/IMP4297). Pursuant to the terms of the agreement, the Company will transfer its 50% equity interest in the JV Company to IMPACT Therapeutics, IMPACT Therapeutics will pay the corresponding share repurchase price to the Company.