Discovery of Selective IRAK4 Degrader based on Protac Technology

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IMMUNOLOGY	Lead
Product Type	Small molecule
Indication	Lupus
Target	IRAK4
MoA(Mechanism of Action)	Blocking TLRs and IL-1R signaling by IRAK4 degradation
Competitiveness	 High IRAK4 degradation potency with DC50 values in single digit nanomolar ranges Confirmed degradation mechanism: Ubiquitin-Proteasome pathway (UPS) High degradation selectivity for CRBN neosubstrates High MLM and HLM stability (> 80% remaining at 30 min), and mouse plasma stability (> 70% remaining at 120 min)
Development Stage	Lead
Route of Administration	Oral