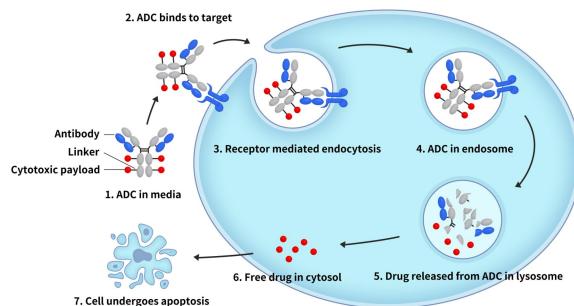


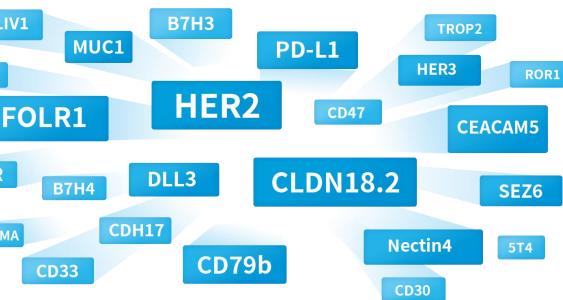


ADC先导分子开发与筛选

抗体偶联药物(antibody-drug conjugate, ADC) 是将单克隆抗体药物的高特异性和小分子细胞毒药物的高活性相结合，用以提高肿瘤药物的靶向性、减少毒副作用。ADC抗体分子的筛选是ADC临床转化前的重要基础，是保证ADC临床有效性和安全性的关键。缔码生物针对每个ADC热门靶点，已经开发出几十种预先开发好的先导分子，同时建立了一整套ADC临床前药效学评价平台。



ADC作用机制



已开发的ADC先导分子热门靶点

平台优势



现货

- 热门ADC靶点的现货分子转让，即刻引进测试
- ADC靶点先导分子库**80+**, B细胞种子库**100+**



丰富的经验

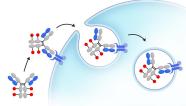
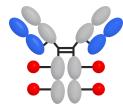
- 已转让ADC项目**10+**
- 正在测试的项目**20+**



赋能ADC开发

- 高灵敏，低背景的内吞试剂
- 抗Payload/Linker单抗 (MMAE, Dxd, Eribulin, SN38, DM1, CL2A (ADC linker))

平台流程



先导分子筛选和抗体生产

人猴稳转株构建

筛选大于10条独特序列的人猴交叉FACS Binder

抗体生产

先导分子初步确定

抗体人源化（可选）

亲和力成熟（可选）

ADC偶联

Benchmark偶联

先导分子偶联

体外药效学评价

先导分子内吞活性测定

天然细胞系杀伤活性测定

最佳先导分子确认

体内药效学评价

小鼠模型构建

体内药效

~8周

1周

4-6周

8-10周



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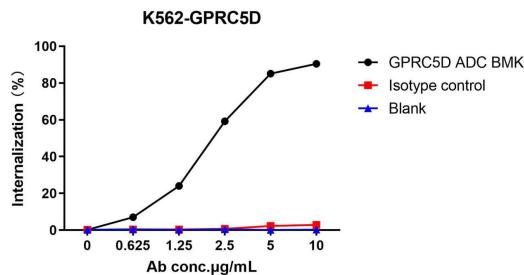
武汉 / Maryland

info@dimabio.com

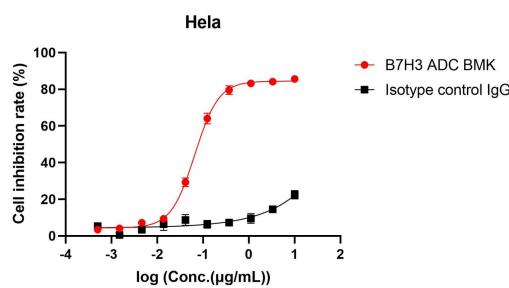
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DiTag™ IgG labeling reagent

pH sensitive IgG labeling reagent

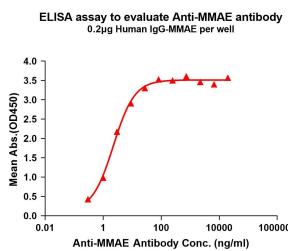


MMAE IgG labeling reagent

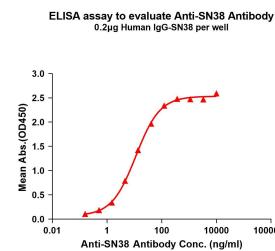


Anti-payload/Linker rabbit antibody

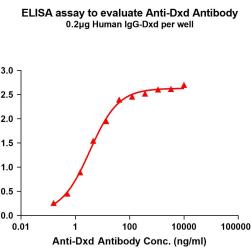
MMAE



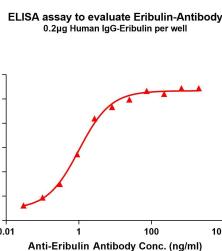
SN38



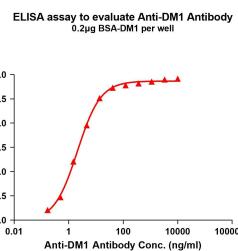
Dxd



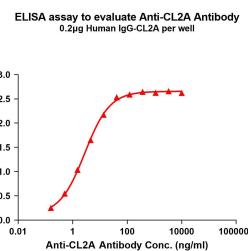
Eribulin



DM1

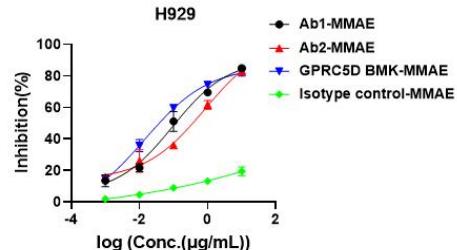
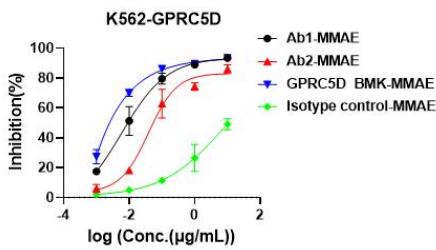
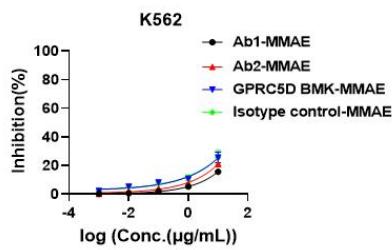


CL2A (ADC linker)



案例展示

Cytotoxicity assay of GPRC5D antibody-drug conjugate (ADC) conjugated with MMAE



Ab1, Ab2: DIMA Lead antibodies against human GPRC5D; Cell line: K562 (negative control), K562-GPRC5D (positive cells), and H929 (GPRC5D-expressing natural cell line)

In this experiment, the data demonstrates the specific cytotoxicity of anti-GPRC5D antibodies against GPRC5D-positive cells following MMAE conjugation.



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