

# 信达生物 Innovent Biologics (1801 HK)

## 稳步推进五年战略规划，国际化战略开始加速

The five-year strategic plan is being steadily advanced, while the internationalization strategy is beginning to accelerate

观点聚焦 Investment Focus

### 维持优于大市 Maintain OUTPERFORM

评级	优于大市 OUTPERFORM
现价	HK\$96.85
目标价	HK\$105.80
HTI ESG	3.6-1.4-3.5
E-S-G: 0-5, (Please refer to the Appendix for ESG comments)	
市值	HK\$165.88bn / US\$21.28bn
日交易额 (3 个月均值)	US\$262.75mn
发行股票数目	1,713mn
自由流通股 (%)	90%
1 年股价最高最低值	HK\$101.90-HK\$30.00
注: 现价 HK\$96.85 为 2025 年 08 月 29 日收盘价	



	1mth	3mth	12mth
绝对值	3.1%	60.9%	119.6%
绝对值 (美元)	3.9%	61.8%	119.7%
相对 MSCI China	1.6%	50.6%	74.2%

Rmb mn	Dec-24A	Dec-25E	Dec-26E	Dec-27E
Revenue	9,422	12,581	15,186	18,691
Revenue (+/-)	52%	34%	21%	23%
Net profit	-206	1,375	1,420	1,817
Net profit (+/-)	n.m.	-767%	3%	28%
Diluted EPS (Rmb)	-0.13	0.84	0.87	1.12
GPM	84.0%	83.7%	84.7%	88.0%
ROE	-1.6%	9.5%	8.9%	10.3%
P/E	n.m.	105	102	79

资料来源: 公司信息, HTI

(Please see APPENDIX 1 for English summary)

### 事件

公司公布 25H1 业绩: 收入 59.5 亿元 (+50.6%), 总产品收入超 52.3 亿元 (+37.3%), 25Q2 产品收入超 27 亿元 (+30%, 环比 +13%)。毛利 51.2 亿元 (+56.3%), 毛利率 86.0% (+3.1pcts), 研发费用 10.0 亿元 (-28.0%), 研发费用率 16.9% (-18.5pcts), 销售费用 23.8 亿元 (+26.4%), 占产品收入 45.4% (-3.9pcts)。期内利润 8.3 亿元 (去年同期-3.9 亿元)。公司在手现金储备超 20 亿 USD。

### 点评

**稳步推进五年战略规划，国际化战略开始加速。**国内商业化表现稳步提升，产品持续增长主要得益于肿瘤板块的稳固基础，以及 CVM 板块的双轮驱动: 1) 肿瘤领域 PD-1 等产品品牌和产品组合优势明显, KRAS G12C、三代 EGFR-TKI 等新产品贡献营收; 2) 综合管线中 PCSK9、IGF-1R 等新产品准入顺利, 快速放量。玛仕度肽也顺利获批。利润端实现 8.3 亿元利润超市场预期, 主要由于确认了罗氏引进 DLL3 ADC 的 8000w USD 首付款以及研发费用效率优化。考虑到 IBI363 (PD-1/IL-2<sup>α</sup>-bias) 的全球 MRCT 临床开展、玛仕度肽、IBI112 (IL-23p19)、IBI128 (XOI) 的 III 期临床开展以及早期管线入组节奏, 我们认为公司下半年研发费用会高于上半年, 全年维持在 21~28 亿元区间。

公司创始人俞德超博士再次强调, 公司正按既定路径稳步推进: 预计 2025 年实现 EBITDA 盈利, 2027 年国内营收达到 200 亿元, 2030 年至少 5 个管线进入全球多中心 III 期临床研究。明确的战略目标彰显了公司迈向全球一流生物医药企业的决心, 也为市场信心提供了有力支撑。公司亦强调国际化布局, 积极在全球范围引进临床开发、运营及供应链人才, 为 IBI363 等产品的海外临床奠定坚实基础。

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**肿瘤领域：IBI363 有望成为下一代 IO 基石药物，新一代 ADC 推动“IO+ADC”组合疗法升级。**

**IBI363 末线到前线/围手术期，IO 经治、“冷肿瘤”全面布局。** IO 初治黑色素瘤（粘膜/肢端）的关键性研究正在进行中；肺癌领域，针对 IO 耐药的 sqNSCLC，公司已启动全球多中心 III 期研究（MRCT），计划在中国、日本、美国、加拿大、欧盟及英国等地招募约 600 例患者，评估 IBI363（3 mg/kg 单药）相较于西他赛在经铂类化疗及抗 PD-1/PD-L1 治疗后进展的无法手术切除局部晚期 nqNSCLC 患者中的疗效与安全性，主要研究终点为 OS。同时，公司在 3L mss CRC 适应症上正筹备开展 III 期临床研究；在一线 NSCLC 及一线 CRC 方面，I/IIb 期研究已在计划中，预计将于 2026 年陆续读出数据。此外，公司还在积极拓展其他适应症，包括铂耐药卵巢癌、EGFR 突变 NSCLC 以及 nsqNSCLC 新辅助治疗等。

**双抗/双 payload ADC、三抗 TCE 蓄势待发。** IBI3001（EGFR/B7H3 ADC）正在中美澳全球爬坡中，确定剂量后会探索联合疗法。IBI3020（CEACAM5 dp ADC）采用自主研发毒素，克服耐药，同时优化 linker 保持稳定性，提高治疗窗口。目前全球 I 期剂量爬坡中，以及爬升到较高剂量，有望 25 年取得 POC 数据。公司也重点强调了 IBI3003（GPC5D/BCMA/CD3）具备同类最佳潜力，目前中澳 I 期临床爬坡中，美国患者即将入组。公司在爬坡中看到不错结果，在 MM 适应症中 ORR、CR 率有望媲美 CAR-T 疗法，有望解决 GPRC5D、BCMA 单靶疗法出现耐药问题。

**CVM 领域：玛仕度肽有望成为国内减重降糖护肝领导品牌，叠加渠道布局加速放量。**

**玛仕度肽有望加速放量。** 玛仕度肽的 T2D 适应症有望 25H2 获批，同时青少年肥胖适应症的 III 期临床即将启动。公司表示玛仕度肽的市场反馈不错。我们认为玛仕度肽定价较双靶竞品低约 30%，叠加线上线下渠道协同，有望实现快速放量。

**GLP-1 口服小分子有差异化优势：** 公司表示 IBI3032 虽然采用 Orforglipron 骨架，但半衰期更长，口服暴露量是同类分子 5-10 倍，并且专利风险较小。其他适应症比如高血压也有潜力，和其他 CVM 管线（XOI 等）形成协同，有望成为下一代基石产品。

**公司 25 年研发催化剂丰富。推荐关注：**

- 1) IBI363: 3L mssCRC 的 III 期临床研究开展；1L NSCLC、1L CRC 的 I/IIb 研究数据有望读出（26 年）。
- 2) IBI343（CLDN18.2 ADC）：2L 胰腺癌全球 MRCT 注册研究有望开展（25H2-26）；1L 胰腺癌的 I 期数据读出。
- 3) IBI354（HER2 ADC）：铂耐药卵巢癌的 III 期数据读出；HER2+ BC 的 III 期有望开展。
- 4) IBI3003（GPC5D/BCMA/CD3）：I 期数据读出（25H2）。
- 5) 玛仕度肽：T2D 适应症获批（25H2）；Dreams-3 研究顶线数据读出（25H2）；GLORY-2 研究顶线数据读出（25H2）；青少年肥胖 I 期数据读出，III 期研究启动（25H2）。
- 6) IBI128（XOI）：II 期数据在亚太风湿病学会年会读出（25.09）以及 III 期研究启动（25H2）。

- 7) IBI3032 (GLP-1 口服小分子): 25Q3 进入中美 I 期, 部分数据有望读出 (25H2)。
  - 8) IBI3002 (TSLP/IL-4R $\alpha$ ): 初步数据读出 (25H2)。
  - 9) IBI356 (OX40L): I 期数据潜在读出, 以及 II 期启动 (25H2)。
- IBI3016 (AGT siRNA): I 期数据在美国 2025 AHA 大会读出 (25H2)。

#### 盈利预测及估值建议:

结合公司 2025 H1 年营收情况、我们上调了公司盈利预测以及更新估值模型。我们预计 2025-27 年总收入为 125.8/151.9/186.9 亿元 (2025-27E 前值: 118.6/143.4/181.7), 同比+34%/+21%/+23%。2025 年扭亏为盈, 实现净利润 13.8 亿元 (前值 3.8 亿元)。我们采用 DCF 模型对公司进行估值, 采用 FY26-37 现金流进行测算, 基于 WACC 9.8% (不变), 永续增长率 3.5% (不变), 假设汇率 RMB:HKD=1:1.14, 调整目标价至 105.8 HKD/股 (前值: 90.1 HKD/股), 维持“优于大市”评级。

**风险提示:** 新药研发风险, 新药审批风险, 新药商业化不及预期风险。

表 1. DCF 估值模型

RMB m	FY24A	FY25E	FY26E	FY27E	FY28E	FY29E	FY30E	FY31E	FY32E	FY33E	FY34E	FY35E	FY36E	FY37E
	2024	2025	2026	2027	2028	2029	2030	2031	2032	2033	2034	2035	2036	2037
Forecast Year		1	2	3	4	5	6	7	8	8	8	8	8	8
Time Factor	0.0	0.3	1.3	2.3	3.3	4.3	5.3	6.3	7.3	8.3	9.3	10.3	11.3	12.3
(fraction of year to next FY end)														
<b>Sales</b>	<b>9422</b>	<b>12581</b>	<b>15186</b>	<b>18691</b>	<b>21488</b>	<b>24934</b>	<b>27961</b>	<b>30382</b>	<b>32604</b>	<b>34216</b>	<b>35599</b>	<b>36607</b>	<b>37568</b>	<b>38500</b>
... Growth	51.8%	33.5%	20.7%	23.1%	15.0%	16.0%	12.1%	8.7%	7.3%	4.9%	4.0%	2.6%	2.6%	2.5%
Gross Profit	7912	10529	12861	16448	18909	21942	24605	26736	28691	30111	31327	32214	33060	33880
... GP Margin	84.0%	83.7%	84.7%	88.0%	88.0%	88.0%	88.0%	88.0%	88.0%	88.0%	88.0%	88.0%	88.0%	88.0%
SG&A	-8668	-9477	-11756	-14876	-16357	-17127	-18146	-19203	-20233	-20858	-21318	-21902	-22460	-23001
... SG&A Margin	92.0%	75.3%	77.4%	79.6%	76.1%	68.7%	64.9%	63.2%	62.1%	61.0%	59.9%	59.8%	59.8%	59.7%
Depreciation & Amortisation	581	473	695	655	636	631	642	665	695	731	770	810	848	885
EBIT	-122	1685	1738	2206	3835	6749	9048	11135	13058	14058	15023	15334	15631	15919
Add: Amortisation	0	0	0	0	0	0	0	0	0	0	0	0	0	0
EBITA	-122	1685	1738	2206	3835	6749	9048	11135	13058	14058	15023	15334	15631	15919
... Margin	-1.3%	13.4%	11.4%	11.8%	17.8%	27.1%	32.4%	36.6%	40.1%	41.1%	42.2%	41.9%	41.6%	41.3%
... Growth														
Add: Depreciation	581	473	695	655	636	631	642	665	695	731	770	810	848	885
EBITDA	458	2,159	2,433	2,861	4,472	7,380	9,690	11,799	13,753	14,790	15,793	16,143	16,479	16,805
... Margin	4.9%	17.2%	16.0%	15.3%	20.8%	29.6%	34.7%	38.8%	42.2%	43.2%	44.4%	44.1%	43.9%	43.6%
Less: Tax	116	-16	-243	-251	-321	-565	-1,002	-1,347	-1,660	-1,949	-2,099	-2,243	-2,290	-2,335
Less: Minority Interests	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Less: Increase of Working Capital	3,047	-1,181	-590	1,181	1,181	2,362	2,362	2,362	2,362	2,362	2,362	2,362	2,362	2,362
Less: Capex	-283	-377	-456	-561	-645	-748	-839	-911	-978	-1,026	-1,068	-1,098	-1,127	-1,155
... Capex/Depreciation	0.5x	0.8x	0.7x	0.9x	1.0x	1.2x	1.3x	1.4x	1.4x	1.4x	1.4x	1.4x	1.3x	1.3x
Less: Acquisitions	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
<b>Free Cash Flow</b>	<b>3,339</b>	<b>584</b>	<b>1,144</b>	<b>3,230</b>	<b>4,687</b>	<b>8,429</b>	<b>10,211</b>	<b>11,903</b>	<b>13,477</b>	<b>14,177</b>	<b>14,988</b>	<b>15,164</b>	<b>15,424</b>	<b>15,677</b>
... FCF Growth	-285.6%	-82.5%	95.8%	182.3%	45.1%	79.8%	21.1%	16.6%	13.2%	5.2%	5.7%	1.2%	1.7%	1.6%
PV of FCF	3,339	566	1,010	2,596	3,431	5,620	6,201	6,583	6,789	6,504	6,263	5,771	5,347	4,950
Bull Case														
WACC														
Risk Free Rate	3.0%													
Market Risk Premium	9.5%													
Equity Beta	0.88													
Cost of Equity	11.4%													
Cost of Debt (Pre-tax)	6.0%													
Cost of Debt (After tax)	5.1%													
Target Debt weight	25.0%													
Target Equity weight	75.0%													
Tax Rate	15.0%													
DCF Valuation														
Sum of PV of FCF														61,065
PV of Terminal Value														81,378
Enterprise Value														142,443
Add: Net Cash FY24														7,508
Equity Value (RMB mn)														149,951
Equity Value (USD mn)														\$ 19,348
Equity Value (HKD mn)														164,946
FX:														1.10
Diluted weighted shares outstanding														1,560
Value per Share, HKD														HK\$ 105.8

资料来源: 公司财报; HTI

表 2.

Key financials	Dec-24A	Dec-25E	Dec-26E	Dec-27E	Profit & Loss (Rmb'm)	Dec-24A	Dec-25E	Dec-26E	Dec-27E
Revenue (Rmbm)	9422	12581	15186	18691	Total turnover	9,422	12,581	15,186	18,691
Operating Profit /Loss (RMBm)	-756	1,052	1,105	1,572	Cost of sales	1,510	2,052	2,325	2,243
Pre-tax profit / Loss (RMBm)	-190	1,618	1,671	2,138	<b>Gross profit</b>	7,912	10,529	12,861	16,448
Net income to ord equity (RMBm)	(206)	1,375	1,420	1,817	Total operating costs	8,668	9,477	11,756	14,876
Revenue growth	52%	34%	21%	23%	<b>Operating profit</b>	(756)	1,052	1,105	1,572
Net profit growth	n.a.	n.a.	3.3%	28.0%	Operating EBITDA	648	2,348	2,622	3,050
Adjusted net income to ord equity (Rmbm)	(206)	1,375	1,420	1,817	<b>Depreciation and amortisation</b>	581	473	695	655
ROE	-1.6%	9.5%	8.9%	10.3%	<b>Operating EBIT</b>	(122)	1,685	1,738	2,206
					Interest income (expense)	190	190	190	190
					Share of loss from an associate/JV	-	-	-	-
<b>Balance Sheet (Rmb'm)</b>	<b>Dec-24A</b>	<b>Dec-25E</b>	<b>Dec-26E</b>	<b>Dec-27E</b>	<b>Pre-tax profit</b>	(190)	1,618	1,671	2,138
Total cash and equivalents	7,508	7,988	9,246	12,528	Taxation	16	243	251	321
Inventories	822.2	623.8	69.9	67.5	<b>Net Income</b>	(206)	1,375	1,420	1,817
Account and other receivables	376	1,883	3,103	961	Minorities	-	-	-	-
Trade receivables	1,184	2,039	2,461	3,029	<b>Net Income to ord equity</b>	(206)	1,375	1,420	1,817
Other current assets	383	383	383	383	One-off expense	-	-	-	-
<b>Total current assets</b>	<b>10,273</b>	<b>12,916</b>	<b>15,263</b>	<b>16,968</b>	<b>Normalized net income</b>	<b>(206)</b>	<b>1,375</b>	<b>1,420</b>	<b>1,817</b>
Property, plant and equipment	5,280	5,184	4,945	4,851					
Other non-current assets	6,050	6,050	6,050	6,050	<b>Per Share Data</b>	<b>Dec-24A</b>	<b>Dec-25E</b>	<b>Dec-26E</b>	<b>Dec-27E</b>
<b>Total non-current assets</b>	<b>11,330</b>	<b>11,234</b>	<b>10,995</b>	<b>10,901</b>	EPS (Rmb)	(0.13)	0.84	0.87	1.12
<b>Total assets</b>	<b>21,603</b>	<b>24,150</b>	<b>26,258</b>	<b>27,869</b>	Revenue per share (Rmb)	5.79	7.73	9.33	11.48
Contract liabilities	256	256	256	256	Operating EBITDA per share (Rmb)	0.28	1.33	1.49	1.76
Trade and other payable	358	673	762	735	BVPS (Rmb)	8.06	8.91	9.78	10.89
Bank borrowing	405	405	405	405	DPS (Rmb)	-	-	-	-
Other current liabilities	3,350	4,497	5,095	4,916	Recurrent cash flow per share (Rmb)	1.61	0.29	0.77	2.02
<b>Total current liabilities</b>	<b>4,369</b>	<b>5,831</b>	<b>6,519</b>	<b>6,313</b>	Shares in issue (million)	1,627	1,627	1,627	1,627
Bank borrowing	2,412	2,412	2,412	2,412	Year end adjusted shares in issue (million)	1,627	1,627	1,627	1,627
Contract liabilities	568	568	568	568					
Other liabilities	1,136	846	846	846	x				
<b>Total non-current liabilities</b>	<b>4,116</b>	<b>3,826</b>	<b>3,826</b>	<b>3,826</b>	<b>Key Ratios</b>	<b>Dec-24A</b>	<b>Dec-25E</b>	<b>Dec-26E</b>	<b>Dec-27E</b>
<b>Total liabilities</b>	<b>8,485</b>	<b>9,657</b>	<b>10,345</b>	<b>10,139</b>	<b>Growth</b>				
<b>Shareholder's equity</b>	<b>13,118</b>	<b>14,493</b>	<b>15,913</b>	<b>17,730</b>	Revenue growth	51.8%	33.5%	20.7%	23.1%
Minority interests	-	-	-	-	Operating profit growth	56.1%	33.1%	22.1%	27.9%
<b>Total equity</b>	<b>13,118</b>	<b>14,493</b>	<b>15,913</b>	<b>17,730</b>	Net profit growth	-80.0%	-767.5%	3.3%	28.0%
<b>Total liabilities &amp; shareholders' equity</b>	<b>21,603</b>	<b>24,150</b>	<b>26,258</b>	<b>27,869</b>	<b>Margins</b>				
					Gross margin	84.0%	83.7%	84.7%	88.0%
<b>Cash flow (Rmb'm)</b>	<b>Dec-24A</b>	<b>Dec-25E</b>	<b>Dec-26E</b>	<b>Dec-27E</b>	Operating EBITDA margin	4.9%	17.2%	16.0%	15.3%
<b>Operating profit</b>	<b>(756)</b>	<b>1,052</b>	<b>1,105</b>	<b>1,572</b>	Operating margin	-8.0%	8.4%	7.3%	8.4%
Deprecation and amortisation	581	473	695	655	Pretax profit margin	-2.0%	12.9%	11.0%	11.4%
Changes in working capital	(3,047)	1,181	590	(1,181)	Tax rate	0.0%	15.0%	15.0%	15.0%
Other operating cash flow	6,901	(1,782)	(609)	2,864	Net profit margin	-2.2%	10.9%	9.4%	9.7%
<b>Cash generated from operations</b>	<b>3,679</b>	<b>925</b>	<b>1,781</b>	<b>3,910</b>	<b>Key Ratios</b>				
Capex	(283)	(377)	(456)	(561)	ROE	-1.6%	9.5%	8.9%	10.3%
Other investing cash flow	-	-	-	-	ROA	-1.0%	5.7%	5.4%	6.5%
<b>Net cash flow from investing activities</b>	<b>(283)</b>	<b>(377)</b>	<b>(456)</b>	<b>(561)</b>	Capex/revenue	-3.0%	-3.0%	-3.0%	-3.0%
Change in borrowings	(704)	-	-	-	Current ratio (x)	2.4	2.2	2.3	2.7
Proceeds from changes in capital	-	-	-	-	Creditor days	120	120	120	120
Other financing cash flow	(68)	(68)	(68)	(68)	Debtor days	59	59	59	59
<b>Net cash flow from financing activities</b>	<b>(772)</b>	<b>(68)</b>	<b>(68)</b>	<b>(68)</b>	Inventory days	111	11	11	11
Cash at beginning of period	7,508	7,988	9,246	12,528	Sales/avg assets	0.8	1.1	1.1	1.2
<b>Net change in cash</b>	<b>2,624</b>	<b>480</b>	<b>1,258</b>	<b>3,282</b>	<b>Credit analysis</b>				
Forex effects	-	-	-	-	Debt/EBITDA (x)	7.7	1.5	1.3	1.1
<b>Implied cash at end of period</b>	<b>7,988</b>	<b>9,246</b>	<b>12,528</b>	<b>17,093</b>	Debt/equity	28%	25%	23%	21%
<b>Free cash flow</b>	<b>3,396</b>	<b>547</b>	<b>1,326</b>	<b>3,350</b>	Net debt to equity	-16%	-10%	-12%	-31%

资料来源: 公司财报; HTI

## APPENDIX 1

### Summary

The company announced its 1H25 results: revenue reached RMB 5.95 billion (+50.6% YoY), with total product sales exceeding RMB 5.23 billion (+37.3% YoY). 2Q25 product sales were over RMB 2.7 billion (+30% YoY, +13% QoQ). Gross profit was RMB 5.12 billion (+56.3% YoY), with a gross margin of 86.0% (+3.1ppts). R&D expenses were RMB 1.0 billion (-28.0% YoY), representing 16.9% of revenue (-18.5ppts). Selling expenses amounted to RMB 2.38 billion (+26.4% YoY), accounting for 45.4% of product revenue (-3.9ppts). Net profit for the period was RMB 830 million (vs. -RMB 390 million in 1H24). The company's cash reserves exceeded USD 2.0 billion.

### Comments

The company is steadily advancing its five-year strategic plan, while its internationalization strategy is gaining momentum. Domestic commercialization performance continued to improve, with product growth mainly driven by: (1) the solid foundation of the oncology portfolio, including brand and combination advantages of PD-1 and other products, as well as revenue contribution from new launches such as KRAS G12C inhibitors and third-generation EGFR-TKIs; (2) the CVM pipeline, where new products such as PCSK9 and IGF-1R achieved smooth market access and rapid uptake. In addition, masitutumide was successfully approved. On the profitability side, the company reported net profit of RMB 830 million, exceeding market expectations. This was primarily due to the recognition of the USD 80 million upfront payment from Roche for the DLL3 ADC licensing deal and improved R&D efficiency. Given the global MRCT of IBI363 (PD-1/IL-2 $\alpha$ -bias), Phase III clinical studies of masitutumide, IBI112 (IL-23p19), and IBI128 (XOI), as well as the enrollment progress of early-stage assets, we expect R&D spending in 2H25 to be higher than in 1H25, with full-year R&D expenses remaining in the RMB 2.1–2.8 billion range.

Founder Dr. Dechao Yu reiterated that the company is steadily progressing along its established pathway: targeting EBITDA breakeven by 2025, domestic revenue of RMB 20 billion by 2027, and at least five assets entering global multi-regional Phase III clinical trials by 2030. These clear strategic milestones demonstrate the company's determination to become a world-class biopharmaceutical enterprise and provide strong confidence to the market. The company also emphasized its internationalization strategy, actively recruiting global talent in clinical development, operations, and supply chain to lay a solid foundation for overseas clinical advancement of assets such as IBI363.

**Oncology: IBI363 is poised to become the next-generation backbone IO therapy, while the company's new ADC portfolio is driving an "IO + ADC" therapeutic upgrade.**

**IBI363 is being developed across the continuum from late-line to front-line and perioperative settings, targeting IO-pretreated and "cold tumor" populations.** A pivotal study in treatment-naïve melanoma (mucosal/acral subtypes) is ongoing. In lung cancer, the company has initiated a global multi-regional Phase III trial (MRCT) in IO-refractory sqNSCLC, enrolling ~600 patients across China, Japan, the US, Canada, the EU, and the UK. The study evaluates IBI363 (3 mg/kg monotherapy) versus docetaxel in unresectable, locally advanced nqNSCLC patients who progressed after platinum chemotherapy and PD-1/PD-L1 treatment, with OS as the primary endpoint. Preparations are underway for a Phase III trial in 3L MSS CRC, while Phase I/IIb trials in first-line NSCLC and CRC are planned, with readouts expected from 2026 onward. Additional indications under exploration include platinum-resistant ovarian cancer, EGFR-mutant NSCLC, and neoadjuvant therapy in nsqNSCLC.

Next-generation modalities are also advancing, including bispecific/dual-payload ADCs and trispecific TCEs. IBI3001 (EGFR/B7H3 ADC) is in global dose-escalation studies (China/US/Australia) and will move into combination regimens once the RP2D is defined. IBI3020 (CEACAM5 dual-payload ADC), leveraging a proprietary cytotoxin and optimized linker technology, is designed to overcome resistance and widen the therapeutic window. Global Phase I dose-escalation is ongoing at higher dose levels, with proof-of-concept (POC) data expected in 2025. Meanwhile, IBI3003 (GPRC5D/BCMA/CD3 trispecific TCE) is advancing in China and Australia with US enrollment starting soon. Early clinical signals are encouraging, with ORR and CR rates in multiple myeloma potentially comparable to CAR-T therapies, while addressing resistance seen in GPRC5D- or BCMA-targeted monotherapies.



**Cardiovascular & Metabolic (CVM): Masitidutide is expected to become a leading domestic brand in weight loss, diabetes, and liver protection, with accelerated ramp-up in sales supported by robust channel expansion.** The T2D indication is anticipated to receive approval in 2H25, while a Phase III trial in adolescent obesity is about to begin. Market feedback has been positive, with pricing set ~30% below competitors and synergistic online/offline channel strategies expected to drive rapid uptake.

The company is also advancing differentiated oral GLP-1 programs. IBI3032, though based on an orforglipton backbone, demonstrates a longer half-life and 5–10x higher oral exposure than peer molecules, while carrying lower patent risk. Beyond diabetes and obesity, IBI3032 has potential in hypertension, complementing other CVM pipeline assets such as the xanthine oxidase inhibitor (XOI), reinforcing its potential as a next-generation cornerstone product.

**The company has a rich pipeline of R&D catalysts in 2025. Key programs to watch include:**

- 1) IBI363: Initiation of a Phase III trial in 3L MSS CRC; Phase I/IIb readouts expected in 1L NSCLC and 1L CRC (2026).
- 2) IBI343 (CLDN18.2 ADC): Global registrational MRCT in 2L pancreatic cancer expected to initiate (2H25–26); Phase I data readout in 1L pancreatic cancer.
- 3) IBI354 (HER2 ADC): Phase III readout in platinum-resistant ovarian cancer; potential initiation of Phase III in HER2+ breast cancer.
- 4) IBI3003 (GPRC5D/BCMA/CD3): Phase I data readout (2H25).
- 5) Masitidutide: T2D indication approval anticipated (2H25); topline results from the DREAMS-3 and GLORY-2 studies expected (2H25); Phase I readout in adolescent obesity and initiation of Phase III (2H25).
- 6) IBI128 (XOI): Phase II data presentation at the APLAR Annual Meeting (Sep 2025) and initiation of Phase III (2H25).
- 7) IBI3032 (oral GLP-1 small molecule): Entry into Phase I in China and the US (Q3 2025), with partial data expected in 2H25.
- 8) IBI3002 (TSLP/IL-4R $\alpha$ ): Preliminary data readout (2H25).
- 9) IBI356 (OX40L): Potential Phase I readout and initiation of Phase II (2H25).
- 10) IBI3016 (AGT siRNA): Phase I data presentation at the American Heart Association (AHA) Scientific Sessions (2H25).

#### **Earnings Forecast and Valuation Recommendation**

Based on the company's 2025 H1 revenue performance, we have raised our earnings forecasts and updated our valuation model. We now project total revenue of RMB 12.58/15.19/18.69 billion in 2025–27 (vs. previous estimates of RMB 11.86/14.34/18.17 billion), representing YoY growth of +34%/+21%/+23%. We expect the company to turn profitable in 2025, with net profit reaching RMB 1.38 billion (vs. previous estimate of RMB 380 million). Using a DCF model based on FY26–37 cash flows, and assuming WACC of 9.8% (unchanged), perpetual growth of 3.5% (unchanged), and an exchange rate of RMB:HKD = 1:1.14, we revise our target price to HKD 105.8 per share (vs. previous HKD 90.1), and maintain our "Outperform" rating.

**Risks:** risks in new drug research and development, risks in new drug approval, risks in commercializing new drugs.

## APPENDIX 2

### ESG Comments

#### **Environmental:**

the overall performance of company on environment is good

#### **Social:**

the overall performance of company on society is good

#### **Governance:**

the overall performance of company on government is good



## 附录 APPENDIX

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卖出，未来 12-18 个月内预期相对基准指数跌幅在 10%以上，基准定义如下

各地股票基准指数：日本 – TOPIX, 韩国 – KOSPI, 台湾 – TAIEX, 印度 – Nifty100; 其他所有中国概念股 – MSCI China.

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#### Haitong International Equity Research Ratings Distribution, as of March 31, 2025

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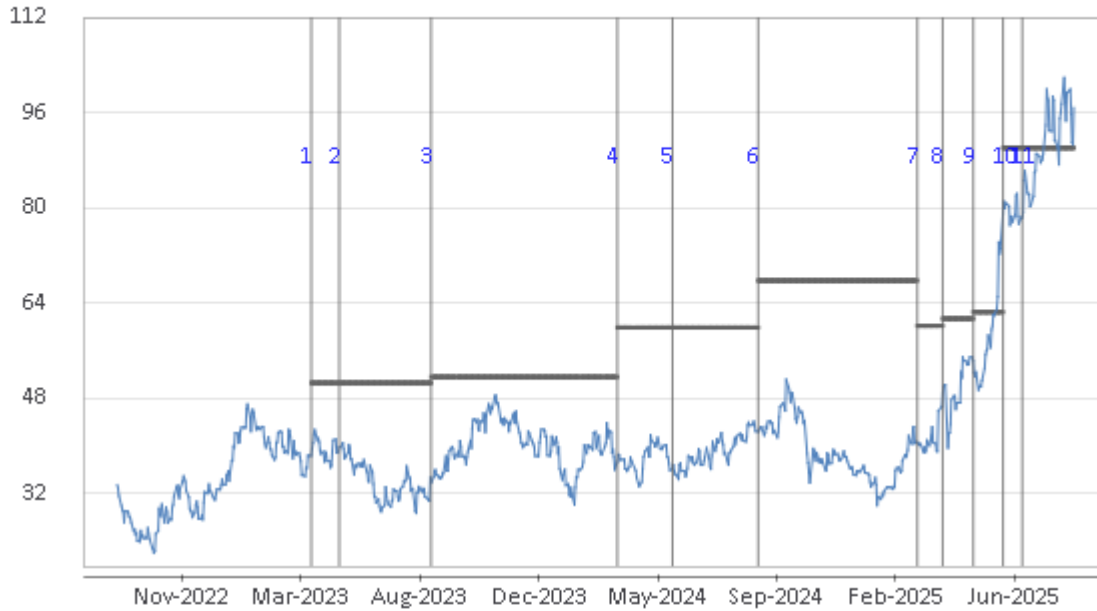
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## Recommendation Chart

### Innovent Biologics - 1801 HK



1. 10 Apr 2023 OUTPERFORM at 38.35 target 50.60.
2. 12 May 2023 OUTPERFORM at 39.75 target 50.60.
3. 25 Aug 2023 OUTPERFORM at 34.65 target 51.60.
4. 25 Mar 2024 OUTPERFORM at 36.00 target 59.90.
5. 27 May 2024 OUTPERFORM at 35.90 target 59.90.
6. 2 Sep 2024 OUTPERFORM at 42.45 target 67.80.
7. 3 Mar 2025 OUTPERFORM at 43.20 target 60.20.
8. 1 Apr 2025 OUTPERFORM at 46.60 target 61.40.
9. 6 May 2025 OUTPERFORM at 54.30 target 62.50.
10. 9 Jun 2025 OUTPERFORM at 73.15 target 90.10.
11. 1 Jul 2025 OUTPERFORM at 78.40 target 90.10.

Source: Company data Bloomberg, HTI estimates