7 Apr 2025



亚盛医药 Ascentage Pharma (6855 HK)

国内商业化表现稳健,全球化创新进入加速期

Domestic commercialization remains solid, while global innovation enters an accelerated phase



观点聚焦 Investment Focus

维持优于	大市M	aintain Ol	JTPERFO	RM
评级			优于大市 0	UTPERFORM
现价				HK\$45.40
目标价				HK\$51.90
HTI ESG				5.0-4.4-5.0
E-S-G: 0-5, (Please refer to	the Appendix fo	or ESG comments)		
市值			HK\$15.81bn	/ US\$2.03bn
日交易额 (3 个月均值)			US\$18.75mn
发行股票数目				348.27mn
自由流通股 (%)				70%
1年股价最高最低值			HK\$47.	70-HK\$15.70
注: 现价 HK\$44.85 为	2025年04	月 01 日收盘价		
_	Price Retu	rn — N	ASCI China	
265 ———		Pm .	•••	
210 ———		Per les	M MY	W
155 —		~		
100			~~~~	MAC.
45 ———				
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Volume	<u></u>	.b.aMl.biba.aasa	التمال ألمحسما	Namada Namada
Apr-24	Aug-2	4 De	:-24	Apr-25
	Aug-2	4 De	c-24	Apr-25
Apr-24	Aug-2			·
Apr-24	Aug-2	4 Dec	3mth -1.3%	Apr-25 12mth 152.5%
Apr-24 <i>资料来源: Factset</i>	Aug-2	1mth	3mth	12mth
Apr-24 <i>资料来源: Factset</i> 绝对值	Aug-2	1mth 19.4%	3mth -1.3%	12mth 152.5%
Apr-24 <i>资料来源: Factset</i> 绝对值 绝对值 (美元)	Aug-2	1mth 19.4% 19.4%	3mth -1.3% -1.5%	12mth 152.5% 154.0%
Apr-24 <i>资料来源: Factset</i> 绝对值 绝对值 (美元)	Aug-2	1mth 19.4% 19.4%	3mth -1.3% -1.5%	12mth 152.5% 154.0%
Apr-24 <i>资料未源: Factset</i> 绝对值 绝对值(美元) 相对 MSCI China		1mth 19.4% 19.4% 17.5%	3mth -1.3% -1.5% -16.0%	12mth 152.5% 154.0% 116.8%
Apr-24 <i>资料未源: Factset</i> 绝对值 绝对值(美元) 相对 MSCI China Rmb mn	Dec-23A	1mth 19.4% 19.4% 17.5%	3mth -1.3% -1.5% -16.0%	12mth 152.5% 154.0% 116.8%
Apr-24 <i>资料未源: Factset</i> 绝对值 绝对值(美元) 相对 MSCI China Rmb mn Revenue	Dec-23A 222	1mth 19.4% 19.4% 17.5% Dec-24A 981	3mth -1.3% -1.5% -16.0% Dec-25E 569	12mth 152.5% 154.0% 116.8% Dec-26E 2,992
Apr-24 <i>资料未源: Factset</i> 绝对值 绝对值 (美元) 相对 MSCI China Rmb mn Revenue Revenue (+/-) Net profit Net profit (+/-)	Dec-23A 222 6%	1mth 19.4% 19.4% 17.5% Dec-24A 981 342% -406 n.m.	3mth -1.3% -1.5% -16.0% Dec-25E 569 -42%	12mth 152.5% 154.0% 116.8% Dec-26E 2,992 426%
Apr-24 <i>资料未源: Factset</i> 绝对值 绝对值(美元) 相对 MSCI China Rmb mn Revenue Revenue (+/-) Net profit Net profit (+/-) Diluted EPS (Rmb)	Dec-23A 222 6% -926 n.m. -3.28	1mth 19.4% 19.4% 17.5% Dec-24A 981 342% -406 n.m. -1.34	3mth -1.3% -1.5% -16.0% Dec-25E 569 -42% -838 n.m. -2.77	12mth 152.5% 154.0% 116.8% Dec-26E 2,992 426% 1,248 -249% 4.13
Apr-24 麥科未源: Factset 绝对值 绝对值 (美元) 相对 MSCI China Rmb mn Revenue Revenue (+/-) Net profit Net profit (+/-) Diluted EPS (Rmb) GPM	Dec-23A 222 6% -926 n.m3.28 86.2%	1mth 19.4% 19.4% 17.5% Dec-24A 981 342% -406 n.m. -1.34 97.0%	3mth -1.3% -1.5% -16.0% Dec-25E 569 -42% -838 n.m. -2.77 90.1%	12mth 152.5% 154.0% 116.8% Dec-26E 2,992 426% 1,248 -249% 4.13 98.1%
Apr-24 <i>资料未源: Factset</i> 绝对值 绝对值(美元) 相对 MSCI China Rmb mn Revenue Revenue (+/-) Net profit Net profit (+/-) Diluted EPS (Rmb)	Dec-23A 222 6% -926 n.m. -3.28	1mth 19.4% 19.4% 17.5% Dec-24A 981 342% -406 n.m. -1.34	3mth -1.3% -1.5% -16.0% Dec-25E 569 -42% -838 n.m. -2.77	12mth 152.5% 154.0% 116.8% Dec-26E 2,992 426% 1,248 -249% 4.13

(Please see APPENDIX 1 for English summary)

事件

2024 年公司实现营收 9.8 亿元(+343%),其中包括武田 1 亿美元选择权付款,耐立克实现营收 2.4 亿元(+52%)。销售和分销开支 2.0 亿元(+0.3%),研发开支 9.5 亿元(+34%),行政开支 1.9 亿元(+3.3%)。亏损 4.1 亿元(同比收窄-56.2%)。在手现金和存款余额 12.6 亿元(+15.3%)。公司在 2025 年完成美股 IPO,募资 1.3 亿美元(9.67 亿元)。

点评

耐立克国内收入表现优异。耐立克实现营收2.4亿元(+52%),其中 H2 收入 1.3 亿元(+149%),主要由于药店、医院渗透率提高以及纳入医保目录后患者可及性增加。截至24年底,耐立克的全国准入医院和DTP药房共达到734家,准入医院同比增长86%。我们预计随着耐立克的所有适应症纳入医保,并且患者治疗持续时间(DOT)不断延长,其市场需求将持续释放,实现增长。

公司近期研发进展梳理:

耐立克: 1) Ⅲ 期研究 POLARIS-2(经治 CML-CP)进行中; 2)Ⅲ 期研究 POLARIS-3(SDH-GIST)进行中; 3)Ⅲ 期研究 POLARIS-1(1L Ph+ ALL)Ⅲ 期研究入组中, 计划向美国 FDA 提交 Ⅲ 期申请。

APG-2575 (BCL-2): 1) 单药治疗 R/R CLL/SLL 的 NDA 获 CDE 受理。2) III 期研究 GLORA-4 (1L MDS) 进行中,计划向 FDA 提交 III 期申请;3) III 期研究 GLORA-3 (1L AML) 进行中;4) III 期研究 GLORA-2 (1L CLL/SLL) 进行中。

APG-2449 (ALK/ROS1 TKI): 1)二代 ALK TKI 耐药/不耐受的 NSCLC: 2)初治 ALK 阳性 NSCLC。

我们认为,耐立克授权武田的交易以及公司美股上市融资有效缓解了公司的现金的需求。建议关注公司后续临床推进和执行情况,以及 FDA 是否批准 APG-2575 用于 1L MDS 的注册性 III 期研究以及海外授权进展等关键催化事件。

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盈利预测及估值建议:

我们预计公司 FY25-27 营收分别为 5.7/29.9/28.0 亿元(不变)。对应 FY25-27 的净利 润为-8.4/12.5/9.1 亿元(FY25-27 前值: -8.5/12.4/9.3 亿元)。我们使用经风险调整的贴现现金流(DCF)模型及 2025-2033 财年的现金流预测对该公司进行估值。基于 WACC 10.0%,永续增长率 3.0%,对应目标价 53.1 HKD/股(前值: 48.0 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, 新药临床推进和商业化不及预期风险。

图1 公司近期里程碑事件梳理 推进POLARIS-2试验患者入组 经FDA批准的、单药治疗CML的注册三期临床试验 推进POLARIS-1试验患者入组 联合化疗治疗1线Ph+ ALL 患者的注册三期临床试验 奥雷巴替尼 推进POLARIS-3试验患者入组 (HQP1351) 单药治疗SDH缺乏症GIST患者的注册三期临床试验 争取FDA批准在美国开展1线治疗Ph+ ALL患者的注册临床试验 武田行使独家选择权协议 计划在2025年在中国商业化,针对r/r CLL 推进GLORA试验患者入组 经FDA批准的注册三期临床试验,联合BTK抑制剂治疗既往接受过BTK 抑制剂治疗的CLL/SLL 推进GLORA-2试验患者入组 联合阿可替尼治疗新诊断的CLL/SLL的注册三期临床试验 Lisaftoclax (APG-2575) 推进GLORA-3试验患者入组 联合AZA治疗新诊断年老或体弱AML的注册三期临床试验 推进GLORA-4试验患者入组 联合AZA治疗新诊断 HR MDS的注册三期临床试验 争取FDA批准在美国开展针对MDS患者的注册临床试验

资料来源:公司公开演示材料; HTI

表 1. DCF 估值模型

DCF Valuation (CNY mn)	2023	2024A	2025E	2026E	2027E	2028E	2029E	2030E	2031E	2032E	2033E
Sales	222	981	569	2,992	2,801	2,080	2,771	1,690	1,887	2,137	2,269
y-y growth		341.8%	-42.0%	425.6%	-6.4%	-25.7%	33.2%	-39.0%	11.7%	13.2%	6.2%
Gross profit	191	952	513	2,934	2,723	1,961	2,610	1,494	1,667	1,895	2,011
y-y growth		397.1%	-46.1%	472.0%	-7.2%	-28.0%	33.0%	-42.8%	11.6%	13.7%	6.1%
EBIT		(288)	(717)	1,383	1,047	390	1,180	298	918	1,466	1,978
Tax rate		15.0%	15.0%	15.0%	15.0%	15.0%	15.0%	15.0%	15.0%	15.0%	15.0%
EBIT*(1-tax rate)		(245)	(609)	1,175	890	331	1,003	253	780	1,246	1,681
+ D&A		82	56	57	58	59	60	61	62	63	64
- Change in working capital		92	(47)	(87)	44	(141)	67	(92)	(146)	136	(122)
- Capx		(92)	(72)	(74)	(75)	(77)	(78)	(79)	(81)	(82)	(84)
FCFF		(163)	(673)	1,072	916	173	1,052	142	616	1,363	1,540
Terminal value											22,527
FCF + Terminal value		(163)	(673)	1,072	916	173	1,052	142	616	1,363	24,066
Discount factor				0.93	0.85	0.77	0.70	0.64	0.58	0.52	0.48
PV of FCF + Terminal value		-	-	998	775	133	735	90	355	715	11,471
WACC	10.0% T e	10.0% Terminal growth rate 3.		3.0%		Р	Present value of enterprise (CNY mn)				15,273
Cost of Equity	11.7%					-	Net debt (CN	IY mn)			(407)
Cost of Debt	4.0%					-	MI (CNY mn)				(10)
Equity Beta	1.10					Е	quity value (CNY mn)			14,855
Risk Free Rate	1.8%					N	lo. of shares				302
Market Risk Premium	9.0%					D	CF per share	(CNY)			49.18
Target Debt to Asset ratio	20%					С	NY/HKD				1.08
Effective Corporate Tax Rate	15.0%					D	CF per share	(HKD)			53.11

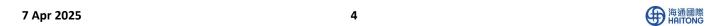
资料来源:公司年报,HTI

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表 2. 财务报表和盈利预测

Key financials	FY2024A	FY2025F	FY2026F	FY2027F	Profit & Loss (CNY mn)	FY2024A	FY2025F	FY2026F	FY2027F
Revenue (CNY mn)	981	569	2,992	2,801	Total turnover	981	569	2,992	2,801
Operating Profit /Loss (CNY mn)	(288)	(717)	1,383	1,047	Cost of sales	(29)	(56)	(58)	(78)
Pre-tax profit / Loss (CNY mn)	(395)	(844)	1,258	913	Gross profit	952	513	2,934	2,723
Net income to ord equity (CNY mn)	(405)	(838)	1,248	905	Total operating costs	(1,240)	(1,230)	(1,551)	(1,676)
Revenue growth	341.8%	-42.0%	425.6%	-6.4%	Operating profit	(288)	(717)	1,383	1,047
Net profit growth	-56.2%	106.5%	-249.0%	-27.5%	Operating EBITDA	(370)	(773)	1,326	989
ROE	-235.3%	-304.2%	138.6%	45.8%	Depreciation and amortisation	(82)	(56)	(57)	(58)
Balance Sheet (CMY mm)	FY2024A	FY2025F	FY2026F	FY2027F	Operating EBIT	(288)	(717)	1,383	1,047
Total cash and equivalents	1,261	1,362	2,581	3,592	Interest income (expense)	(64)	(124)	(131)	(139)
Inventories	7	39	7	55	Pre-tax profit	(395)	(844)	1,258	913
Account and other receivables	83	222	67	304	Taxation	(10)	7	(10)	(7)
Trade receivables	N.A.	N.A.	N.A.	N.A.	Net Income	(406)	(838)	1,248	905
Other current assets	123	92	94	96	Minorities	(0)	(0)	0	0
Total current assets	1,474	1,715	2,749	4,046	Net Income to ord equity	(405)	(838)	1,248	905
Property, plant and equipment	849	862	2,743 875	888	One-off expense	N.A.	N.A.	N.A.	N.A.
	294	273	282	292	Normalized net income	N.A.	N.A.	N.A.	N.A.
Other non-current assets Total non-current assets	1,144	1,135	1,157	1,180					
Total assets	2,618	2,850	3,906	5,226	Per Share Data	FY2024A	FY2025F	FY2026F	FY2027F
Contract liabilities	2,618 37	2,850 38	3,906 39	40	EPS (CNY)	(1.3)	(2.8)	4.1	3.0
					Revenue per share (CNY)	3.2	1.9	9.9	9.3
Trade and other payable	258	263	269	274	Operating EBITDA per share (CNY)	(1.2)	(2.6)	4.4	3.3
Bank borrowing	779	826	875	928	BVPS (CNY)	0.9	0.9	5.0	8.0
Other current liabilities	92	216	(57)	272	DPS (CNY)	N.A.	N.A.	N.A.	N.A.
Total current liabilities	1,167	1,344	1,126	1,513	Recurrent cash flow per share (CNY)	(0.5)	0.0	4.0	3.3
Bank borrowing	248	263	279	296	Shares in issue (million) Year end adjusted shares in issue	302	302	302	302
Contract liabilities	N.A.	N.A.	N.A.	N.A.	(million)	N.A.	N.A.	N.A.	N.A.
Other liabilities	929	966	977	987	Kara Bartlan	EV20244	FV202FF	FW202CF	FV2027F
Total non-current liabilities	1,177	1,230	1,256	1,283	Key Ratios	FY2024A	FY2025F	FY2026F	FY2027F
Total liabilities	2,344	2,573	2,381	2,796	Growth	244.00/	42.00/	425.60/	C 40/
Shareholder's equity	264	267	1,515	2,420	Revenue growth	341.8%	-42.0%	425.6%	-6.4%
Minority interests	10	10	10	10	Operating profit growth	-65.4%	148.9%	-292.9%	-24.3%
Total equity	274	277	1,525	2,430	Net profit growth	-56.2%	106.5%	-249.0%	-27.5%
Total liabilities & shareholders' equity	2,618	2,850	3,906	5,226	Margins	07.00/	22.42/	00.40/	07.00/
Cash flow (CNY mn)	FY2024A	FY2025F	FY2026F	FY2027F	Gross margin	97.0%	90.1%	98.1%	97.2%
Operating profit	(288)	(717)	1,383	1,047	Operating EBITDA margin	-37.7%	-135.8%	44.3%	35.3%
Deprecation and amortisation	(82)	(56)	(57)	(58)	Operating margin	-29.4%	-125.9%	46.2%	37.4%
Changes in working capital	(47)	(87)	44	(141)	Pretax profit margin	-40.3%	-148.3%	42.1%	32.6%
Other operating cash flow	306	154	(21)	297	Tax rate	-0.8%	-0.8%	-0.8%	-0.8%
Cash generated from operations	(111)	(706)	1,349	1,145	Net profit margin	-41.4%	-147.1%	41.7%	32.3%
Capex	(76)	(71)	(72)	(73)	Key Ratios				
Other investing cash flow	(286)	15	15	16	ROE	-235.3%	-304.2%	138.6%	45.8%
other myesting cush now	(200)	13	13	10	ROA	-15.9%	-30.6%	37.0%	19.8%
Net cash flow from investing activities	(362)	(56)	(57)	(57)	Capex/revenue	-7.8%	-12.4%	-2.4%	-2.6%
Change in borrowings	(127)	56	59	62	• •				
Proceeds from changes in capital	#REF!	840	-	-	Current ratio (x)	1.3	1.3	2.4	2.7
Other financing cash flow	#REF!	(124)	(131)	(139)	Creditor days Debtor days	800.0 110.0	1000.0 105.0	500.0 100.0	500.0 95.0
	315	772	(73)	(78)	Debitor days	110.0	103.0	100.0	33.0
Net cash flow from financing activities	313	879	889	2,109	Inventory days	146.0	145.0	160.0	158.0
	1 038		000	2,100	Sales/avg assets	0.4	0.2	0.9	0.6
Cash at beginning of period	1,038 (159)			1 011					
Cash at beginning of period Net change in cash	(159)	10	1,219	1,011	Credit analysis				
Net cash flow from financing activities Cash at beginning of period Net change in cash Forex effects	(159)	10 -	1,219 -	-	· · ·	-2.8	-1.4	0.9	1.2
Cash at beginning of period Net change in cash	(159)	10	1,219		Credit analysis			0.9 76%	1.2 50%

资料来源: HTI



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APPENDIX 1

Summary

Events. In 2024, the company achieved revenue of RMB 980 million (+343% YoY), including a \$100 million option payment from Takeda, while HQP1351 generated revenue of RMB 240 million (+52% YoY). Selling and distribution expenses were RMB 200 million (+0.3% YoY), R&D expenses reached RMB 950 million (+34% YoY), and administrative expenses amounted to RMB 190 million (+3.3% YoY). The company reported a net loss of RMB 410 million, narrowing by 56.2% YoY, with a cash and deposit balance of RMB 1.26 billion (+15.3% YoY). In 2025, the company completed its U.S. IPO, raising RMB 967 million.

Comments. HQP1351 delivered strong domestic revenue performance, reaching RMB 240 million (+52% YoY), with H2 revenue of RMB 130 million (+149%), driven primarily by increased penetration in pharmacies and hospitals, as well as improved patient accessibility following its inclusion in the national reimbursement drug list. By the end of 2024, HQP1351 had secured market access in 734 hospitals and DTP pharmacies nationwide, with the number of hospitals increasing by 86% YoY.

Recent R&D Progress Overview:

HQP1351:

- 1) Phase III trial POLARIS-2 (pretreated CML-CP) is ongoing.
- 2) Phase III trial POLARIS-3 (SDH-GIST) is ongoing.
- 3) Patient enrollment is underway for the Phase III trial POLARIS-1 (1L Ph+ ALL), with plans to submit a Phase III application to the U.S. FDA.

APG-2575:

- 1) NDA for monotherapy in R/R CLL/SLL has been accepted by the CDE.
- 2) Phase III trial GLORA-4 (1L MDS) is ongoing, with plans to submit a Phase III application to the FDA.
- 3) Phase III trial GLORA-3 (1L AML) is ongoing.
- 4) Phase III trial GLORA-2 (1L CLL/SLL) is ongoing.

APG-2449 (ALK/ROS1 TKI):

- 1) NSCLC patients resistant or intolerant to second-generation ALK TKIs.
- 2) Treatment-naïve ALK-positive NSCLC.

We think that the licensing deal between Ascentage Pharma and Takeda, along with the company's U.S. IPO financing, has effectively alleviated its cash needs. We recommend closely monitoring the company's clinical progress and execution, as well as key catalysts such as FDA approval for the registrational Phase III trial of APG-2575 in 1L MDS and the progress of license-out deal.

Earnings Forecast and Valuation. We estimate the company's FY25-27 revenue to be RMB 5.7/29.9/28.0 billion (unchanged), with corresponding net profit of RMB -8.4/12.5/9.1 billion (previous estimates: RMB -8.5/12.4/9.3 billion). We value the company using a risk-adjusted discounted cash flow (DCF) model based on cash flow projections for FY25-33. Applying a WACC of 10.0% and a terminal growth rate of 3.0%, we arrive a target price of HKD 53.1 per share (previously HKD 48.0), maintaining our "Outperform" rating.

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Risks: New drug development risk, regulatory approval risk, and commercialization underperformance risk.

APPENDIX 2

ESG Comments

Environmental:

overall good

Social:

overall good

Governance:

overall good



附录 APPENDIX

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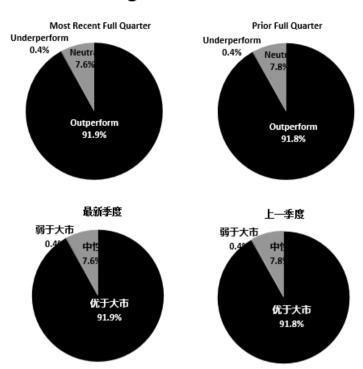
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		(持有)				
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各地股票基准指数: 日本-TOPIX,韩国-KOSPI,台湾-TAIEX,印度-Nifty100;其他所有中国概念股-MSCI China.

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		(hold)	
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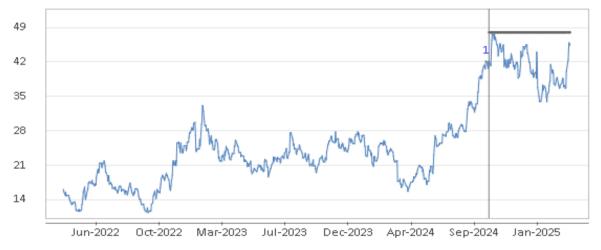
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Ascentage Pharma - 6855 HK



1. 11 Oct 2024 OUTPERFORM at 40.70 target 48.00.

Source: Company data Bloomberg, HTI estimates

