Hong Kong

招股概略

上海复宏汉霖生物技术股份有限公司 (2696)

发行数据

集资额:	3,208.9 –3,739.4 百万港元
发售股份数目:	64,695,400 股 H 股
价格:	49.60 –57.80 港元
每手股数:	100
入场费:	5,838.25 港元
历史市盈率:	不适用
每股有形资产净值:	6.09 –7.04 港元
市值(招股后):	7,889.5 –9,193.8 百万港元
招股开始:	2019年9月12日
招股截止:	2019年9月17日中午12时
上市:	2019年9月25日
	中国国际金融香港证券有限公司、Merrill Lynch Far East Limited、招银国际融资有限公司、复星恒利证
保荐人:	券有限公司及花旗环球金融亚洲有限公司

12月 31 日止年度	人民币千元	按年变动
来自持续经营业务的收入		
2017 年	33,910	N/A
2018年	7,421	-78.1%
年内 / 期间利润		
2017年	-384,327	N/A
2018年	-504,789	31.3%

背景

- 集团是一家中国领先的生物制药公司,旨在为全球患者提供质高价优的创新药物。
- 集团是首家根据现行中国有关生物类似药评价及上市审批的法规《生物类似药指导原则》,就单克隆抗体生物类似药自国家药监局取得新 药药证申请批准的生物制药公司,且亦为中国首家商业化推出生物类似药产品的生物制药公司。
- 于最后实际可行日期,除集团已商业化推出的生物类似药产品(即 HLX01 汉利康)外,集团在管线中自主开发 20 种以上候选生物药及 多种肿瘤免疫联合疗法,其中

(i) 两种候选单抗获国家药监局接纳新药药证申请,包括一种候选单抗获欧洲药品管理局接纳营销授权申请;

(ii) 两种候选单抗正在进行 3 期临床试验及六种候选单抗正在进行 1/2 期临床试验,以及两种肿瘤免疫联合疗法正在进行 3 期临床试验;及

(iii) 已在不同司法权区取得 31 项新药临床试验申请批准。

业务策略

- 进一步巩固集团在生物类似药持续开发方面的领导地位,并把握先发优势;
- 利用集团强大而全面的生物药物管线及成熟的单抗开发平台,开发以肿瘤免疫联合疗法为主的创新产品组合;
- 在维持高质量标准的同时扩大产能及提高成本效益;
- 通过内部销售及市场团队及合作伙伴关系提升商业化能力;及
- 选择性地进行战略合作,以扩大集团的全球版图。

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竞争优势

- 具有吸引力的商业模式: 兼备近期可实现性及长期增长性;
- 高效的一体化全球研发平台,开发出强大的产品管线;
- 具有加强成本效益和健全质量管理体系的大规模生产能力;
- 强大的全球商业化能力;及
- 远见卓识的联合创始人及领导团队。

主要风险

- 与集团的财务前景及额外资金需求有关的风险;
- 与集团候选药物开发、临床试验及监管批准有关的风险;
- 与集团的候选药物商业化有关的风险;
- 与知识产权有关的风险;
- 与集团的经营有关的风险;
- 与在中国经营业务有关的风险;及
- 与全球发售有关的风险

股息政策

▪ 无固定股息政策。

所得款项用途

	(百万港元)	比重
将用于拨付持续进行的核心产品的临床试验、监管备案及注册	1,341.7	40.0%
将用于拨付持续进行的候选生物类似药(包括 HLX12、HLX11 及 HLX14)的临床试验、监管备案及注册	503.1	15.0%
将用于拨付持续进行的生物创新药的临床试验、监管备案及注册以及开 发免疫肿瘤联合疗法	1,174.0	35.0%
将分配至营运资金及一般企业用途	335.4	10.0%
合计:	3,354.1	100.0%

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