

3/18/2019

招股概略

康希諾生物股份公司 (6185 HK)

發行數據

集資額: 1,202.2 - 1,259.5 百萬港元

發售股份數目: 57,248,600 股 H 股 **價格:** 21.00- 22.00 港元

每手股數: 200

入場費: 4,444.34 港元

歷史市盈率: 不適用

每股有形資產淨值: 7.62-7.88 港元

市值(招股後): 2,692.6-2,820.9 百萬港元 (H 股)

招股開始: 2019年3月18日

招股截止: 2019年3月21日中午12時

上市: 2019年3月28日

保薦人: 摩根士丹利亞洲有限公司及中信里昂證券資本市場有限公司

12月31日止年度	人民幣千元	
經營虧損		
2017年	-63,796	21.1%
2018年	-138,578	117.2%
年內虧損及全面虧損總額		
2017年	-64,450	29.3%
2018年	-138,281	114.6%

背景

- 康希諾的使命是研發、生產和商業化高質量、創新及經濟實惠的疫苗。
- 肩負此使命的,是集團傑出的創始人和高級管理層團隊,包括多位擁有於賽諾菲巴斯德、阿斯利康和惠氏(現為輝瑞)等全球製藥公司領導創新國際疫苗研發經驗的世界頂尖科學家。其他管理層成員均來自領先的跨國及國內生物製藥公司,亦為疫苗行業的資深人士。
- 集團的疫苗產品線在戰略上旨在針對中國龐大且供不應求的市場,可總結為三個類別:
 - (i)全球創新疫苗(如 Ad5-EBOV、集團的在研結核病加強疫苗及在研 PBPV),以迎合中國尚未滿足的醫療需求;
 - (ii)研發潛在的中國首創疫苗,憑藉質量更高的世界級疫苗取代現時的主流疫苗(如集團的在研 DTcP 疫苗及在研 MCV4);及(iii)研發出與中國市場的進口產品競爭的中國潛在最佳疫苗(如集團的在研 PCV13i)。
- 集團正為 12 個疾病領域研發 15 種在研疫苗。除了集團預防腦膜炎球菌感染及埃博拉病毒病的三項臨近商業化疫苗產品之外,集團有六種在研疫苗處於臨床試驗階段或臨床試驗申請階段。集團亦有六種臨床前在研疫苗,包括一種在研聯合疫苗。
- 為籌備不久將來的商業化生產,集團已建成一座位於天津的生產廠房,其設計、驗證和運營均達到國際標準。該廠房目前使集團的年原液產能達到約7,000萬至8,000萬劑,集團相信該廠房在可預見未來將充分支持集團臨近商業化在研疫苗的商業化計劃以及支持集團臨床試驗材料的生產。集團對複雜的疫苗生產工藝關鍵步驟擁有重要的專業知識。

業務策略

- 推進臨近商業化疫苗產品的研發和商業化;
- 迅速推動其他在研疫苗產品線的開發;
- 建立並鞏固集團的商業化基礎設施;及
- 通過全球合作及收購機會鞏固集團的優勢。



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競爭優勢

- 臨近商業化資產的巨大潛力;
- 全面且強大的疫苗產品線以應付供不應求的龐大市場;
- 先進的疫苗研發平台技術;
- 達全球標準的疫苗生產能力及質量管理體系;及
- 來自全球領先的生物製藥公司的世界頂尖科學和管理團隊。

主要風險

- 自成立以來,集團已蒙受重大虧損,並預計在未來幾年將繼續虧損,並有可能永遠無法達致或維持盈利能力;
- 集團的財務前景取決於集團的臨床階段及臨床前階段疫苗產品線獲成功開發及批准;
- 集團可能面臨疫苗市場的重大競爭; (iv)集團可能無法獲得在研疫苗的監管批准;
- 集團可能無法獲得目標省份省級疾控中心的預審資格或取得其後的產品訂單;及
- 集團的任何疫苗及在研疫苗的商業成功取決於接種者、國家及地方疫控中心、KOL 及疫苗或疾病預防行業其他人士的市場認可度。

股息政策

■ 無固定股息收入。

所得款項用途

	(百萬港元)	比重 (%)
將用於集團核心產品以及集團疫苗產品線中其他主要產品的研發及商業 化	898.70	80.0%
將用於集團臨床前在研疫苗的持續研發	112.40	10.0%
將用於營運資金及其他一般企業用途	112.40	10.0%
合計	1,123.50	100.00%



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