

IPO FACT SHEET
Innovent Biologics, Inc. (1801 HK)
ISSUE STATISTICS

Offer Size:	HK\$2,954.4m – HK\$3,308.9m
Placement Tranche:	236.35m
Price:	HK\$12.50 – HK\$14.00
Board lot:	500
Entry fee:	HK\$7,070.54
Historical PE	N/A
Net tangible asset per share:	HK\$4.54 – HK\$4.84
Market Cap (post-IPO):	HK\$13.98b – HK\$15.65b
Open:	18 Oct 2018
Close:	12.00 noon on 23 Oct 2018
Trading:	31 Oct 2018
Sponsor:	Morgan Stanley Asia Limited, Goldman Sachs (Asia) L.L.C., J.P. Morgan Securities (Far East) Limited and China Merchants Securities (HK) Co., Limited

Year ended 31 Dec	(RMB'000)	yoy % chg
Revenue		
2016	-	N/A
2017	18,538	N/A
(Loss) profit and total comprehensive (expenses) income for the year/period attributable to owners of the Company		
2016	-504,204	N/A
2017	-562,318	11.5%

BACKGROUND

- They were founded in 2011 by their visionary leader, Dr. De-Chao Michael Yu, a highly accomplished scientist, innovator and entrepreneur. Dr. Yu invented the world's first oncolytic virus-based immunotherapeutic product, Oncorine, and also co-invented and led the development of the first domestic innovative fully human antibody-like therapeutic approved for marketing in China, Conbercept.
- They have developed their fully integrated platform which boasts advanced research, discovery, development, manufacturing and commercialization capabilities. These capabilities have enabled them to build a robust pipeline of innovative and commercially promising monoclonal antibodies and other biologics in the fields of oncology, ophthalmology, and autoimmune and metabolic diseases.
- Leveraging their platform, they have built up a pipeline of 17 antibody drug candidates in the last seven years, led by their four core products that are in late-stage clinical development in China, including sintilimab (IBI-308), their novel PD-1 antibody; IBI-305, their bevacizumab (Avastin) biosimilar; IBI-301, their rituximab (MabThera/Rituxan) biosimilar; and IBI-303, their adalimumab (Humira) biosimilar. In addition, out of their pipeline of 17 antibody drug candidates, six are in clinical development in China. Four other drug candidates in their pipeline.
- They have three bi-specific monoclonal antibody candidates based on sintilimab (IBI-308) in co-development with Eli Lilly, two of which, IBI-318 and IBI-319, are under development in China.
- In addition to developing their pipeline drug candidates in China, they have obtained FDA approval for their IND applications for sintilimab (IBI-308) and IBI-188 and plan to initiate a multi-center Phase 1b/2 clinical trial for sintilimab (IBI-308) and a Phase 1a clinical trial for IBI-188 in the U.S.

BUSINESS STRATEGY AND FUTURE PLANS

- Expedite regulatory approval and commercialization of their lead product candidates
- Rapidly advance their clinical programs for pipeline products
- Continue to enhance their fully-integrated platform
- Maximize the value of their fully-integrated platform through a global strategy of organic growth and collaboration

COMPETITIVE STRENGTHS

- Fully-integrated biological therapeutics platform
- Potentially best-in-class innovative PD-1 monoclonal antibody with NDA accepted and priority review status granted by the NMPA
- Three biosimilar drug candidates in Phase 3 clinical trials in China
- Robust pipeline of innovative monoclonal antibody and bi-specific antibody drug candidates
- State-of-the-art manufacturing facilities designed to, built to and operating at international standards
- Strategic partnerships with leading global companies, such as Eli Lilly and Adimab
- Senior management with a proven track record of success, led by their co-founder, the co-inventor and developer of the first innovative fully human antibody-like drug in China

KEY RISKS

- The price and trading volume of their Shares could be volatile, which may lead to substantial losses to investors.
- They have incurred significant net losses since their inception and anticipate that they will continue to incur net losses for the foreseeable future and may never become profitable.
- They have a limited operating history, which may make it difficult to evaluate their current business and predict their future performance.
- They depend substantially on the success of their drug candidates, all of which are in pre-clinical or clinical development, and their ability to identify additional drug candidates. If they are unable to successfully identify new drug candidates, complete clinical development, obtain regulatory approval and commercialize their drug candidates, or experience significant delays in doing so, their business will be materially harmed.
- Clinical drug development involves a lengthy and expensive process with an uncertain outcome, and results of earlier studies and trials and non-head-to-head analyses (e.g., comparisons with competing drugs based on their publicly available study and trial data) may not be predictive of future trial results.
- They have no experience in launching and marketing drug candidates. If they are unable to further develop marketing and sales capabilities or enter into agreements with third parties to market and sell their drug candidates, they may not be able to generate product sales revenue.
- They face substantial competition, which may result in others discovering, developing or commercializing competing drugs before or more successfully than they do.
- If they are unable to obtain and maintain patent protection for their drug candidates through intellectual property rights, or if the scope of such intellectual property rights obtained is not sufficiently broad, third parties could develop and commercialize products and technologies similar or identical to their and complete directly against them, and their ability to successfully commercialize any product or technology may be adversely affected.
- Their rights to develop and commercialize our drug candidates are subject, in part, to the terms and conditions of licenses granted to them by others.
- Their future success depends on their ability to retain key executives and to attract, retain and motivate qualified personnel.

DIVIDEND POLICY

- No fixed dividend policy

USE OF PROCEEDS

	HK mn	As a percentage of gross proceeds from the Invitation (%)
To fund ongoing and planned clinical trials, preparation for registration filings and planned commercial launches (including sales and marketing) of sintilimab IBI-308, IBI-305, IBI-301 and IBI-303.	1,919.10	65.0%
To fund ongoing and planned clinical trials, preparation for registration filings and potential commercial launches (including sales and marketing) of the other drug candidates in their pipeline.	738.10	25.0%
For working capital and general corporate purposes.	295.20	10.0%
Total:	2,952.40	100.00%

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