

IPO FACT SHEET
TOT BIOPHARM International Company Limited (1875)
ISSUE STATISTICS

Offer Size:	HK\$589.5m – HK\$679.5m
Placement Tranche:	90m
Price:	HK\$6.55 – HK\$7.55
Board lot:	400
Entry fee:	HK\$3,050.43
Historical PE	N/A
Net tangible asset per share:	HK\$1.94 – HK\$2.10
Market Cap (post-IPO):	HK\$3,733.5m – HK\$4,303.5m
Open:	29 Oct 2019
Close:	12.00 noon on 1 Nov 2019
Trading:	8 Nov 2019
Sponsor:	ICBC International Capital Limited

Year ended 31 Dec	(RMB'000)	yoy % chg
Revenue		
2017	39,219	-24.0%
2018	9,218	-76.5%
Loss for the year/period attributable to the equity holders of the Company		
2017	-268,263	80.4%
2018	-70,290	-73.8%

BACKGROUND

- They are a clinical-stage biopharmaceutical company dedicated to developing and commercializing innovative oncology drugs and therapies.
- They have a comprehensive portfolio of oncology drug candidates, which include monoclonal antibodies (mAbs), antibody drug conjugates (ADCs), oncolytic virus products and specialty oncology drugs such as liposome drugs, targeting various types of cancers.
- They have built and established a fully integrated in-house platform of discovery, process development, quality management, pre-clinical and clinical development, as well as commercial-scale manufacturing facilities and proven sales and marketing capabilities, which provides flexibility and scalability for their business to expand along the innovative drug industry value chain.
- Their comprehensive product pipeline consists of seven biological and five chemical drug candidates, 11 of which are in-house developed. Their strategy is to develop innovative drugs that have high viability for commercialization and clear market demand. They focus on achieving a diverse product mix with a sustainable launch schedule, targeting to start from 2020.
- As of the Latest Practicable Date, they had four biological drug candidates in the clinical stage. Their most advanced biological drug candidate and Core Product, TAB008, is undergoing Phase III clinical trials. TAB008 is a bevacizumab biosimilar. Bevacizumab has been approved for the treatment of non-squamous non-small-cell lung cancer (nsNSCLC) and metastatic colorectal cancer (mCRC) in China. They currently expect to launch TAB008 between late 2020 and early 2021.

BUSINESS STRATEGIES

- Commercialize TAB008.
- Rapidly advance their clinical trials for drug candidates.
- Further enrich product portfolio via self-development and collaboration focusing on immune-oncology combination therapies and seeking innovative cancer treatment solutions.
- Strengthen their in-house sales and marketing force and commercial-scale manufacturing capacities.
- Continue to attract, train and retain quality talent to support their rapid growth and maximize the value of their integrated platform.

COMPETITIVE STRENGTHS

- Robust product pipeline with sustainable launch schedule, covering a wide variety of cancer types and extended applications.

- Well-established and advanced technology platforms focusing on oncology drugs.
- Cost-efficient commercial-scale and state-of-the-art manufacturing facilities, built to and operating at international standards.
- Proven open platform business model empowered by strong and integrated capabilities covering the full oncology drug industry value chain.
- Industry-leading, experienced and professional management team supported by a strong talent base.

KEY RISKS

- They depend substantially on the success of their drug candidates, all of which are undergoing pre-clinical or clinical development. If they are unable to successfully complete clinical development, obtain regulatory approval and commercialize our drug candidates, or experience significant delays in doing so, their business will be materially harmed.
- The regulatory approval processes of regulatory authorities are lengthy, time-consuming and inherently unpredictable. If they are ultimately unable to obtain regulatory approval for their drug candidates, their business will be substantially harmed.
- They currently do not generate revenue from sales of any drug products developed by us and may not become profitable in the foreseeable future, or at all.
- They have incurred significant net losses and net operating cash outflows since their inception, and they anticipate that they will continue to incur net losses and net operating cash outflows until they successfully commercialize their drug candidates.
- Their success depends on the ability to retain their research and development, manufacturing, clinical trial and sales and marketing team and other key executives, and to attract, train, retain and motivate qualified and highly skilled personnel.
- They have no experience in manufacturing their drug candidates on a large commercial scale, which is a highly exacting and complex process, and have not yet begun utilizing their manufacturing facilities for commercial purposes.
- They have limited experience in marketing drugs. If they are unable to develop sufficient capabilities to market and sell their drug candidates, they may not be able to generate product sales revenue.
- They may not be successful in developing, enhancing or adapting to new technologies and methodologies.
- If they are unable to obtain and maintain patent protection for their drug candidates, primarily their novel 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 theirs and compete directly against us, and their ability to successfully commercialize any product or technology may be adversely affected.
- If they are sued for infringing, misappropriating, or otherwise violating intellectual property rights of third parties or engaging in unfair competition, such litigation could be costly and time-consuming and could prevent or delay us from developing or commercializing their drug candidates.
- They face substantial competition, which may result in others discovering, developing or commercializing competing drugs before or more successfully than they do.

DIVIDEND POLICY

- No fixed dividend policy.

USE OF PROCEEDS

	HK mn	As a percentage of gross proceeds from the Invitation
Will be used to fund ongoing and planned clinical trials, preparation for registration filings and the potential commercial launch (including sales and marketing) of TAB008.	166.3	30.0%
Will be used to fund ongoing and planned pre-clinical and clinical trials, expansion of facilities, preparation for registration filings and potential commercial launches (including sales and marketing) of the other drug candidates in their pipeline.	277.2	50.0%
Will be used for non-project specific capital expenditure.	83.2	15.0%
Will be used to fund continued expansion of their product portfolio in cancer and potentially other therapeutic areas through internal research and external licenses and business development collaborations.	11.1	2.0%
Will be used for their working capital and other general corporate purposes.	16.6	3.0%
Total:	554.4	100.0%

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