UNITED STATES SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

Form 8-K

Current Report Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

September 18, 2019

Date of Report (Date of earliest event reported)

<u>Chardan Healthcare Acquisition Corp.</u> (Exact Name of Registrant as Specified in its Charter)

Delaware	001-38762	82-3364020
(State or other jurisdiction	(Commission File Number)	(I.R.S. Employer
of incorporation)		Identification No.)
45 00 00 00 00 00		
17 State Street, 21st Floor		10004
New York, NY (Address of Principal Executive Office		(Zip Code)
(Address of Principal Executive Office	28)	(Zip Code)
Registr	rant's telephone number, including area code: (646) 465-	9000
	N/A	
(For	rmer name or former address, if changed since last repor	rt)
Check the appropriate box below if the Form 8-K filing is int	tended to simultaneously satisfy the filing obligation of	the registrant under any of the following provisions:
☐ Written communications pursuant to Rule 425 under	r the Securities Act	
Soliciting material pursuant to Rule 14a-12 under the	e Exchange Act	
☐ Pre-commencement communications pursuant to Ru	ıle 14d-2(b) under the Exchange Act	
☐ Pre-commencement communications pursuant to Ru	ıle 13e-4(c) under the Exchange Act	
Securities registered pursuant to Section 12(b) of the Act:		
Title of each class	Trading Symbols	Name of each exchange on which registered
Units, each consisting of one share of common stock, \$0.00	±	
one Warrant	CHAC.U	NYSE American
Common stock, \$0.0001 par value per share		NYSE American
Warrants to purchase common stock	CHAC.WS	NYSE American
Indicate by check mark whether the registrant is an emerging Securities Exchange Act of 1934 (17 CFR §240.12b-2).	growth company as defined in Rule 405 of the Securitie	es Act of 1933 (17 CFR §230.405) or Rule 12b-2 of the
Emerging growth company ⊠		
If an emerging growth company, indicate by check mark if th accounting standards provided pursuant to Section 13(a) of the		n period for complying with any new or revised financial

IMPORTANT NOTICES

Participants in the Solicitation

BiomX Ltd. ("BiomX"), Chardan Healthcare Acquisition Corp. ("CHAC"), and their respective directors, executive officers and employees and other persons may be deemed to be participants in the solicitation of proxies from the holders of CHAC common stock in respect of the proposed transaction described herein. Information about CHAC's directors and executive officers and their ownership of CHAC's common stock is set forth in CHAC's Prospectus dated December 14, 2018 filed with the SEC, as modified or supplemented by any Form 3 or Form 4 filed with the Securities and Exchange Commission (the "SEC") since the date of such filing. Other information regarding the interests of the participants in the proxy solicitation will be included in the proxy statement pertaining to the proposed transaction when it becomes available. These documents can be obtained free of charge from the sources indicated below.

Additional Information and Where to Find It

In connection with the transaction described herein, CHAC will file relevant materials with the SEC, including a proxy statement on Schedule 14A. Promptly after filing its definitive proxy statement with the SEC, CHAC will mail the definitive proxy statement and a proxy card to each stockholder entitled to vote at the special meeting relating to the transaction. INVESTORS AND SECURITY HOLDERS OF CHAC ARE URGED TO READ THESE MATERIALS (INCLUDING ANY AMENDMENTS OR SUPPLEMENTS THERETO) AND ANY OTHER RELEVANT DOCUMENTS IN CONNECTION WITH THE TRANSACTION THAT CHAC WILL FILE WITH THE SEC WHEN THEY BECOME AVAILABLE BECAUSE THEY WILL CONTAIN IMPORTANT INFORMATION ABOUT CHAC, BIOMX AND THE TRANSACTION. The definitive proxy statement, the preliminary proxy statement and other relevant materials in connection with the transaction (when they become available), and any other documents filed by CHAC with the SEC, may be obtained free of charge at the SEC's website (www.sec.gov) or by writing to Chardan Healthcare Acquisition Corp., 17 State Street, 21st Floor, New York, NY 10004.

Forward-Looking Statements

This Current Report on Form 8-K and the documents incorporated by reference herein (this "Current Report") contain certain "forward-looking statements" within the meaning of "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995. Forward-looking statements can be identified by words such as: "target," "believe," "expect," "will," "shall," "may," "anticipate," "estimate," "would," "positioned," "future," "forecast," "intend," "plan," "project" and other similar expressions that predict or indicate future events or trends or that are not statements of historical matters. Examples of forward-looking statements include, among others, statements made in this Current Report regarding the proposed transactions contemplated by the merger agreement (the "Merger Agreement") among CHAC, CHAC Merger Sub Ltd. and BiomX (the "Merger"), including the anticipated initial enterprise value and post-closing equity value, the benefits of the Merger, integration plans, expected synergies and revenue opportunities, anticipated future financial and operating performance and results, including estimates for growth, the expected management and governance of the combined company, and the expected timing of the Merger. Forward-looking statements are neither historical facts nor assurances of future performance. Instead, they are based only on CHAC and BiomX managements' current beliefs, expectations and assumptions. Because forward-looking statements relate to the future, they are subject to inherent uncertainties, risks and changes in circumstances that are difficult to predict and many of which are outside of our control. Actual results and outcomes may differ materially from those indicated in the forward-looking statements. Therefore, you should not rely on any of these forward-looking statements. Important factors that could cause actual results and outcomes to differ materially from those indicated in the forward-looking statements include, among others, the following: (1) the occurrence of any event that could give rise to the termination of the Merger Agreement; (2) the outcome of any legal proceedings that may be instituted against CHAC, the combined company, or others following the announcement of the Merger and the Merger Agreement; (3) the inability to complete the Merger due to the failure to obtain approval of CHAC's stockholders or to satisfy other conditions to closing in the Merger Agreement; (4) changes to the proposed structure of the Merger that may be required or appropriate as a result of applicable laws; (5) the ability to meet NYSE American listing standards following the consummation of the Merger; (6) the risk that the Merger disrupts current plans and operations of BiomX as a result of the announcement and consummation of the Merger; (7) the ability to recognize the anticipated benefits of the Merger, which may be affected by, among other things, competition, the ability of the combined company to grow and manage growth profitably, maintain relationships with third parties and partners, obtain adequate supply of raw materials and retain its management and key employees; (8) costs related to the Merger; (9) changes in applicable laws or regulations; (10) the possibility that BiomX or the combined company may be adversely affected by other economic, business, regulatory, and/or competitive factors; (11) BiomX estimates of expenses; (12) the impact of foreign currency exchange rates and interest rates fluctuations on the results of BiomX or the combined company; and (13) other risks and uncertainties indicated in the proxy statement of CHAC to be filed by CHAC with the SEC in connection with the Merger, including those under "Risk Factors" therein, and other documents filed or to be filed from time to time with the SEC by CHAC.

A further list and description of risks and uncertainties can be found in CHAC's Prospectus dated December 14, 2018 filed with the SEC and in the proxy statement on Schedule 14A that will be filed with the SEC by CHAC in connection with the proposed transaction, and other documents that the parties may file or furnish with the SEC, which you are encouraged to read. Any forward-looking statement made by us in this Current Report is based only on information currently available to CHAC and BiomX and speaks only as of the date on which it is made. CHAC and BiomX undertake no obligation to publicly update any forward-looking statement, whether written or oral, that may be made from time to time, whether as a result of new information, future developments or otherwise, except as required by law.

Item 7.01 Regulation FD Disclosure

Attached as Exhibit 99.1 to this Current Report on Form 8-K and incorporated into this Item 7.01 by reference is the investor presentation dated September 2019 that will be used by Chardan Healthcare Acquisition Corp. ("CHAC") in making presentations to certain existing and potential stockholders of CHAC with respect to the proposed transaction with BiomX Ltd. ("BiomX").

Item 9.01. Financial Statements and Exhibits

(d) Exhibits.

Exhibit No. Description

99.1* <u>Investor Presentation dated September 2019</u>

*Furnished but not filed.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

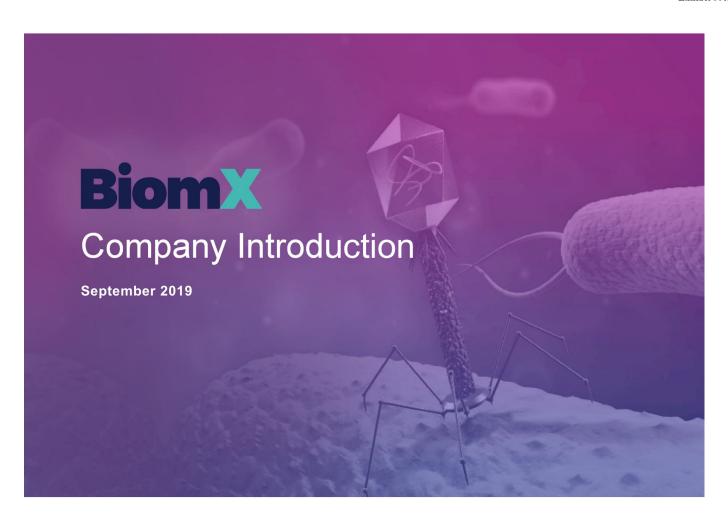
Dated September 18, 2019

CHARDAN HEALTHCARE ACQUISITION CORP.

/s/ Jonas Grossman Jonas Grossman By:

Name:

Title: President and Chief Executive Officer



Disclaimer

This investor presentation ("Investor Presentation") is for informational purposes only and does not constitute an offer to sell, solicitation of an offer to buy, or a recommendation to purchase any equity, debt or other financial instruments of Chardan Healthcare Acquisition ("CHAC") or BiomX Ltd. ("BiomX") or any of BiomX or CHAC's affiliate securities (as such term is defined under the U.S. federal securities laws). The information contained herein does not purport to be all-inclusive and is qualified in its entirety by the definitive merger agreement entered into July 16, 2019 filed by CHAC on a Current Report on Form 8-K on July 16, 2019. The data contained herein is derived from various internal and external sources. No representation is made as to the reasonableness of the assumptions made within or the accuracy or completeness of any other information contained herein. All levels, prices and spreads are historical and do not represent current market levels, prices or spreads, some or all of which may have changed since the issuance of this document. Any data on past performance, modeling contained herein is not an indication as to future performance. CHAC and BiomX assume no obligation to update the information in this Investor Presentation. Neither CHAC or BiomX accepts any liability whatsoever for any losses arising from the use of this Investor Presentation or reliance on the information contained herein. Nothing herein shall be deemed to constitute investment, legal, tax, financial, accounting or other advice. This Investor Presentation is being provided for use only by the intended recipient.

No representation or warranty (whether expressed or implied) has been made by CHAC, BiomX or any of their respective affiliates with respect to the matters set forth in this Investor Presentation, and the recipient disclaims any such representation or warranty. Only those particular representations and warranties of CHAC, BiomX or any of their respective affiliates made in a definite written subscription agreement, if any, regarding the matters set forth in this Investor Presentation (which will not contain any representation or warranty relating to this Investor Presentation or information contained in or omitted from this Investor Presentation) when and if executed, and subject to such limitations and restrictions as specified therein, shall have any legal effect. At any time upon the request of CHAC for any reason, recipient shall promptly deliver to CHAC or securely destroy this Investor Presentation and any other documents furnished to recipient by or on behalf of CHAC or BiomX without keeping any copies, in whole or part, thereof.



Disclaimer Continued

Forward-Looking Statements

This Investor Presentation includes "forward-looking statements" within the meaning of the "safe harbor" provisions of the US Private Securities Litigation Reform Act of 1995. Forward-looking statements can be identified by the use of words such as "forecast," "intend," "target," "believe," will," "expect," "estimate," "plan," and "project" and other similar expressions that predict or indicate future events or trends or that are not statements for historical information and other projections contained herein. Examples of forward-looking statements include, among others, statements made in this Investor Presentation regarding the revenues, earnings, performance, strategies, prospects and other aspects of the business of BiomX, CHAC or the combined company after completion of the proposed transaction ("the Business Combination"). Forward-looking statements are neither historical facts nor assurances of future performance. Instead, they are based only on management's current beliefs, expectations and assumptions. Because forward-looking statements relate to the future, they are esubject to inherent uncertainties, risks and changes in circumstances that are difficult to predict and many of which are outside of our control. Actual results and outcomes to differ materially from those indicated in the forward-looking statements. Therefore, you should not rely on any of these forward-looking statements include, among others, the following: (1) the occurrence of any event that could give rise to the termination of the merger agreement between CHAC and BiomX (the "Merger Agreement") with respect to the Business Combination; (2) the outcome of any legal proceedings that may be instituted against CHAC, the combined company, or others following the announcement of the Business Combination and the Merger Agreement; (3) the inability to complete the Business Combination disrupts current plans and operations of BiomX as a result of the ability to meet NYSE American listing standards following the consummation of the Business

Industry and Market Data

In this Investor Presentation, BiomX relies on and refers to information and statistics in the sectors in which it intends to compete. BiomX obtained this information and statistics from third-party sources believed to be reliable, including reports by market research firms. BiomX has supplemented this information where necessary with its own internal estimates, taking into account publicly available information about other industry participants and its management's best view as to information that is not publicly available. Neither Bioma Or CHAC has independently enrified the accuracy or completeness of any such third-party information.



Chardan Healthcare Investment Criteria



Venture-backed healthcare company in the biopharma or digital health sector



Public-company-ready management team



Actively considering public listing



\$200-\$500mm IPO valuation



Upcoming catalysts to drive valuation post-business combination



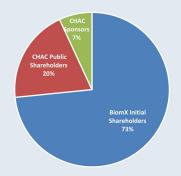
Transaction Overview: CHAC Merging with BiomX Ltd.

Business combination creates a leading publicly traded microbiome discovery company

Transaction Summary¹

- CHAC to merge with BiomX Ltd. to create a publicly-listed company focused on delivering novel microbiome-based technologies designed to improve the appearance of acne-prone skin² and treat conditions such as inflammatory bowel disease, primary sclerosing cholangitis, and cancer
- Pro forma valuation of \$254 million
- ~\$99 million of post-transaction cash
- BiomX securityholders to roll 100% of their vested equity holdings in the transaction, resulting in ~73% pro forma ownership post merger
- 16.625 million shares³ to be issued to BiomX securityholders at \$10.00 per share
- Earn-out provision provides certain shareholders of BiomX additional potential milestone-based equity consideration as follows:
 - 6,000,000 in total potential earn-out shares if the share price exceeds each of \$16.50, \$22.75 and \$29.00 by FYE21, FYE23 and FYE25, respectively⁴

Post Transaction Ownership





- (1) Assuming no redemptions from the CHAC shareholders.
- (2) For brevity, from here forward in this presentation we use the term "acne" to mean acne-prone skin.
- (3) Includes vested securities convertible into CHAC shares.
- (4) Based on VWAP during any 20 out of a 30-day period.

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Transaction Summary¹

Pro forma valuation	
Illustrative share price (per share)	\$10.00
Vested securities outstanding (million)	25.375
Equity Value	\$253,750,000
Estimated post-close cash on Balance Sheet	\$99,000,000

Sources of Funds	
CHAC Cash in Trust	\$70,000,000
BiomX Shareholder Equity Rollover	\$166,250,000
Sponsor Promote	\$17,500,000
Total Sources	\$253,750,000

Uses of Funds	
Equity Issued to BiomX Shareholders	\$166,250,000
Additional cash to BiomX Balance Sheet	\$69,000,000
CHAC Estimated Transaction Costs	\$1,000,000
Sponsor Promote	\$17,500,000
Total Uses	\$253,750,000

Pro Forma Ownership with Earn-out to BiomX and % total ownership (1) (Millions of shares)								
	Pro Forma Share Price, per share							
	\$10.00		\$16.50		\$22.75	\$29.00		
	<u>Shares</u>	<u>%</u>	Shares	<u>%</u>	Shares %	Shares	<u>%</u>	
BiomX Initial Shareholders	18.625		18.625		18.625	18.625		
Earn-out Shares, cumulative			2.000		4.000	6.000		
BiomX Initial Shareholders	18.625 7	3%	20.625	70%	22.625 70%	24.625	70%	
CHAC Public Shareholders	5.000 2	20%	6.061	21%	6.731 20%	7.112	20%	
CHAC Sponsors	1.750	7%	2.629	9%	3.184 10%	3.500	10%	
Pro Forma Shares Outstanding	25.375 10	00%	29.315	100%	32.540 100%	35.237	100%	



(1) Assuming no redemptions from the CHAC shareholders, giving effect to warrants to exercisable at \$11.50 per share, using treasury method to calculate fully diluted shares outstanding and 2 million shares of currently outstanding CHAC shares purchased at closing by current BiomX investors.

Executive Summary

Phage: A disruptive technology targeting significant market opportunities

- The customized (natural and engineered) phage platform is disruptive and has a long history of use in humans and animals ¹
- · Investment momentum in phage modality has been seen in recent months
- Acne is a significant initial opportunity, with a potential cosmetic commercialization path (Approximately a \$4 bn global cosmetic market²)
- · IBD, primary sclerosing cholangitis (an orphan indication), and oncology are significant opportunities

Investors: Robust science fuels support from leading life sciences and strategic investors

- · The platform and IP portfolio originate from leading scientists at MIT and the Weizmann Institute
- Strong support exists from institutional life sciences investors
- · BiomX insiders include big biopharma investors, which could support meaningful future partnerships

Catalysts: Several value inflection points (acne, IBD, PSC) over an expected 2-3-year period

- ~\$100mm of post-transaction cash offers certain clinical valuation inflection points
- Platform PoC acne data read out expected in 1Q20, with PoC expected to follow in IBD (2020) and PSC (2021)
- · Performance is likely on microbiome sector macro, given modest valuation and upcoming clinical data



- (1) Kortright et al. (2019), Cell Host & Microbe; Schmidt et al. (2019), Nature Biotechnology
- (2) Global Anti Acne Cosmetics Industry Market Research Report 2019, Wise Guy Reports

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Leadership Team

Management Team Proclara BiomX Jonathan Solomon CEO and Board Member POC Ltd ChondroSite Assaf Oron CBO Pfizer DURATA Allergan II TERUM BiomX Sailaja Puttagunta, MD CMO Ectel Ltd Trans Pharma Sigal Fattal¹ CFO **Scientific Founders** מכון ויצמן למדע שנסאא אאדווודס אמאס Prof. Rotem Sorek מכון ויצמן למדע מאוווא אאאצאי Prof. Eran Elinav MITBE Prof. Timothy K. Lu

(1) Following the closing of the transaction Sigal Fattal will transition from the CFO role.



Leadership Team

Board of Directors





Mission Statement



We develop precision medicines in the microbiome sector using customized phage therapies that target harmful bacteria in chronic diseases such as IBD and cancer.

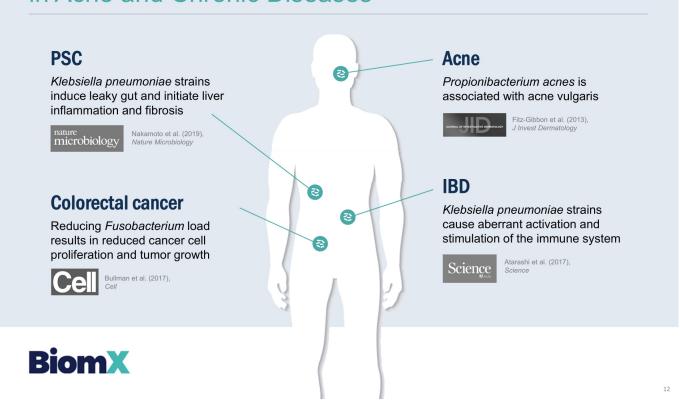


Unique Position

Technology	 Phage discovery platform Proprietary synthetic biology capabilities Cutting-edge data science In-house manufacturing
Pipeline	 Initial 4 programs: acne, IBD, PSC (a liver disease), colorectal cancer Phase I data in acne expected in Q1 2020 Phase I data in IBD expected in H2 2020 Phase I/II data in PSC expected in 2021
Exclusive access to novel targets	 Proprietary targets in IBD and PSC Target discovery and validation platform steered by cutting-edge research of scientific founders
Partnerships	Acne collaboration with leading global cosmetic company Biomarker discovery in IBD for key Janssen (J&J) IBD drug Janssen
Leading life science and strategic investors	OrbiMed Johnson-Yohnson Takeda Seventure 8VC



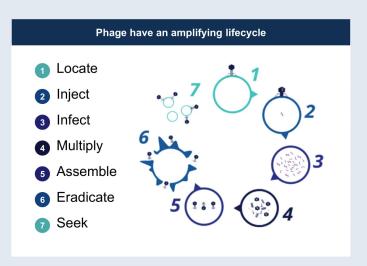
Growing Evidence of Harmful Bacteria Role in Acne and Chronic Diseases



Phage • Nature's Precision Targeting Vector

bacterial strains

Phage bind only to specific



Source: Kortright et al. (2019), Cell Host & Microbe



Phage Technology Platform

Phage Hunting

- · Sample sourcing
- · Automated sample processing
- · SynBio prophage extraction



Phage Engineering (SynBio)

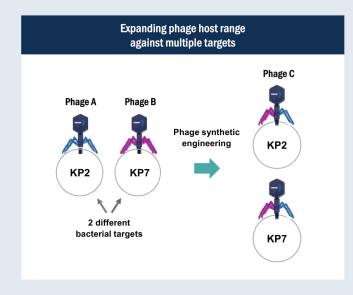
- · Host range expansion
- · Lysogenic to lytic
- · Payload incorporation

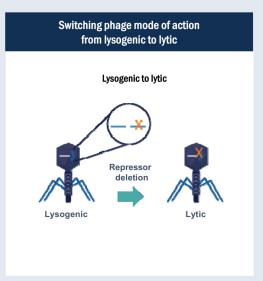
Cocktail Optimization

 Multi-dimensional optimization in vitro and in vivo: host range, biofilm, resistance, receptor analysis, toxic genes



Proprietary Synthetic Biology Capabilities





Source: Internal experiments



Pipeline

	Phage discovery	Preclinical	Phase I	Phase II	Partners
Product Candidates					
Acne • BX001¹				Phase I results expected 1Q20 Phase II results expected 2H20	Global cosmetics company
IBD • BX002				Pre-IND meeting expected 2H19 Phase I results expected 2H20	
PSC • BX003				Pre-IND meeting expected 2H20 Phase I/II results expected 2H21	
Colorectal cancer					
	Biomarker discovery	Validation	Developmo	ent	
Diagnostics					
IBD (responder/ non-responder)					janssen T

(1) BX001 is intended to be developed and commercialized as a cosmetic



Acne • Indication Offers Quick Phage Platform POC

- Strong rationale for phage approach: Biology underpins efficient eradication of *P. acnes*
- Will run clinical trial in collaboration with a leading multi-national cosmetic company
- Approximate \$4 billion global cosmetic market exists¹



(1) Global Anti Acne Cosmetics Industry Market Research Report 2019, Wise Guy Reports



Acne • BX001 Product Attributes

- Active in-vitro on 96% of tested P. acnes clinical strains
- Active in-vitro on antibiotic-resistant strains
- Self-amplifying: 50-100 phage per bacteria killed
- Penetrates biofilm (in contrast to antibiotic erythromycin)
- Highly specific: Does not affect other skin microbiome bacteria
- · Proprietary gel formulation

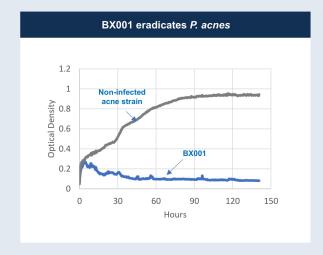
BX001
A topical gel containing natural phage against
P. acnes to modulate skin microbiome

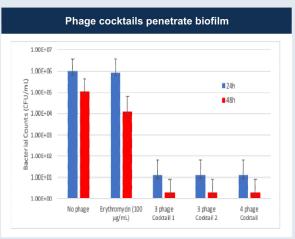


Sources: Internal experiments;



Acne • BX001 Pre-clinical Results



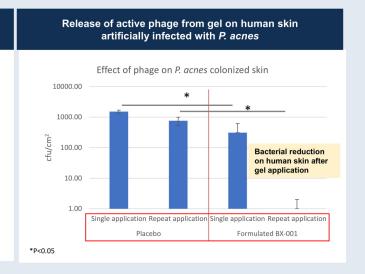


Source: Internal experiments



Acne • BX001 Pre-clinical Results

Release of active phage from gel on *P. acnes* bacterial lawn Cleared area after gel application Black: Where gel with BX001 was applied Red: Area of phage activity (eradicated *P. acnes*)



Source: Internal experiments

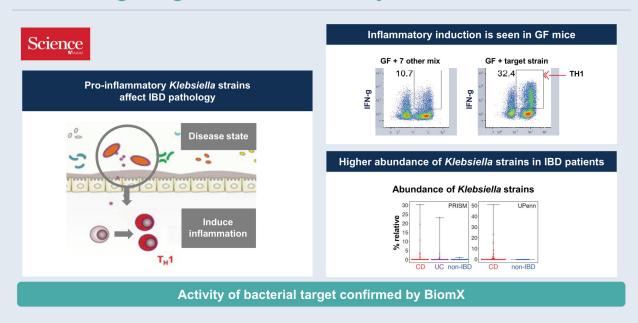


Acne • BX001 Planned Clinical Trials

Phase I Phase II 4-week study (placebo-controlled) 8-week study (placebo-controlled) Objectives Objectives i. Safety and efficacy i. Safety Exploratory endpoints Endpoints i. Reduction of P. acnes (efficacy) i. Reduction of *P. acnes* (efficacy) ii. Skin microbiome evaluation ii. Skin microbiome evaluation iii. Lesion numbers (efficacy, trend) iii. IGA and lesion numbers (efficacy) • 75 patients total over 3 cohorts · 100 patients total over 2 cohorts - 2 doses + 1 placebo (vehicle) - 1 dose + 1 placebo (vehicle) - 25 patients per cohort - 50 patients per cohort 2Q19 3Q19 4Q19 1Q20 2Q20 3Q20 4Q20



IBD • Targeting Pro-inflammatory Klebsiella Strains

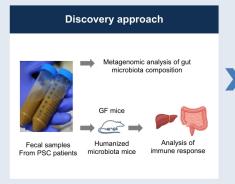


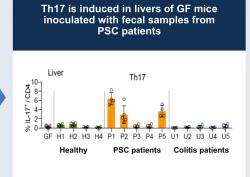
Source: Atarashi et al. (2017), Science

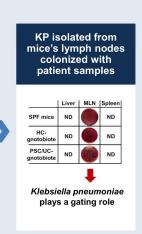


PSC • Klebsiella Identified as Novel Pathobiont

nature microbiology







Klebsiella pneumoniae (KP) is a specific gut pathobiont of PSC that is an intestinal barrier disrupter and is pro-inflammatory ("leaky gut")

Source: Nakamoto et al. (2019), Nature Microbiology



PSC • Bacterial Pathogens Contribute to Orphan Liver Disease





PSC (primary sclerosing cholangitis)

Characterized by stricturing of bile ducts that impedes the flow of bile to the intestines and gradually leading to cirrhosis of the liver and liver failure.

- ~30,000 patients in the US
- 10–15 years until liver transplant is required
- No existing therapy to avoid eventual liver transplant

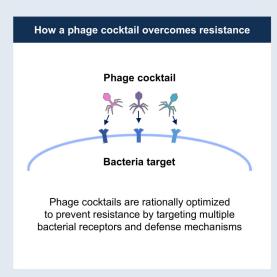
- Evidence that manipulation of the microbiome impacts the disease
- High abundance of bacteria found in bile fluid of patients
- A majority of PSC patients suffer from ulcerative colitis

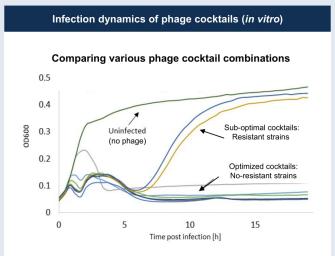
Hepatology. (2013) Dec;58(6):2045-55, UpToDate, MedScape

Source: NEJM 2016, PSC Review, LaRusso and Lazaridis



BX002 for IBD • Looking to Address Resistance by Using Phage Cocktails

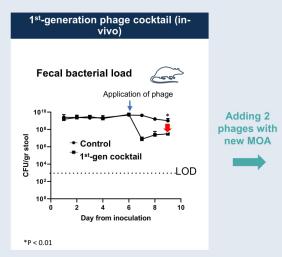


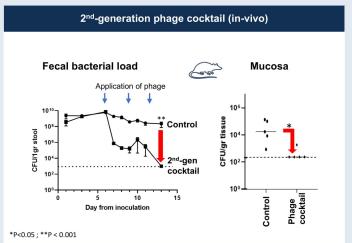


Source: Internal experiments



BX002 for IBD • Cocktail Composition Drives Activity



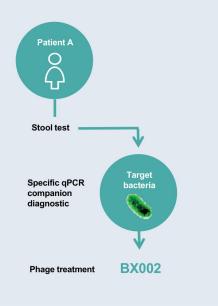


Source: Internal experiments



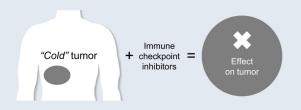
IBD • Planned Clinical Development - Phase 1

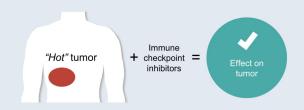
Study	Phase 1a/b
Study objectives	 Primary: Safety and tolerability of orally-administered BX002 Secondary: Reduction of target bacteria levels in stool Evaluation of microbial composition in stool Exploratory: Local inflammatory
Population	■ Target bacteria carriers – patients or healthy individuals
Cohorts	 30-45 patients across 3 cohorts 2 dose levels + placebo (vehicle) 10-15 patients per cohort
Treatment route, duration	Oral route4 weeks, daily administration





CRC • Most Colorectal Cancer (CRC) Patients Do Not Respond to Immunotherapy

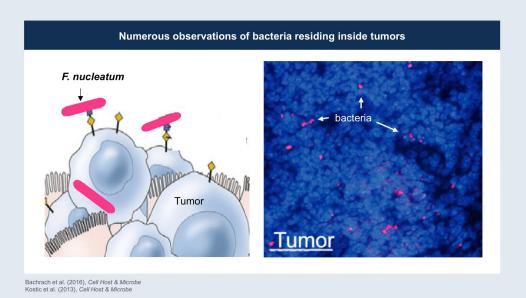




Sources: Vareki (2018), Journal for immunotherapy of Cancer; Galon et al. (2019), Nature Reviews/Drug Discovery

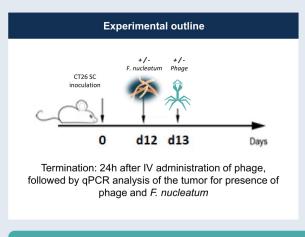


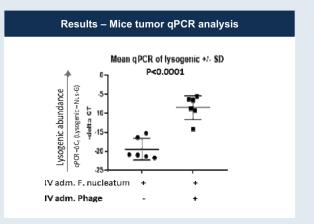
CRC • Bacteria Residing Inside Tumors Offer a Novel Targeted Intervention to "Uncloak" Tumors to "Hot"





CRC • IV Delivery of Phage to Intra-tumor Bacteria has been Demonstrated



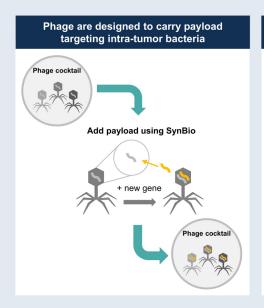


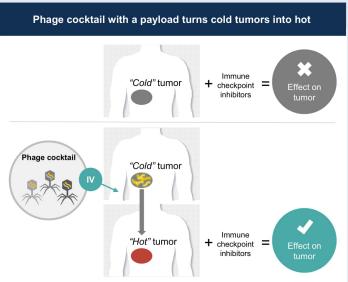
Detection of lysogenized intra-tumor *F. nucleatum* demonstrates phage delivered IV reached bacteria within tumor microenvironment and integrated stably into host bacteria genome

Source: Internal experiments



CRC • Engineered Phage Designed to Bring Immunestimulating Payload to Bacteria in Tumors







Key Catalysts

	1H19	2H19	1H20	2H20	1H21	2H21
Acne	Mfg.	Phase I starts	Phase I Results	Phase II results	Prep large- scale mfg.	Prep large-scale mfg.
IBD	Phage cocktail	Pre-IND meeting	Mfg.	Phase I results	Phase II starts	Interim results
PSC	Phage discovered	Phage cocktail	Pre-IND meeting	Mfg.	Phase I/II starts	Interim results





Transaction Highlights¹

- · Pro forma valuation of \$254 million
 - ~\$99 million of post-transaction cash on the combined company balance sheet to pursue clinical and commercial development
- BiomX securityholders to roll 100% of their vested equity holdings in the transaction
 - Holders exchange all BiomX vested securities for 16.625 million CHAC shares² at \$10.00 per share
 - Certain BiomX Shareholders are also purchasing an additional 2.000 million CHAC shares directly from current CHAC shareholders via purchase and sale agreements
 - Results in BiomX vested securityholders with approximately 73% ownership post-merger
- Earn-out provision provides certain shareholders of BiomX additional potential milestone-based equity consideration as follows³:
 - Additional 2.0 million shares if share price exceeds \$16.50 by fiscal year 2021
 - Additional 2.0 million shares if share price exceeds \$22.75 by fiscal year 2023
 - Additional 2.0 million shares if share price exceeds \$29.00 by fiscal year 2025



- (1) Assuming no redemptions from the CHAC shareholders.
- $(2) \ \ Includes \ vested \ securities \ convertible \ into \ CHAC \ shares.$
- (3) Based on VWAP during any 20 out of a 30-day period.

Risks Related to Projections and Pro Forma Presentation

The inclusion of the CHAC projections for BiomX in the Proxy Statement and the inclusion in this Investor Presentation of certain analyses referencing such projections should not be regarded as an indication that BiomX, CHAC or their respective advisors or other representatives considered or consider the projections to be necessarily predictive of actual future financial performance or events, and the projections should not be relied upon as such.

BiomX is a preclinical stage microbiome company expecting to start a first clinical trial by the end of 2019, and as such does not yet have any revenue-generating products, and thus does not make public its long-term financial forecasts driven by the potential of its phage products. In connection with BiomX's Board of Directors' evaluation of the Business Combination, BiomX's management did not prepare long-range, risk-adjusted revenue projections but did provide expense estimates for the years 2019 through 2022, which were based on numerous assumptions and qualifications believed by BiomX to be reasonable. CHAC conducted additional analyses, independently of BiomX, to assess the risk-adjusted revenue prospects of acne (assuming the product candidate will be marketed as a cosmetic), IBD, PSC, and colorectal cancer products by relying on assumptions, none of which were approved by BiomX, about the robustness of BiomX's technologies, the likelihood of the emergence of phage as a therapeutic platform, regulatory postures around phage technology, and individual product probabilities of success, launch timing, pricing, pricing growth, market growth, phage market penetration, BiomX product candidates market share, effects from competition and certain other factors affecting the commercial prospects of BiomX's product candidates.

Factors considered and assumptions made by CHAC are extremely uncertain and difficult to predict, with many being beyond the control of BiomX or its competitors. CHAC thought it appropriate to prepare forecasts representing three cases (A, B, and C) reflecting a range of risk-adjusted commercial outcomes on BiomX's portfolio of product candidates, none of which has received any regulatory or marketing approvals. The three cases represent three potential outcomes with revenue variations over the three cases (A, B, and C) driven only by differences in various assumptions for product launch probabilities. As such, there can be no certainty that the projections presented will be realized or that BiomX will ever receive regulatory approvals required in connection with any product candidates or achieve profitability.

Case A: Lower product launch probabilities (BX001 40%, BX002 15%, BX003 10%)
Case B: Moderate product launch probabilities (BX001 60%, BX002 25%, BX003 15%)
Case C: Higher product launch probabilities (BX001 100%, BX002 40%, BX003 25%)



Risks Related to Projections and Pro Forma Presentation (continued)

The projections below were not prepared for the purposes of public disclosure, nor for adherence to compliance with published guidelines of the SEC, nor for U.S. generally accepted accounting principles or other foreign or international accounting standards. In addition, the projections below were prepared by CHAC without the assistance, compilation, examination, or other review by independent accountants.

There may be differences between actual and projected results, and the differences may be material. The risk that these uncertainties and contingencies could cause the assumptions to fail to be reflective of actual results is further increased by the length of time over which these assumptions apply. The failure to achieve assumptions and projections in early periods could have a compounding effect on the projections shown for the later periods. Thus, any such failure of an assumption or projection to be reflective of actual results in an early period could have a greater effect on the projected results failing to be reflective of actual events in later periods.

BiomX is a preclinical stage company, without a regulatorily-approved product, and as discussed in the Proxy Statement, its business is subject to numerous risks. In the context of a preclinical stage company projections are inherently unreliable given the many variables, especially in later years, that may affect results

All projections are "forward-looking statements" within the meaning of the "safe harbor" provisions of the US Private Securities Litigation Reform Act of 1995. See "Forward-Looking Statements" in this Investor Presentation.



Merger Valuation versus DCF Analysis

Discounted Cash Flow Analysis, based on forecasts prepared by CHAC using a range of assumptions regarding clinical success and commercial acceptance. Information on this page should be read in conjunction with prior pages and with disclosures in the Proxy Statement regarding the risk and uncertainties related to pro forma forecasts.





