Zacks Small-Cap Research
Sponsored – Impartial - Comprehensive

August 6, 2020
David Bautz, PhD
312-265-9471
dbautz@zacks.com

scr.zacks.com
10 S. Riverside Plaza, Chicago, IL 60606

Antibe Therapeutics Inc.

(ATBPF - OTCQB)

ATBPF: Commercial Studies Show Blockbuster Potential for ATB-346…

Based on our probability adjusted DCF model that takes into account potential future revenues from otenaproxesul, ATBPF is valued at $2.25 per share. This model is highly dependent upon continued clinical success of otenaproxesul and will be adjusted accordingly based upon future clinical results and the company's execution.

Current Price (08/06/2020)	$0.34
Valuation	**$2.25**

OUTLOOK

Antibe Therapeutics Inc. (ATBPF) is developing safer medicines for pain and inflammation utilizing the company's proprietary hydrogen-sulfide releasing technology. Antibe recently announced positive topline results for the Phase 2b dose-ranging efficacy study of the company's lead development compound, ATB-346 (otenaproxesul) in patients with osteoarthritis. The company also recently announced the findings of market and commercial studies showing the blockbuster potential for otenaproxesul. Near-term milestones include filing an IND for otenaproxesul before the end of 2020, meeting with the FDA for an 'End-of-Phase 2' meeting before the end of 2020, and initiating a pivotal Phase 2/3 adaptive design trial late in the first quarter or early in the second quarter of 2021.

SUMMARY DATA

52-Week High	**$0.57**
52-Week Low	**$0.25**
One-Year Return (%)	**9.97**
Beta	**0.25**
Average Daily Volume (sh)	**261,039**
Shares Outstanding (mil)	**386**
Market Capitalization ($mil)	**$93**
Short Interest Ratio (days)	**1**
Institutional Ownership (%)	**0**
Insider Ownership (%)	**14**
Annual Cash Dividend	**$0.00**
Dividend Yield (%)	**0.00**
5-Yr. Historical Growth Rates	
Sales (%)	**N/A**
Earnings Per Share (%)	**N/A**
Dividend (%)	**N/A**
P/E using TTM EPS	**N/A**
P/E using #Lin Estimate	**N/A**
P/E using #Lin Estimate	**N/A**

Risk Level	**Above Avg.**
Type of Stock	**Small-Growth**
Industry	**Med-Biomed/Gene**

ZACKS ESTIMATES

Revenue
(In millions of $)

	Q1 (Jun)	Q2 (Sep)	Q3 (Dec)	Q4 (Mar)	Year (Mar)
2020	2.1 A	1.7 A	2.0 A	1.7 A	7.1 A
2021	1.5 E	1.5 E	1.9 E	1.9 E	6.8 E
2022					8.3 E
2023					8.3 E

Earnings per Share

	Q1 (Jun)	Q2 (Sep)	Q3 (Dec)	Q4 (Mar)	Year (Mar)
2020	-$0.01 A	-$0.01 A	-$0.02 A	-$0.02 A	-$0.05 A
2021	-$0.01 E	-$0.01 E	-$0.01 E	-$0.01 E	-$0.04 E
2022					-$0.03 E
2023					-$0.03 E

© Copyright 2020, Zacks Investment Research. All Rights Reserved.

WHAT'S NEW

Business Update

Advancing Toward Phase 3 Program for ATB-346 (Otenaproxesul)

Antibe Therapeutics, Inc. (ATBPF) is preparing to move its lead development compound, otenaproxesul (the nonproprietary name for ATB-346), into a Phase 3 program. Otenaproxesul is being developed as a solution to the dose-related gastrointestinal (GI) side effects associated with nonsteroidal anti-inflammatory drugs (NSAIDs). It uses naproxen as a base molecule with a hydrogen sulfide moiety covalently attached. Hydrogen sulfide (H_2S) is an important gasotransmitter, a gas that serves an important signaling molecule in the body.

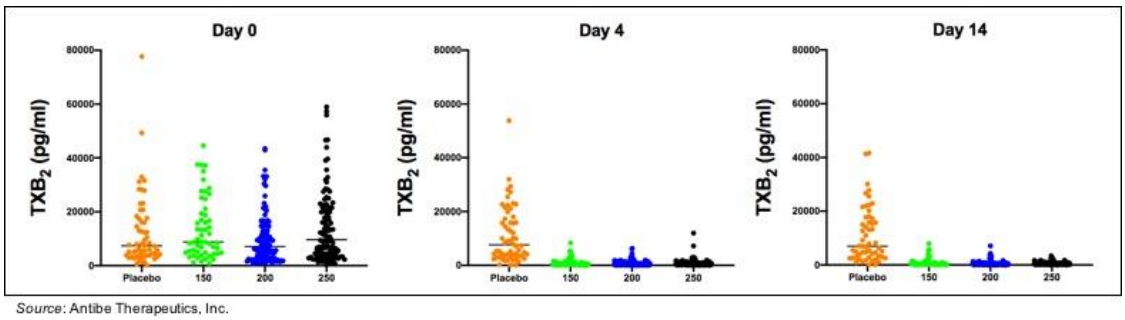
In 2018, Antibe announced positive topline results for the Phase 2 GI safety study of otenaproxesul. The double-blind, study enrolled 244 healthy volunteers, in which participants were administered either 250 mg otenaproxesul once daily or 500 mg naproxen twice daily. All subjects had an endoscopic examination of the upper GI tract prior to drug treatment and at the end of the 14-day study period.

The results showed that 42.1% (53/126) of naproxen-dosed subjects had GI ulceration compared to only 2.5% (3/118) of subjects treated with otenaproxesul, which was a statistically significant difference (P<0.001). Importantly, there were no safety signals and transient increases in liver enzymes were seen at levels comparable to what is seen with the most commonly prescribed NSAIDs.

In June 2020, Antibe announced positive topline results from the Phase 2b dose-ranging, efficacy trial of otenaproxesul. The randomized, double blind, placebo controlled trial enrolled 385 patients with osteoarthritis (OA) of the knee to once daily placebo or one of the three doses of otenaproxesul: 250 mg, 200 mg, and 150 mg. The primary endpoint of the trial was the change from baseline in the WOMAC pain subscale score following 14 days of treatment. Secondary efficacy endpoints included the WOMAC stiffness subscale score and the WOMAC difficulty performing daily activities (DPDA) subscale score.

Patients administered otenaproxesul showed a statistically significantly greater decrease in the change in the WOMAC pain subscale score compared to those administered placebo at both 250 mg (P=0.01) and 200 mg (P=0.007). The 150 mg cohort was not powered for statistical significance, however that dose showed robust efficacy in the primary endpoint and likely would have been statistically significant if it had been adequately powered like the other treatment arms. The full results are discussed in detail in our previous report. The adverse event profile was similar between placebo and all three doses of ATB-346, and consisted of events typically seen with the use of NSAIDs (e.g., dyspepsia, acid reflux, and dizziness).

In addition to the WOMAC results, the company also recently reported that otenaproxesul treatment resulted in a marked inhibition of cyclo-oxygenase enzymes (TXB_2 is a product of COX enzyme activity). The following figure shows that all three doses of otenaproxesul resulted in a profound inhibition by Day 4 and this continued through Day 14. In addition, there was little difference between the three doses, and likely indicates that the company will be able to decrease the dose before seeing any decrease in efficacy.



Source: Antibe Therapeutics, Inc.

Now that the efficacy and GI safety data for otenaproxesul is in hand, the company has begun planning the Phase 3 development program. We anticipate an IND being filed before the end of 2020 and an 'End-of-Phase 2' meeting occurring with the FDA (and the equivalent meeting with the EMA) on a similar timeframe. While we await word on the specific details of the Phase 3 program, we believe it will involve:

- Two 12-week efficacy trials that will include lower doses of otenaproxesul than were seen in the Phase 2b dose-ranging efficacy trial (tested against placebo) such that the lowest effective dose can be determined. The first of those trials is likely to be a Phase 2/3 adaptive design trial, whereby an interim analysis is conducted following 50% enrollment to ensure that an adequate number of patients are enrolled to achieve statistical significance. The second efficacy trial will likely commence following the interim analysis of the first trial.

- Two six-month GI-safety trials will be conducted so that the company can obtain a label for otenaproxesul that does not include the 'black box' warning regarding GI toxicity that is found on the labels for all NSAIDs.

- Safety data out to one year that will likely be collected from an open-label extension trial of the efficacy studies.

We anticipate partnering discussions with potential global and regional partners to accelerate now that the Phase 2 trials are completed. Antibe will be actively advancing toward initiation of a Phase 3 trial while simultaneously engaging with potential partners. While we continue to believe a partnership with a major, global pharmaceutical company is likely, we are unsure of the timing for such an agreement knowing that these arrangements can take a considerable amount of time to pull together.

Market Analysis Points to Blockbuster Potential for Otenaproxesul

Antibe recently reported on two commercial studies for otenaproxesul that examined its potential in the largest global markets (e.g., the U.S., the 'E.U. 5', and Japan). The studies were conducted by Shift Health (for the U.S. and E.U.) and LEK (for Japan). Following interviews with clinicians, payors, and pharmacy benefit managers, the analysis indicated that peak sales of up to CAD$5.3 billion are possible for otenaproxesul in OA. This forecast was based on a daily price of US$6 per day in the U.S., which is similar to branded NSAIDs (e.g., Celebrex®).

Financial Update

Raises CAD$28.75 Million in Bought Deal Public Offering

On June 30, 2020, Antibe announced it closed a bought deal public offering of 62.5 million units at a price of CAD$0.40 per unit. Each unit consisted of one share of common stock and one-third of one common share purchase warrant, with each full warrant exercisable at a price of CAD$0.60 prior to June 30, 2022. The underwriter's fully exercised their over-allotment option of 9.375 million units, thus the aggregate gross proceeds from the offering were CAD$28.75 million.

FY20 Financial Results

On July 24, 2020, Antibe announced financial results for fiscal year 2020 that ended Mar. 31, 2020. The company reported revenues of CAD$10.0 million in fiscal year 2020 compared to revenues of CAD$9.5 million for fiscal year 2019.

General and administrative, selling and marketing, research and development, stock-based compensation, and amortization and depreciation totaled CAD$21.5 million for fiscal year 2020 compared to CAD$15.7 million in fiscal year 2019. The increase was primarily related to the following:

- G&A expenses were CAD$5.7 million in fiscal year 2020 compared to CAD$4.9 million in fiscal year 2019. The increase was primarily due to higher salaries and professional and consulting fees.
- Selling and marketing costs were CAD$3.8 million in fiscal year 2020 compared to CAD$3.5 million in fiscal year 2019. The increase was due to higher salaries, commissions, travel, and entertainment costs.
- R&D expenses were CAD$8.1 million in fiscal year 2020 compared to CAD$3.9 million in fiscal year 2019. The increase was primarily due to higher development costs for ATB-346 partially offset by lower salary and professional costs.
- Stock-based compensation in fiscal year 2020 was CAD$3.4 million compared to CAD$3.0 million in fiscal year 2019 due to expensing of previously granted RSUs.
- Amortization and depreciation was CAD$0.6 million in fiscal year 2020 compared to CAD$0.4 million in fiscal year 2019.

As of March 31, 2020, Antibe reported cash and cash equivalents of approximately CAD$6.2 million. From April 1, 2020 through July 24, 2020, the company raised approximately CAD$1.2 million from exercised stock options and CAD$3.3 million from exercised warrants. When including the June 2020 financing we estimate that the company currently has approximately CAD$34 million.

We estimate that the company currently has approximately 385.3 million shares outstanding and, when considering stock options and warrants, a fully diluted share count of approximately 451 million shares. Antibe is preparing for the optimal time to uplist the company's shares to a senior exchange, most likely either the Nasdaq and/or the Toronto Stock Exchange. In addition, the company will be expanding its investor outreach in the coming months to include institutional investors in an effort to expand its shareholder base.

Conclusion

Antibe has a plan in place to get otenaproxesul into a global Phase 3 program, with the first Phase 3 clinical trial likely to start in the Spring of 2021, and subsequent trials initiating over the subsequent months. The market analyses are very encouraging and show that an effective NSAID without GI side effects would be a welcome addition to the treatment options for those suffering from OA. We are looking forward to continued updates as the company meets with regulatory authorities in the second half of 2020 in preparation for the Phase 3 program. Our model had accounted for additional financings, thus even after the recent offering our valuation remains at $2.25 per share.

PROJECTED FINANCIALS

Antibe Therapeutics Inc.
Income Statement

Antibe Therapeutics, Inc. Fiscal Year Ends Mar. 31	FY 2020 A	Q1 '21 E	Q2 '21 E	Q3 '21 E	Q4 '21 E	FY 2021 E	FY 2022 E	FY 2023 E
ATB-346 (royalty)	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0
ATB-352 (royalty)	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0
Citagenix	$7.5	$1.5	$1.5	$1.9	$1.9	$6.8	$8.3	$8.3
Licensing / Development	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0
Total Revenues	$7.5	$1.5	$1.5	$1.9	$1.9	$6.8	$8.3	$8.3
YOY Growth	*3.5%*	*-20.7%*	*-8.9%*	*-3.2%*	*4.9%*	*-4.2%*	*16.7%*	*0.0%*
Cost of Goods Sold	$4.6	$1.0	$1.0	$1.2	$1.2	$4.4	$5.2	$5.2
Product Gross Margin	*29%*	*26%*	*26%*	*27%*	*27%*	*27%*	*28%*	*28%*
SG&A	$7.1	$1.5	$1.7	$1.8	$2.0	$6.9	$7.0	$7.1
R&D	$6.1	$1.5	$1.5	$1.7	$1.8	$6.5	$6.8	$7.5
Stock-based compensation	$2.5	$0.8	$0.8	$0.6	$0.6	$2.3	$2.3	$2.3
Impairment of goodwill	$1.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0
Amortization and Depreciation	$0.4	$0.1	$0.1	$0.1	$0.1	$0.4	$0.4	$0.4
% Other	*4.3%*	*5.4%*	*5.6%*	*4.2%*	*2.0%*	*4.2%*	*3.4%*	*3.4%*
Operating Income	($14.2)	($3.4)	($3.5)	($3.5)	($3.7)	($13.6)	($13.3)	($14.2)
Operating Margin	-142%	-169%	-177%	-138%	-149%	-151%	-121%	-129%
Interest Income / Net	($0.3)	($0.1)	($0.1)	($0.0)	($0.2)	($0.6)	($0.6)	($0.6)
Pre-Tax Income	($14.5)	($3.5)	($3.6)	($3.5)	($3.9)	($14.2)	($13.9)	($14.8)
Taxes	($0.0)	$0.0	$0.0	$0.0	$0.0	$0.0	($0.0)	($0.0)
Tax Rate	*$0.0*	*$0.0*	*$0.0*	*$0.0*	*$0.0*	*$0.0*	*$0.0*	*$0.0*
Net Income	($14.5)	($3.5)	($3.6)	($3.5)	($3.9)	($14.2)	($13.9)	($14.8)
Reported EPS	($0.05)	($0.01)	($0.01)	($0.01)	($0.01)	($0.04)	($0.03)	($0.03)
Fully Diluted Shares	272.7	294.0	360.0	385.0	390.0	357.3	450.0	500.0

Source: David Bautz, PhD - Zacks Investment Research, Inc.

© Copyright 2020, Zacks Investment Research. All Rights Reserved.

HISTORICAL STOCK PRICE



Source: Zacks Small Cap Research

© Copyright 2020, Zacks Investment Research. All Rights Reserved.

DISCLOSURES

The following disclosures relate to relationships between Zacks Small-Cap Research ("Zacks SCR"), a division of Zacks Investment Research ("ZIR"), and the issuers covered by the Zacks SCR Analysts in the Small-Cap Universe.

ANALYST DISCLOSURES

I, David Bautz, PhD, hereby certify that the view expressed in this research report accurately reflect my personal views about the subject securities and issuers. I also certify that no part of my compensation was, is, or will be, directly or indirectly, related to the recommendations or views expressed in this research report. I believe the information used for the creation of this report has been obtained from sources I considered to be reliable, but I can neither guarantee nor represent the completeness or accuracy of the information herewith. Such information and the opinions expressed are subject to change without notice.

INVESTMENT BANKING AND FEES FOR SERVICES

Zacks SCR does not provide investment banking services nor has it received compensation for investment banking services from the issuers of the securities covered in this report or article.
Zacks SCR has received compensation from the issuer directly, from an investment manager, or from an investor relations consulting firm engaged by the issuer for providing non-investment banking services to this issuer and expects to receive additional compensation for such non-investment banking services provided to this issuer. The non-investment banking services provided to the issuer includes the preparation of this report, investor relations services, investment software, financial database analysis, organization of non-deal road shows, and attendance fees for conferences sponsored or co-sponsored by Zacks SCR. The fees for these services vary on a per-client basis and are subject to the number and types of services contracted. Fees typically range between ten thousand and fifty thousand dollars per annum. Details of fees paid by this issuer are available upon request.

POLICY DISCLOSURES

This report provides an objective valuation of the issuer today and expected valuations of the issuer at various future dates based on applying standard investment valuation methodologies to the revenue and EPS forecasts made by the SCR Analyst of the issuer's business.
SCR Analysts are restricted from holding or trading securities in the issuers that they cover. ZIR and Zacks SCR do not make a market in any security followed by SCR nor do they act as dealers in these securities. Each Zacks SCR Analyst has full discretion over the valuation of the issuer included in this report based on his or her own due diligence. SCR Analysts are paid based on the number of companies they cover. SCR Analyst compensation is not, was not, nor will be, directly or indirectly, related to the specific valuations or views expressed in any report or article.

ADDITIONAL INFORMATION

Additional information is available upon request. Zacks SCR reports and articles are based on data obtained from sources that it believes to be reliable, but are not guaranteed to be accurate nor do they purport to be complete. Because of individual financial or investment objectives and/or financial circumstances, this report or article should not be construed as advice designed to meet the particular investment needs of any investor. Investing involves risk. Any opinions expressed by Zacks SCR Analysts are subject to change without notice. Reports or articles or tweets are not to be construed as an offer or solicitation of an offer to buy or sell the securities herein mentioned.

CANADIAN COVERAGE

This research report is a product of Zacks SCR and prepared by a research analyst who is employed by or is a consultant to Zacks SCR. The research analyst preparing the research report is a resident outside of Canada and is not an associated person of any Canadian registered adviser and/or dealer and, therefore, the analyst is not subject to supervision by a Canadian registered adviser and/or dealer, and is not required to satisfy the regulatory licensing requirements of any Canadian provincial securities regulators, the Investment Industry Regulatory Organization of Canada and is not required to otherwise comply with Canadian rules or regulations.