

MabVax Therapeutics (MBVX - \$ 4.20)

MVT-5873 and MVT-2163 Updates with Conference Call Scheduled for This Afternoon

MBVX reported this morning an interim analysis from MVT-5873 in PDAC Phase I dose finding study and MVT-2163 in PDAC PET imaging analysis. The company is scheduled to host a conference call this afternoon at 4:15 pm for 3Q16 financial results and other updates (dial-in #: (877) 407-8293). MBVX also recently reported ending 3Q16 with a net loss of (\$4.4MM) or (\$0.86)/share. MBVX ended 3Q16 with cash of ~\$6.9MM.

- MVT-5873 in PDAC Phase I/II and MVT-2163 study updates.** MBVX announced that MVT-5873 monotherapy dose escalating study has completed three doses and have established the safety and tolerability enabling the trial to enter the combination portion of the study. A total of 16 patients have been dosed for the monotherapy so far and additional up-titration is ongoing with possibly two or more doses. Patients eligible for the monotherapy are those heavily pre-treated with multiple regimens and still with recurrences; while those eligible for the recommended Phase II dose (RP2D) for combination with nab-paclitaxel plus gemcitabine treatment are treatment naïve. The RP2D study outcomes will be compared to that of historical chemotherapy alone. The initial dose of MVT-5873 for the combo study will start from the second tested in the monotherapy. The combo portion of the study could potentially enroll up to 60 patients to determine the potential of MVT-5873 as part of a first line therapy. For MVT-2163 Phase I trial, the study has completed two doses with the second of a 17mg cold blocking. The objective of cold blocking is to sequester circulating antigens. A third cohort of 47 mg cold blocking is underway. After a conventional computerized tomography (CT) scans verification, the MVT-2163 results so far appeared to be very promising and we anticipate more updates later in 2017. MBVX also reiterated that they will start a MVT-1075 Phase I study in 1H17 as they expect to file an IND before YE16. Together, with the ongoing progress and the interim safety and specificity results we remain encouraged also given the company could advance the MVT-5873 program more rapidly than initially planned.
- Action.** We are reiterating our Buy rating and \$18 price target to reflect our view on highly encouraging developments of the broad HuMab-5B1-based platform. Valuation is based on peer comparable, probability adjusted DCF and sum-of-the-parts analyses.

Earnings Estimates: (per share)

(Dec)	1Q	2Q	3Q	4Q	FY	P/E
FY-16E	-0.15A	-0.92A	-0.86A	-0.86	-3.71	NM
FY-15A	-6.25	-0.29	-0.20	-1.03	-13.44	NM
FY-14A	NA	NA	NA	NA	-9.51	NM
FY-13A	NA	NA	NA	NA	NA	NM

Source: Laidlaw & Company estimates

Healthcare/Biotechnology

Ticker:	MBVX
Rating:	Buy
Price Target:	\$ 18.00

Trading Data:

Last Price (11/14/2016)	\$ 4.20
52-Week High (11/19/2015)	\$ 7.55
52-Week Low (2/16/2016)	\$ 3.03
Market Cap. (MM)	\$ 26
Shares Out. (MM)	6

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Anticipated milestones in 2016 and beyond

Product	Indication	Event	Timing	Importance
MVT-5873	Pancreatic adenocarcinoma (PDAC)	Report top-line results of the single agent Phase I/II study	1H17	***
		Report interim results of the combination Phase I/II study	1H17	****
		Report results of the combination Phase I/II study	2H17	****
MVT-1075	Pancreatic adenocarcinoma (PDAC)	Potentially start Phase I study	1H17	***
HuMab-5B1 ADC	Pancreatic adenocarcinoma (PDAC)	Potentially start Phase I study	2018	***
MVT-2163	Pancreatic adenocarcinoma (PDAC)	Potentially report Phase I study top-line results	Mid-2017	****
Sarcoma vaccine	Sarcoma	Potentially report Phase II study OS results	4Q16	***
Ovarian cancer vaccine	Ovarian cancer	Potentially report Phase II study OS results	1H17	***

**** / ***** Major catalyst event that could impact share price very significantly while *** event is more informative

Source: Laidlaw & Company estimates and company presentation.

Major risks

Clinical study failure could have a major impact on MBVX share value. Despite promising pre-clinical results of the company's lead products, HuMab-5B1 and HuMab-5B1-PET, it remains too early to predict the longer term safety and efficacy from the upcoming clinical studies. Given that clinical validation has not been established, it would be critical for these studies to demonstrate efficacy and a positive safety profile in order to increase the assets and shareholder value. Negative results of Phase I and future clinical studies could have a materially negative impact on the shareholder value; especially since the company has a very diverse-limited pipeline profile.

Yet-to-be-validated vaccinated patient derived monoclonal antibody (Mab) screening platform could remain uncertain. Although monoclonal antibodies have been established as a validated cancer treatment modality; currently there is no Mab derived from vaccinated patients that has been approved or is in a late clinical development stage to demonstrate efficacy. As such, clinical risks for monoclonal antibody based cancer therapy derived from successfully vaccinated patients are higher than similar products generated from other more proven development platforms.

Product may not be approved or reach anticipated sales. Although MBVX's current pipeline products have exhibited the potential to generate positive clinical outcomes from current and future trials; it remains too early to project whether any of these products would be approved by regulatory agencies. Even if the products were to enter the market, sales could be significantly below projections due to the specific product label under approval, physician consensus for prescribing the drug, changes of treatment paradigms, entrance of competitors, and the possible changes in pricing flexibility and payer reimbursement. A revenue outlook below expectations could also negatively affect MBVX shareholder value.

Additional financings could dilute shareholder value. Although the company currently has ~\$10MM (pro forma) cash after its recent financing, MBVX most likely would need more financial resources going forward if they want to expand and further develop their pipeline. Should the future operational expenses, especially from R&D, increase significantly, products not receive FDA approval, or product revenue not reach expectations; the company might need to issue new equity to raise additional cash. Under such a scenario, the share value of existing shareholders could be diluted.

Limited trading liquidity limits shareholder options. Given that MBVX shares only entered the public market recently; daily trading volume and name recognition are relatively modest. As such, shareholders wanting to increase or reduce their positions more substantially in a volatile stock market may face constraints.

Figure 1: Income Statement

MabVax Therapeutics – Income Statement												
(\$ '000)	2014	2015	1Q16	2Q16	3Q16	4Q16E	2016E	2017E	2018E	2019E	2020E	2021E
Revenue												
Grants	304	1,267	148	-	-	0	148	1	1	1	0	0
Product revenue	10	0	0	-	-	0	0	0	0	0	1,500	3,383
Total revenues	314	1,267	148	0	0	0	148	1	1	1	1,500	3,383
Gross revenue											1,500	3,383
Research and development	3,503	9,597	1,701	1,596	1,671	1,705	6,672	8,207	9,766	11,622	14,062	16,453
General and administrative	5,204	9,795	2,652	1,929	2,421	2,469	9,470	10,796	11,876	12,945	13,980	14,959
Marketing and sales												
Total operating costs and expenses	8,707	19,392	4,352	3,525	4,092	4,174	16,143	19,003	21,642	24,567	28,043	31,412
Operating Incomes (losses)	(8,393)	(18,125)	(4,204)	(3,525)	(4,092)	(4,174)	(15,995)	(19,002)	(21,641)	(24,566)	(26,543)	(28,029)
Interest and other income (expense)	(0)	(0)	(200)	(263)	(266)	(400)	(1,129)	(1,600)	(1,600)	(1,600)	(1,600)	(1,600)
Change in fair value of warrant liability	475	20	-	0	0	0	0	0	0	0	0	0
Tax												
Net Income (Loss)	(7,918)	(18,105)	(4,405)	(3,788)	(4,358)	(4,574)	(17,124)	(20,602)	(23,241)	(26,166)	(28,143)	(29,629)
Deemed dividend on Series A-1 preferred-stock	(2,215)	(9,018)	-	0	0	0	0	0	0	0	0	0
Deemed dividend on Series A-1 warrant		(179)	-	0	0	0	0	0	0	0	0	0
Deemed dividend on Series B preferred stock		(8,656)	-	0	0	0	0	0	0	0	0	0
Accretion of preferred stock dividends	(445)	(93)	-	0	0	0	0	0	0	0	0	0
Net loss allocable to common stockholders	(10,578)	(36,051)	(4,405)	(3,788)	(4,358)	(4,574)	(17,124)	(20,602)	(23,241)	(26,166)	(28,143)	(29,629)
Basic and diluted net loss per share	(\$9.51)	(\$13.44)	(\$0.15)	(\$0.92)	(\$0.86)	(\$0.86)	(\$3.71)	(\$2.81)	(\$2.25)	(\$1.96)	(\$1.83)	(\$1.71)
Shares used to calculate the basic and diluted net loss per share	1,112	2,682	3,947	4,129	5,041	5,341	4,615	7,341	10,341	13,341	15,341	17,341
Margin Analysis (% of Sales/Revenue)												
Costs of goods										15%	15%	15%
R&D	1115%	757%	1149%	NA	NA	NA	4507%	820693%	976624%	1162183%	937%	486%
SG&A	1657%	773%	1791%	NA	NA	NA	6397%	1079631%	1187594%	1294477%	932%	442%
Operating Income (loss)	-2671%	-1430%	-2840%	NA	NA	NA	-10803%	-1900224%	-2164118%	-2456560%	-1769%	-828%
Pretax	-2520%	-1429%	-2975%	NA	NA	NA	-11566%	-2060224%	-2324118%	-2616560%	-1876%	-876%
Tax Rate												
Net Income	-705%	-2845%	-2975%	NA	NA	NA	-11566%	-2060224%	-2324118%	-2616560%	-1876%	-876%
Financial Indicator Growth Analysis (YoY%)												
Total Revenue	NA	303%	-38%	-100%	-100%	-100%	-88%	-99%	0%	0%	149904%	126%
R&D	NA	174%	-1%	-31%	-47%	-30%	-30%	23%	19%	19%	21%	17%
SG&A	NA	88%	170%	-54%	6%	6%	-3%	14%	10%	9%	8%	7%
Operating Income (Losses)	NA	116%	70%	-45%	-23%	5%	-12%	19%	14%	14%	8%	6%
Pretax Income	NA	241%	-78%	-41%	-17%	15%	-53%	20%	13%	13%	8%	5%
Net Income	NA	241%	-78%	-41%	-17%	15%	-53%	20%	13%	13%	8%	5%
EPS	NA	41%	-98%	216%	332%	-17%	-72%	-24%	-20%	-13%	-6%	-7%
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Source: Bloomberg LP; Company reports; Laidlaw & Company estimates.

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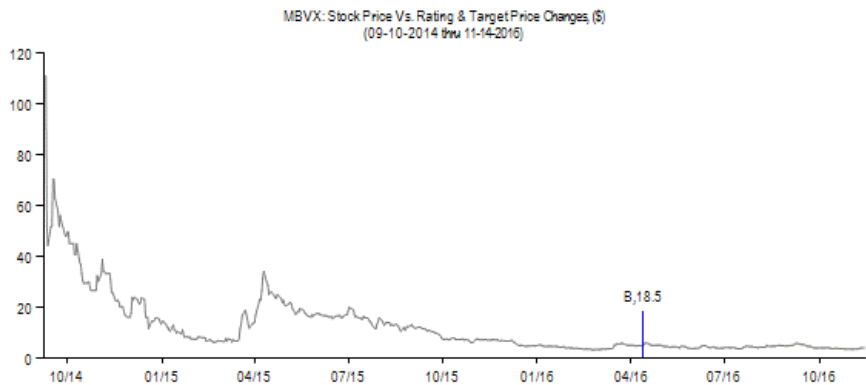
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Rating and Price Target Change History



3 Year Rating Change History

Date	Rating	Closing Price (\$)
04/13/2016	Buy (B)	5.18

3 Year Price Change History

Date	Target Price (\$)	Closing Price, (\$)
04/13/2016	18.50**	5.18

** Split Adjusted

Source: Laidlaw & Company

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			Investment Banking	Brokerage
Strong Buy (SB)	Expected to significantly outperform the sector over 12 months.	2.56%	2.56%	0.00%
Buy (B)	Expected to outperform the sector average over 12 months.	58.97%	28.21%	2.56%
Hold (H)	Expected returns to be in line with the sector average over 12 months.	5.13%	0.00%	0.00%
Sell (S)	Returns expected to significantly underperform the sector average over 12 months.	2.56%	0.00%	0.00%

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