September 17, 2015

ADMA Biologics (ADMA - \$ 9.49)

FDA Approved the Second Plasma Collection Center Ahead of Projection

This morning, ADMA reported that the FDA has approved ADMA's second plasma center, located at Marietta, GA to sell human source plasma within the U.S.

- **Details.** As expected, the FDA approved ADMA's second plasma center, located at Marietta, GA to sell human source plasma in the U.S. The timeline, however, is earlier than we projected (4Q15). The center could potentially support over 50 donor beds with 50 staff members at peak capacity. We anticipate the new center could start to generate revenue immediately (in 3Q15) and we believe the revenue generating capability of the 2nd center is overall similar to that of the first one.
- Implication. The approval of the 2nd plasma collection center is another milestone achieved because it not only provides ADMA additional modest revenue but also fulfills a strategic objective to supply a portion of raw material for RI-002. This helps insulate the company from certain supply and pricing fluctuations of external plasma supplies. ADMA filed a BLA on July 31st, 2015 for RI-002 as an IVIG to treat patients suffering from PIDD. We expect the FDA to continue the reviewing process, and the agency will likely decide to accept the application by end of September or early October. If so, we anticipate the potential PDUFA date could slate to August 2016 and RI-002 would likely be approved. The competition for RI-002 as an IVIG treating PIDD and organ transplantation patients is very limited, based on the unique attribute of consistently having high anti-RSV antibody titer (5.3x ↑ from the baseline, p=0.0001) and antibodies against other respiratory infectious pathogens. In addition, RI-002 recently has been granted patent of compositions and methods for the treatment of immunodeficiency (see our 2015-06-30 note). Together, we view RI-002 potentially is a very impressive and distinctive asset and makes ADMA, in our opinion, a potential acquisition target.
- **Action.** We are reiterating our Buy rating and \$20 target price to reflect the company's transition to a commercial-stage company. Our target price is supported by P/E, peer comparable and risk-adjusted cash flow sum-of-theparts analyses.

Earnings Estimates: (per share)

(Dec)	1Q	2Q	3Q	4Q	FY	P/E
FY-15E	-0.37A	-0.44A	-0.46	-0.47	-1.74	NM
FY-14A	-0.64	-0.43	-0.36	-0.38	-1.93	NM
FY-13A	-0.55	-0.83	-0.46	-0.55	-2.38	NM
FY-12A	-0.18	-0.20	-0.70	-0.68	-1.76	NM

Source: Laidlaw & Company estimates

Healthcare/Biotechnology

Ticker:	ADMA
Rating:	Buy
Price Target:	\$ 20.00

Trading Data:

Last Price (09/16/2015)	\$ 9.49
52-Week High (12/3/2014)	\$ 14.00
52-Week Low (5/8/2015)	\$ 7.51
Market Cap. (MM)	\$ 102
Shares Out. (MM)	11

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Anticipated milestones in 2015 and beyond

Program	Indication	Event	Timing	Importance
		Potentially build commercialization structure	2015	***
RI-002	RSV infection prevention in PIDD	Potential U.S. approval	2H16	****
		Potential U.S. product launch	2H16	****
VZIG (Varitect)	Vicella Zoster virus infection	Potential commence Phase II/III study	2015	***
BioCenters		FDA approval of 3rd BioCenter	2016	***

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

Source: Laidlaw & Company estimates and company presentation.

Major Risks

Risks of clinical study failure could have a major impact on ADMA share value. Despite a well-established path for IVIG approval, risks still exist as RI-002 might not be approved by the FDA if the pivotal study does not meet its endpoints. Given that the majority of upside for ADMA shares is currently based on the assumption that the product can be approved before its commercial potential can be realized, an unsuccessful approval application would have a significant negative impact on ADMA share value.

Commercial success of the RI-002 in PIDD and potentially in transplantation is less predictable. We believe that the potential product label for RI-002, if approved, would likely to indicate as a regular IVIG; and higher titer of anti-RSV antibodies could appear on the label if the pivotal study met the relevant secondary endpoint. As such, the company may not promote the product directly for the prevention or treatment of RSV infection. Instead it may be based on the understanding that receiving high titer RSV antibodies should reduce probability of RSV infection. With more limited sales and marketing tactics available, the sales ramp up could be slower than projected. There is also risk that more rapid sales expansion might only occur after the company conducts more clinical studies and demonstrates positive clinical outcomes.

Developments by competitors may render RI-002 or relevant technologies obsolete or un-competitive. Although the manufacturing processes of RI-002 are protected by proprietary technology, trade secrets and know-how, it is possible that other competitors develop similar processes to produce similar or even better anti-RSV IVIG. As such, the company might not enjoy the competitive edge and potentially damage RI-002's commercial outlook

Plasma collection center operations might not perform as expected. The company currently operates one and expects to expand into three plasma collection centers over the next 12 months. Although the plasma collection operation is a business with relatively sustainable positive cash flows, and ADMA management appears to have substantial experience; risks of mismanagement or other factors could result in sub-par business performance. Less successful performance in this area could negatively impact the expected cash flow and strategic objective of diversifying plasma sources for RI-002 production.

Limited product diversity could increase overall risk. Given the nascent stage of the corporate development, the majority of the product pipeline value mainly resides on RI-002. The second potential pipeline product, an anti-Vicella Zoster virus immunoglobin, is in very early development stage with market potential possibly much smaller. As such, we believe the company at the current stage has very limited diversification potential in its product pipeline.

Lack of cash could impede corporate development. Despite the company's recent successful IPO, which raised \$26.5MM of cash, ADMA could potentially

need more financial resources going forward if they want to expand and further develop its pipeline. Should the product not receive FDA approval or product revenue does not reach expectations, the company might have 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 ADMA shares only entered the public market recently; daily trading volume and name recognition are relatively low. With relatively illiquid trading volume, shareholders wanting to increase or reduce their positions in a volatile stock market may face constraints.

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(\$ '000)	2012	2013	2014					2015E	2016E	2017E	2018E	2019E	2020E
	2012	2013	2014	1Q15	2Q15	3Q15E	4Q15E	2015E	20100	2017	2010E	2019E	2020E
Revenue								· ·					
Product revenue	1,118	3,024	5,840	1,484	1,291	1,511	1,556	5,842	8,996	12,235	12,846	13,489	14,25
RI-002 revenue	-	-	-						17,318	52,409	90,303	143,560	185,04
License revenue	-	44	76	19	19	19	19	76	300	300	300	300	30
Total Revenue	1,118	3,068	5,916	1,503	1,310	1,529	1,575	5,917	26,614	64,944	103,449	157,349	199,60
Cost of product revenue	669	2,023	3,742	910	786	1,065	1,097	3,858	6,117	8,320	8,736	9,172	9,69
Cost of RI-002		·	, i			,		,	8,659	26,205	45,151	71,780	92,52
Gross revenue (RI-002)									8,659	26,205	45,151	71,780	92,52
Gross revenue (Biocenter)	449	1,000	2,076	575	505	446	459	1,984	2,879	3,915	4,111	4,316	4,5
Total gross revenue	449	1,000	2,076	575	505	446	459	1,984	11,538	30,120	49,262	76,097	97,0
Research and development	3,469	9,303	9,517	1,402	1,506	2,018	2,038	6,964	7,242	7,532	7,908	8,304	8,7
Plasma center operating expenses	1,747	2,418	3,851	1,048	1,097	1,108	1,130	4,383	4,558	4,877	5,072	5,224	5,3
General and administrative	3,142	4,365	4,824	1,346	1,437	1,969	2,540	7,293	7,585	7,888	8,156	8,434	8,7
Marketing and sales									8,400	8,904	9,438	9,948	10,4
Total Operating Expenses	8,358	16,087	18,192	3,796	4,040	5,095	5,708	18,640	27,785	29,201	30,575	31,910	33,2
perating Income (loss)	(7,909)	(15,042)	(16,019)	(3,202)	(3,517)	(4,631)	(5,231)	(16,580)	(1,171)	35,743	72,874	125,439	166,3
Interest income	21	(10,012)	14	5	10	(,, 55 .)	6	27	43	69	82	99	1
Interest income	(31)	(618)	(1,286)	(476)	(453)	(322)	(322)	(1,573)	(1,573)	(1,573)	(1,573)	(1,573)	(1,5
Change in fair value of stock warrants	(31)	43	(74)	(476)	(433)	(322)	(40)	(1,373)	(1,373)	(1,373)	(1,373)	(1,373)	(1,5 (1
Other income		82	(74)	-	(719)	(40)	(40)	(718)	(718)	(718)	(718)	(718)	(7
Total other expenses	(10)	(485)	(1,346)	(403)	(1,163)	(356)	(355)	(2,277)	(2,349)	(2,323)	(2,309)	(2,293)	(2,2
Income (loss) before tax expense	(7,919)	(15,527)	(17,340)	(3,606)	(4,679)	(4,987)	(5,586)	(18,857)	(3,520)	33,420	70,565	123,146	164,1
Income tax expense-State income tax benefit	618	(15,527)	(17,363)	(3,606)	(4,679)	(4,967)	(3,366)	(10,057)	(3,320)	12,365	26,109	45,564	60,7
let Incomes (Losses)	(7,301)	(15,527)	(17,917)	(3,606)	(4,679)	(4,987)	(5,586)	(18,857)	(3,520)	21,055	44,456	77,582	103,3
Net Earnings (Losses) Per Share—Basic	(\$1.76)	(\$2.38)	(\$1.93)	(\$0.37)	(\$0.44)	(\$0.46)	(\$0.52)	(\$1.78)	(\$0.27)	\$1.57	\$3.21	\$5.45	\$7.07
Net Earnings (Losses) Per Share—Basic Net Earnings (Losses) Per Share—Diluted	(\$1.76)	(\$2.38)	(\$1.93)	(\$0.37)	(\$0.44)	(\$0.46)	(\$0.52)	(\$1.78)	(\$0.27)	\$1.57 \$1.57	\$3.21	\$5.45 \$5.45	\$7.07
• ' '	4,146	6,531	9,292	9,855	10,706	10,756	10,806	10,531	13,031	13,431	13,831	14,231	14,631
Shares outstanding—basic Shares outstanding—diluted	4,146	6,531	9,292	9,855	10,706	10,756	10,806	10,531	13,031	13,431	13,831	14,231	14,631
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Margin Analysis (% of Revenue)													
Gross	40%	33%	36%	39%	30%	30%	30%	34%	32%	32%	32%	32%	32%
Cost of RI-002									50%	50%	50%	50%	50%
R&D	310%	303%	161%	93%	115%	132%	129%	118%	27%	12%	8%	5%	4%
Plasma operation	156%	80%	66%	71%	85%	73%	73%	75%	51%	40%	39%	39%	37%
G&A	281%	142%	82%	90%	110%	129%	161%	123%	28%	12%	8%	5%	4%
M&S	20170	270	0270	0070	,	.2070	10170	.2070	32%	14%	9%	6%	5%
Operating Income (loss)	-707%	-490%	-271%	-213%	-268%	-303%	-332%	-280%	-4%	55%	70%	80%	83%
Net Income	-653%	-506%	-303%	-240%	-357%	-326%	-355%	-319%	-13%	32%	43%	49%	52%
Inancial Indicator Growth Analysis (Y/Y)	470/	4700/ I	020/	407	400/	100/	60/	001	E 40/	200/	E0/	E0/	00/
Product (Biocenter) revenue	47%	170%	93%	-4% N. A	-13%	12%	6% N. A	0% N. A	54% N. A	36% 203%	5%	5% 50%	6%
RI-002 revenue (projected)	N.A.	N.A.	N.A.	N.A.	N.A.	N.A.	N.A.	N.A.	N.A.		72%	59%	29%
Total Revenue	47%	174%	93%	-4%	-13%	12%	6%	0%	350%	144%	59%	52%	27%
Research and development	436%	168%	2%	-68%	-16%	36%	6%	-27%	4%	4%	5%	5%	5%
Plasma center operating expenses	50%	38%	59%	31%	34%	9%	-7%	14%	4%	7%	4%	3%	2%
General and administrative	119%	39%	11%	19%	-7%	90%	128%	51%	4%	4%	3%	3%	3%
Marketing and sales] _,,,]	2021	00.	4.451	001	5001	4451	40.	0000	6%	6%	5%	5%
Operating incomes	71%	90%	6%	-44%	-2%	52%	41%	4%	-93%	-3152%	104%	72%	33%
Pretax Income	27%	96%	12%	-39%	18%	47%	36%	9%	-81%	-1049%	111%	75%	33%
Net Income	24%	113%	15%	-39%	18%	47%	57%	5%	-81%	-698%	111%	75%	33%
EPS - Basic	-89%	35%	-19%	-42%	3%	27%	35%	-8%	-85%	-680%	105%	70%	30%
EPS - Diluted	-89%	35%	-19%	-42%	3%	27%	35%	-8%	-85%	-680%	105%	70%	30%
Shares outstanding—basic	1074%	58%	42%	6%	15%	16%	16%	13%	24%	3%	3%	3%	3%
Shares outstanding—diluted	1074%	58%	42%	6%	15%	16%	16%	13%	24%	3%	3%	3%	3%

September 17, 2015

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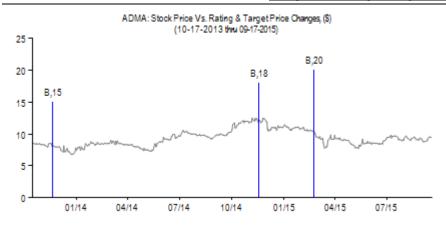
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Rating and Price Target Change History



3 Year Rating Change History							
Date	Rating	Closing Price (\$)					
11/21/2013	Buy (B.)	8.10					

3 Year Price Change History

Date	Target Price (\$)	Closing Price, (\$)
11/21/2013	15.00	8.10
11/18/2014	18.00	12.05
02/23/2015	20.00	10.40

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		With This Rating	Investment Banking	Brokerage		
Strong Buy (SB)	Expected to significantly outperform the sector over 12 months.	0.00%	0.00%	0.00%		
Buy (B)	Expected to outperform the sector average over 12 months.	74.19%	25.81%	6.45%		
Hold (H)	Hold (H) Expected returns to be in line with the sector average over 12 months.		0.00%	0.00%		
Sell (S)	Returns expected to significantly underperform the sector average over 12 months.	0.00%	0.00%	0.00%		

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