

# Chimerix

## Outperform (1)

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## Reports Q1:13; Dosing In SUPPRESS On Track To Begin During Q3

**Conclusion:** This morning Chimerix reported Q1:13 earnings and provided an update on its pipeline. We believe that Chimerix is undervalued based just on CMX001's potential as a CMV prophylactic in HSCT patients, with no contribution from other indications or other pipeline programs. We expect Chimerix's stock to outperform over the next 12 months as CMX001 progresses through development.

- **Q1:13 Financials.** Chimerix reported \$1.8MM in BARDA-associated revenue, a net loss of \$9.1MM and \$25.5MM in accretion of redeemable convertible preferred, equating to a net loss of \$22.58/share. Following Chimerix's April IPO the company had \$134.1MM in cash, \$14.3MM in debt, and 25.7MM shares outstanding.
- **Dosing In SUPPRESS To Begin During Q3.** The Ph. III SUPPRESS trial will test CMX001 in CMV prophylaxis. It will enroll 450 CMV seropositive adult patients and randomize them 2:1 to 100mg BIW CMX001 or placebo for 14 weeks. The primary endpoint of the study is failure to prevent CMV reactivation through week 24. The trial is powered to detect a 50% decrease in CMV reactivation in CMX001 vs. placebo. A single measure of CMV in the blood greater than or equal to 1,000 copies/mL (by quantitative PCR) will be considered a failure of CMV prevention. In addition, subjects at risk for rapid progression to CMV disease (recipients of umbilical cord blood stem cells, for example) will have a lower threshold of 150 copies/mL (the lower limit of quantification of the assay) for the initiation of pre-emptive therapy. Data from the SUPPRESS trial are expected in 2015.
- **CMRX To Define Pediatric Development Plan During H2.** Chimerix will unblind the results from the Phase II of CMX001 as preemptive therapy for adenoviral disease in HSCT during H2, and will define a pediatric registration plan after.

<b>CMRX (05/10)</b>	<b>\$20.05</b>	<b>Revenue \$MM</b>							
<b>Mkt cap</b>	<b>515.3MM</b>	<b>FY</b>	<b>2012</b>	<b>2013E</b>	<b>2014E</b>	<b>2015E</b>			
Dil shares out	25.7MM	<b>Dec</b>	<b>Actual</b>	<b>Prior</b>	<b>Current</b>	<b>Prior</b>	<b>Current</b>	<b>Prior</b>	<b>Current</b>
Avg daily vol	102.5K	Q1	0.0	—	1.8A	—	—	—	—
52-wk range	\$15.1-22.3	Q2	0.0	—	0.0	—	—	—	—
Dividend	Nil	Q3	0.0	—	0.0	—	—	—	—
Dividend yield	Nil	Q4	0.0	—	0.0	—	—	—	—
BV/sh	\$4.24	Year	<b>33.7</b>	<b>0.0</b>	<b>1.8</b>	—	<b>5.0</b>	—	<b>15.5</b>
Net cash/sh	\$4.26	EV/S	—	—	230.6x	—	83.0x	—	26.8x
Debt/cap	2.8%								
ROIC (LTM)	NA								
5-yr fwd EPS growth (Norm)	NA								
<b>S&amp;P 500</b>	<b>1630.8</b>	<b>EPS \$</b>	<b>2012</b>	<b>2013E</b>	<b>2014E</b>	<b>2015E</b>			
		<b>FY</b>	<b>Actual</b>	<b>Prior</b>	<b>Current</b>	<b>Prior</b>	<b>Current</b>	<b>Prior</b>	<b>Current</b>
		<b>Dec</b>							
		Q1	0.00	—	(22.58)A	—	—	—	—
		Q2	0.00	(0.42)	(0.40)	—	—	—	—
		Q3	0.00	(0.43)	(0.42)	—	—	—	—
		Q4	0.00	(0.45)	(0.44)	—	—	—	—
		Year	<b>(1.62)</b>	<b>(2.10)</b>	<b>(2.00)</b>	—	<b>(1.55)</b>	—	<b>(1.20)</b>
		P/E	—	—	—	—	—	—	—

## Investment Thesis

Chimerix is a biopharmaceutical company focused on the discovery and development of novel antivirals. Chimerix has a propriety lipid technology that has been shown to improve the potency of antivirals, and has produced two clinical stage candidates. Lead candidate CMX001 is a phospholipid derivative of GILD's cidofovir that can potentially kill a wide range of dsDNA viruses. It has successfully completed a Phase II trial for the prophylaxis against CMV reactivation in hematopoietic stem cell transplant (HSCT) patients. A Phase III trial is expected to begin in Q3:13, supporting an U.S. launch by 2016. Our consultants think CMX001 is safe, well tolerated, and potent, and is consequently likely to succeed in its Phase III SUPPRESS trial. Moreover, they think there is a need for a prophylactic to prevent infection with CMV and other dsDNA viral infections in transplant patients, and therefore expect CMX001 to be widely adopted once available. We project that CMX001 will achieve worldwide sales of \$330MM in HSCT alone by 2019, with Chimerix achieving profitability in 2017. CMX001 is also in development for the prevention of viral infection in solid organ transplant patients, and as a bioterrorism measure to prevent smallpox. Behind CMX001 is CMX157, a phospholipid derivative of GILD's tenofovir that partner Merck is developing for the treatment of HIV. We believe that Chimerix is undervalued based just on CMX001's potential as a CMV prophylactic in HSCT patients, with no contribution from other indications or other pipeline programs. We expect Chimerix's stock to outperform over the next 12 months as CMX001 progresses through development.

### Upcoming Chimerix Milestones

Event	Timing
Initiate Phase III SUPPRESS trial of CMX001 as prophylactic against CMV in adult HSCT	Q3:13
Data from Phase II Study 202 of CMX001 as preemptive therapy for adenoviral disease in HSCT	H2:13
Define pediatric development plan for CMX001	H2:13
Data from Study 350 of CMX001 in transplant patients with severe, life threatening dsDNA infections	2013
Negotiation with BARDA over continued funding of CMX001's smallpox program	2013

Source: Cowen and Company

## Chimerix Quarterly P&amp;L (\$MM)

	2012A	Q1:13A	Q2:13E	Q3:13E	Q4:13E	2013E
CMX-001		-				-
CMX-157 Royalty		-				-
Collaboration and Licensing Revenue	17.4	-	-	-	-	-
Contract And Grant Revenue	16.3	1.8				1.8
Total Revenue	33.7	1.8	-	-	-	1.8
COGS		-	-	-	-	-
<i>Gross Margin</i>						
R&D	27.8	6.5	7.2	7.8	8.2	29.7
SG&A	8.7	1.8	2.8	3.0	3.1	10.7
Other						
Operating Expenses	36.5	8.3	10.0	10.8	11.3	40.4
Operating Income / (Loss)	(2.8)	(6.5)	(10.0)	(10.8)	(11.3)	(38.6)
Interest Income, net	(0.8)	(0.4)	(0.2)	(0.2)	(0.2)	(1.0)
Other Income	(0.8)	(2.2)				
Pretax net income	(4.4)	(9.1)	(10.2)	(11.0)	(11.5)	(39.6)
Accretion of redeemable convertible preferred stock	(4.4)	(25.5)				
Taxes		-	-	-	-	-
<i>Tax Rate</i>		0%	0%	0%	0%	0%
GAAP Net Income	(8.8)	(34.6)	(10.2)	(11.0)	(11.5)	(39.6)
<b>GAAP EPS</b>	<b>\$(1.62)</b>	<b>\$(22.58)</b>	<b>\$(0.40)</b>	<b>\$(0.42)</b>	<b>\$(0.44)</b>	<b>\$(2.00)</b>
Diluted Shares Outstanding (MM)	5.4	1.5	25.7	25.9	26.0	19.8

Source: Cowen and Company

## Chimerix Annual P&amp;L (\$MM)

	2012A	2013E	2014E	2015E	2016E	2017E	2018E	2019E
CMX-001	-	-	-	-	45.0	135.0	240.0	330.0
CMX-157 Royalty	-	-	-	-	-	-	-	-
Collaboration and Licensing Revenue	17.4	-	5.0	15.5	20.0	20.0	20.0	20.0
Contract And Grant Revenue	16.3	1.8	-	-	-	-	-	-
Total Revenue	33.7	1.8	5.0	15.5	65.0	155.0	260.0	350.0
COGS	-	-	0.5	1.6	3.6	10.8	19.2	26.4
<i>Gross Margin</i>		0%	0%	0%	0%	0%	0%	0%
R&D	27.8	29.7	34.5	38.0	45.0	55.0	65.0	75.0
SG&A	8.7	10.7	12.0	13.0	28.0	34.0	40.0	45.0
Other	-	-	-	-	-	-	-	-
Operating Expenses	36.5	40.4	47.0	52.6	76.6	99.8	124.2	146.4
Operating Income / (Loss)	(2.8)	(38.6)	(42.0)	(37.1)	(11.6)	55.2	135.8	203.6
Interest Income, net	(0.8)	(1.0)	0.5	1.0	1.0	2.0	6.0	6.0
Other Income								
Pretax net income	(4.4)	(39.6)	(41.5)	(36.1)	(10.6)	57.2	141.8	209.6
Accretion of redeemable convertible preferred stock								
Taxes	-	-	-	-	-	-	-	-
<i>Tax Rate</i>	-	0%	0%	0%	0%	0%	0%	0%
GAAP Net Income	(8.8)	(39.6)	(41.5)	(36.1)	(10.6)	57.2	141.8	209.6
<b>GAAP EPS</b>	<b>(1.62)</b>	<b>(2.00)</b>	<b>(1.55)</b>	<b>(1.20)</b>	<b>(0.35)</b>	<b>1.65</b>	<b>3.95</b>	<b>5.65</b>
Diluted Shares Outstanding (MM)	5.4	19.8	26.8	30.0	30.5	34.8	35.9	37.1

Source: Cowen and Company



## Addendum

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Ticker	Company Name
CMRX	Chimerix

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(a) Assumptions: Time horizon is 12 months; S&P 500 is flat over forecast period.

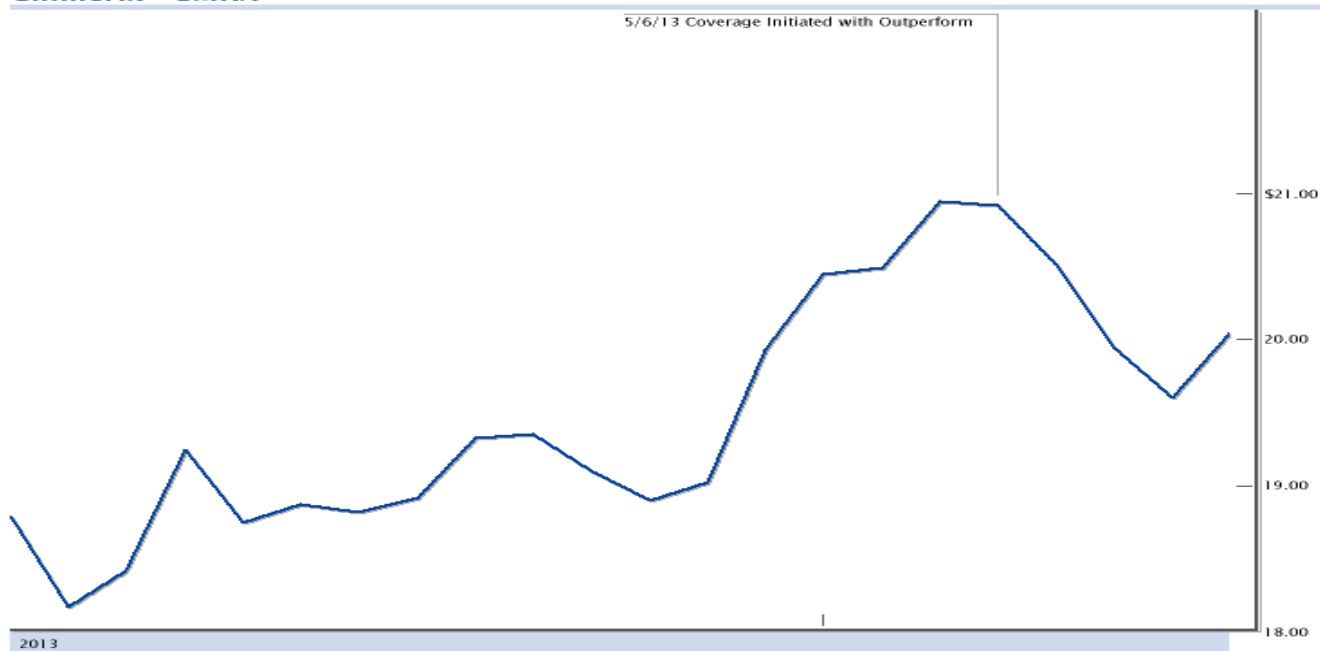
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**Chimerix - CMRX**



Pricing data provided by Reuters America. Chart as of 5/10/13 in USD.