

Celladon Corp.

Mydicar on Track

What's New? CLDN just announced 4Q13 and full year earnings and provided a review of its pipeline.

CUPID-2 on track to read-out in April 2015: Mydicar is currently being evaluated in a phase 2b trial (CUPID-2) for systolic heart failure. The trial completed enrollment in February and results are on track to be released in April 2015. If results show significant benefits with Mydicar, management believes it could be accepted as a pivotal trial, and an FDA approval could be granted in 2018. The EMA has also indicated to Celladon that it would accept CUPID-2 as a pivotal trial if the primary endpoint is met. We expect the FDA approval threshold for Mydicar to be high, so we have assigned a low probability of success (20%). However, we do believe Mydicar has a lower level of risk than other gene therapies, given the EMA and FDA's familiarity with and recognition of AAV1 vectors as a potentially safe delivery mechanism.

Significant room for upside: CLDN estimates the initial US target market for Mydicar at ~350,000 patients. We are using more conservative assumptions and estimate ~100,000 patients are initially eligible for treatment. We also assume a low penetration level in our model. We are estimating peak US sales of ~\$390mn and EU sales of ~\$240mn (royalties of~\$36mn). Nevertheless, we see significant upside potential to our estimates if CUPID-2 results demonstrate a significant improvement in heart failure symptoms and reduction of clinical events and hospitalizations.

Diabetes will be the first indication from the small molecules program: In February, CLDN entered into an options agreement with Servier to develop and commercialize SERCA2b modulators for type 2 diabetes and other metabolic diseases. The diabetes indication will be the first from the small molecule program and Servier will have the option to license the product for ex-US regions. Both companies will support the discovery phase but Servier will be responsible for further development efforts if the option is exercised.

CLDN: Quarterly and Annual EPS (USD)

	2013	2014			2015			Change y/y	
FY Dec	Actual	Old	New	Cons	Old	New	Cons	2014	2015
Q1	N/A	-0.36E	-0.37E	N/A	N/A	-0.40E	N/A	N/A	-8%
Q2	N/A	-0.36E	-0.40E	N/A	N/A	-0.35E	N/A	N/A	13%
Q3	N/A	-0.36E	-0.41E	N/A	N/A	-0.25E	N/A	N/A	39%
Q4	N/A	-0.36E	-0.44E	N/A	N/A	-0.27E	N/A	N/A	39%
Year	-26.13A	-1.43E	-1.62E	N/A	N/A	-1.27E	N/A	94%	22%
P/E	N/A		N/A			N/A			

Source: Barclays Research.

Consensus numbers are from Thomson Reuters

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Equity Research

OVERWEIGHT

Exchange-Nasdaq

Healthcare | U.S. Biotechnology 31 March 2014

J. G. C.	Unchanged
Industry View	NEUTRAL
	Unchanged
Price Target	USD 15.00
	Unchanged
Price (28-Mar-2014)	USD 11.26
Potential Upside/Downside	+33%
Tickers	CLDN
Market Cap (USD mn)	208
Shares Outstanding (mn)	18.50
Free Float (%)	68.18
52 Wk Avg Daily Volume (mn)	0.2
Dividend Yield (%)	N/A
Return on Equity TTM (%)	N/A
Current BVPS (USD)	-2.59
Source: Thomson Reuters	

Stock Rating



13-Feb-2014 28-Feb-2014 15-Mar-2014

Link to Barclays Live for interactive charting

U.S. Biotechnology

Price Performance

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U.S. Biotechnology Industry View: NEUTRAL							
Celladon Corp. (CLDN) Stock Rating: OVERWEIGHT							
Income statement (\$k)	2013A	2014E	2015E	2016E	CAGR	Price (28-Mar-2014) US	D 11.26
Revenue	0	0	0	N/A	N/A		D 15.00
EBITDA (adj)	-19,897	-28,544	-24,766	-29,621	N/A	Why Overweight? We view Celladon as having	
EBIT (adj)	-19,964	-28,698	-26,828	N/A	N/A	favorable risk/reward profile for investors. The	
Pre-tax income (adj) -23,		-28,777	-26,729	N/A	N/A	company is developing a gene therapy, Mydic	
Net income (adj)	-23,100	-28,777	-26,729	N/A	N/A	treat chronic heart failure. While there are	,
EPS (adj) (\$) -26.		-1.62	-1.27	N/A	N/A	considerable risks associated gene therapy, w	e also
Diluted shares (k)	884.2	17,808.0	21,675.6	N/A	N/A	view the market opportunity as significant if the	
DPS	N/A	N/A	N/A	N/A	N/A	company is successfully develop the product	
DF3	IN/ A	IN/ A	IN/ A	IN/ A	IN/ A		
Margin and return data					Average		D 42.00
EBITDA (adj) margin (%)	N/A	N/A	N/A	N/A	N/A	We see significant upside potential for Cellado phase 2b results are positive, we see valuation	
EBIT (adj) margin (%)	N/A	N/A	N/A	N/A	N/A	increasing to \$42/share.	1
Pre-tax (adj) margin (%)	N/A	N/A	N/A	N/A	N/A	increasing to \$427 share.	
Net (adj) margin (%)	N/A	N/A	N/A	N/A	N/A	Downside case U	JSD 2.00
ROIC (%)	-148.1	-82.3	-33.1	-61.5	-81.2	We see downside risks at approximately \$2/sl	
ROA (%)	-94.4	-65.0	-30.0	-52.1	-60.4	Mydicar gene therapy does not receive approx	
ROE (%)	-158.7	-84.9	-33.5	-62.7	-84.9	assumes some value attributed to the early sta	
Balance sheet and cash flow (\$k)				CAGR	small molecule platform and some cash.	
Tangible fixed assets	308	5,154	11,092	16,319	275.6%	Upside/Downside scenarios	
Intangible fixed assets	N/A	N/A	N/A	N/A	N/A		
Cash and equivalents	18,370	36,136	75,588	42,874	32.6%	Price History Price Target Prior 12 months Next 12 months	
Total assets	21,154	44,160	89,513	62,137	43.2%	High Upside	
Short and long-term debt	1,044	1,044	1,044	1,044	0.0%	42.00	
Other long-term liabilities	N/A	N/A	N/A	N/A	N/A	42.00	
Total liabilities	6,597	10,246	9,723	11,281	19.6%		
Net debt/(funds)	-6,859	-24,625	-64,077	-31,363	N/A		
Shareholders' equity	14,557	33,914	79,791	50,856	51.7%	17.16	
Change in working capital	-19,169	14,511	39,938	-34,162	N/A	Current	get
Cash flow from operations	-16,196	-23,841	-23,474	-25,833	N/A	11.26	00
Capital expenditure	-87	-5,000	-8,000	-8,000	N/A	7.45	
Free cash flow	1,664	-23,841	-23,474	-25,833	N/A	2.00	
Tree easi now	1,001	25,011	23,171	23,033	14//(
Valuation and leverage metric	s				Average	Low Downside	
P/E (adj) (x)	N/A	N/A	N/A	N/A	N/A		
EV/EBITDA (adj) (x)	-0.2	0.5	2.2	0.7	0.8	POINT® Quantitative Equity Scores	
Equity FCF yield (%)	N/A	N/A	N/A	N/A	N/A	Value	
EV/sales (x)	N/A	N/A	N/A	N/A	N/A		
P/BV (x)	N/A	N/A	N/A	N/A	N/A		
Dividend yield (%)	N/A	N/A	N/A	N/A	N/A	Quality	
Total debt/capital (%)	-89.1	-265.1	-407.8	-160.9	-230.7	N/A	
Selected operating metrics					Average	Sentiment	
SG&A/sales (%)	N/A	N/A	N/A	N/A	N/A	N/A	
R&D/sales (%)	N/A	N/A	N/A	N/A	N/A	14/74	
R&D growth (%)	27.1	40.0	-10.0	N/A	19.0		
SG&A growth (%)	15.4	64.6	10.0	N/A	30.0	Low	High
	.5.1	00	10.0	1771	30.3	Source: POINT®. The scores are valid as of the date report and are independent of the fundamental analyviews. To view the latest scores, please go to the equicompany page on Barclays Live.	ysts'

Source: Company data, Barclays Research Note: FY End Dec

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Primary Stocks (Ticker, Date, Price)

Celladon Corp. (CLDN, 28-Mar-2014, USD 11.26), Overweight/Neutral, A/C/D/J/K/L/M

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Arrowhead Research Corp. (ARWR)	Biogen Idec (BIIB)	BioMarin Pharmaceutical (BMRN)
Celgene Corp. (CELG)	Celladon Corp. (CLDN)	Dendreon Corp. (DNDN)
Fibrocell Science Inc. (FCSC)	Gilead Sciences (GILD)	GlycoMimetics Inc. (GLYC)
Halozyme Therapeutics Inc. (HALO)	Idenix Pharmaceuticals (IDIX)	Incyte Corp. (INCY)
Intrexon Corp. (XON)	Medivation Inc. (MDVN)	Regeneron Pharmaceuticals (REGN)
Tetraphase (TTPH)	Trevena Inc. (TRVN)	Vertex Pharmaceuticals (VRTX)

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Celladon Corp. (CLDN)

USD 11.26 (28-Mar-2014)

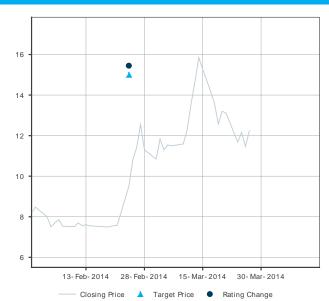
Stock Rating **OVERWEIGHT**

Industry View

NEUTRAL

15.00

Rating and Price Target Chart - USD (as of 28-Mar-2014)



Currency=USD Date **Closing Price** Rating **Adjusted Price Target** 24-Feb-2014 9.50

Overweight

Source: Thomson Reuters, Barclays Research

Historical stock prices and price targets may have been adjusted for stock splits and dividends.

Source: IDC, Barclays Research

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Valuation Methodology: Our price target is based on a probability-adjusted NPV analysis. We have only included the systolic heart failure indication in our model which contributes ~\$12/share. Including cash of ~\$3/share, we arrive at our price target of \$15.

Risks which May Impede the Achievement of the Barclays Research Price Target: Downside risks include failure of CUPID-2 trial, need for phase 3 trial, and inability to receive FDA approval for Mydicar.

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