

# **Conatus Pharmaceuticals Inc.** (CNAT)

Conatus to Initiate Two New Studies in 1H14

MARKET DATA	
Price	\$7.25
52-Week Range:	\$5.83 - \$11.24
Shares Out. (M):	16.0
Market Cap (\$M):	\$116.0
Average Daily Vol. (000):	105.0
Cash (M):	\$60
LT Debt (M):	\$1
Source: Thomson Reuters and JMP Securities LLC	

FY DEC		2012A	2013E	2014E
Revenue (\$M)	1Q		\$0.0A	
, , , , , , , , , , , , , , , , , , ,	2Q		\$0.0A	
	3Q		\$0.0A	
	4Q		\$0.0	
	FY	\$0.0	\$0.0	\$0.0
EPS	1Q		(\$0.03)A	
	2Q		\$0.16A	
	3Q		(\$0.28)A	
	4Q		(\$0.38)	
	FY	(\$1.04)	(\$0.50)	(\$1.62)
Source: Company reports and JMP Securities LLC				



MARKET OUTPERFORM | Price: \$7.25 | Target Price: \$14.00

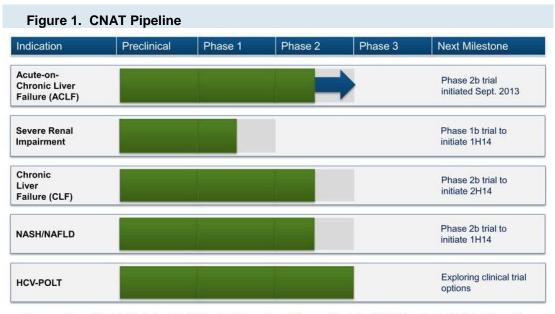
### **INVESTMENT HIGHLIGHTS**

Conatus to initiate two new studies in 1H14; reiterate Market Outperform rating and \$14 price target based on a risk-adjusted, discounted cash flow analysis. Today, Conatus announced plans to initiate two additional clinical trials for its pan-caspase inhibitor emricasan -- a Phase 2b trial in NASH and a Phase 1b trial in liver failure patients with severe renal impairment (Figure 1). The NASH study is designed to establish a dosing strategy for this population. The severe renal impairment study is designed to broaden the scope of the liver failure program to include patients with severe renal impairment. Together, we view these upcoming studies as positive signs that Conatus is aggressively moving forward with emricasan development, providing additional incremental catalysts for the stock next year.

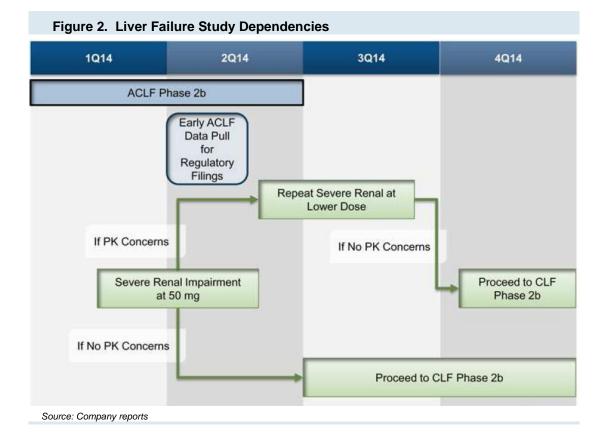
Phase 2b trial in NASH. Conatus plans to enroll ~40 patients for 28 days of dosing and to use the study to characterize the effect of emricasan on key biomarkers (cCK18 and ALT). If the biomarkers perform as expected based on studies of emricasan in other settings, development in NASH could be more straightforward. Management commented that if the initial study is successful, the next step could be a trial with chronic dosing for a full year; however, Conatus would likely wait for data from the placebo arms of a competitive program from Gilead to gain information about the natural history of the disease before moving into a larger NASH trial itself. Although there are some mechanistic safety concerns related to long-term dosing, emricasan is currently cleared by the FDA for dosing for up to two years and recent cCK18 studies have shown that while emricasan reduces unnaturally high apoptosis rates it does not reduce rates below normal levels.

**Phase 1b trial in severe renal impairment.** The trial is designed as an open-label study in the U.S. with eight patients on a 50 mg dose and eight matching controls. Conatus plans to understand the pharmacokinetic and safety/tolerability profile with the goals of broadening eligibility requirements in future liver failure studies and identifying any dose adjustments that may be needed in this population. Results are expected in 2Q14 and, along with the acute on chronic liver failure (ACLF) study, will feed into the next chronic liver failure (CLF) study, slated to start in 2H14 (Figure 2).





Also supporting a pilot clinical study funded by National Institute on Alcohol Abuse and Alcoholism (NIAAA) in patients with alcoholic hepatitis Source: Company reports



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# **Company Description**

Conatus Pharmaceuticals is a San Diego-based biopharmaceutical company focused on the development of emricasan, a pan-caspase inhibitor with the potential to be used to treat liver fibrosis, particularly in areas of large unmet need, including cirrhosis-induced liver failure.

#### **Investment Risks**

Clinical Risk. Emricasan has not yet been evaluated in clinical trials longer than 12 weeks. Longer trials may result in unanticipated safety concerns, which could cause emricasan to underperform in clinical trials. Although successful in the regulation of biomarkers, emricasan may not impact clinical outcomes.

Regulatory Risk. Conatus hopes to use emricasan in acute and chronic indications. However, due to potential on-target activities that may increase cancer risk, FDA maybe unwilling to approve the drug in a chronic setting.

Intellectual Property Risk. The composition of matter patent for emricasan expires in 2017 and has not yet received a Hatch Waxman extension. As such, Conatus may have to rely on a polymorph composition and method patent, which expires in 2027, for long-term market exclusivity. If Conatus does not receive orphan exclusivity for emricasan, protection may also be limited.

Commercial Risk. As a small company, Conatus may have difficulty educating healthcare payers and providers on the benefits of a novel drug. As such, emricasan adoption may be slowed.

Sector Risk. Valuation of biopharmaceutical stocks is subject to both investor assessments of the prospects of the underlying companies, as well as risk tolerance and the level of confidence in the prospects of pharmaceutical stocks as a group. Therefore, Conatus' stock price may fall even while the company meets or exceeds investor expectations.



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JMP Securities was manager or co-manager of a public offering, and received compensation for doing so, for Conatus Pharmaceuticals Inc. in the past 12 months.

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Market Outperform (MO): JMP Securities expects the stock price to outperform relevant market indices over the next 12 months.

Market Perform (MP): JMP Securities expects the stock price to perform in line with relevant market indices over the next 12 months.

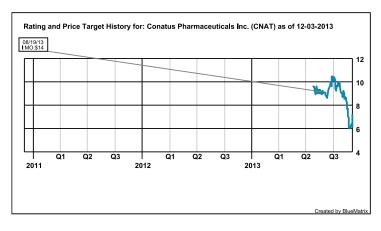
Market Underperform (MU): JMP Securities expects the stock price to underperform relevant market indices over the next 12 months.

JMP Securities Research Ratings and Investment Banking Services: (as of December 3, 2013)

							# Co's	
							Receiving	
		# Co's	%		# Co's	%	Services in	% of Co's
	Regulatory	Under	of	Regulatory	Under	of	Past 12	With This
JMP Rating	Equivalent	Coverage	Total	Equivalent	Coverage	Total	Months	Rating
MARKET OUTDERSONA	_	207	5 4 700/		207	5 4 700/	o <del>-</del>	00 000/
MARKET OUTPERFORM	Buy	227	54.70%	Buy	227	54.70%	87	38.33%
MARKET PERFORM	Hold	138	33.25%	Hold	138	33.25%	25	18.12%
MARKET UNDERPERFORM	Sell	5	1.20%	Sell	5	1.20%	0	0%
COVERAGE IN TRANSITION		45	10.84%		45	10.84%	0	0%
TOTAL:		415	100%		415	100%	112	26.99%

#### **Stock Price Chart of Rating and Target Price Changes:**

Note: First annotation denotes initiation of coverage or 3 years, whichever is shorter. If no target price is listed, then the target price is N/A. In accordance with NASD Rule 2711, the chart(s) below reflect(s) price range and any changes to the rating or price target as of the end of the most recent calendar quarter. The action reflected in this note is not annotated in the stock price chart. Source: JMP Securities.



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## **Conatus Pharmaceuticals Inc. (CNAT)**



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