

# **Enanta Pharmaceuticals, Inc.** (ENTA)

Competition Posts Phase 2 Data Supporting Eight-Week Regimen

MARKET DATA	
Price	\$19.57
52-Week Range:	\$14.31 - \$22.40
Shares Out. (M):	17.4
Market Cap (\$M):	\$340.5
Average Daily Vol. (000):	51.0
Cash (M):	\$109
LT Debt (M):	\$0
Source: Thomson Reuters and JMP Securities LLC	

FY SEP		2012A	2013E	2014E
Revenue (\$M)	1Q		\$27.9	
	2Q		\$1.4	
	3Q		\$1.4	
	4Q		\$26.4	
	FY	\$41.7	\$57.1	\$48.8
EPS	1Q		\$1.53	
3	2Q		(\$0.35)	
	3Q		(\$0.30)	
	4Q		\$0.94	
	FY	\$1.13	\$1.99	\$0.59
	CY	\$2.38		
Source: Company re	ports and	d JMP Securities	LLC	



MARKET OUTPERFORM | Price: \$19.57 | Target Price: \$25.00

### **INVESTMENT HIGHLIGHTS**

Competition posts Phase 2 data supporting an eight-week regimen; reiterate Market Outperform rating on Enanta Pharmaceuticals and \$25 price target based on a risk-adjusted, discounted cash flow analysis. Yesterday, Gilead released interim data from the LONESTAR study. The study evaluated the benefit of sofosbuvir (SFS) and ledipasvir (LED), with and without ribavirin for eight and 12 weeks in treatment naive patients, and for 12 weeks in experienced patients who failed a protease inhibitor, half of which were compensated cirrhotics. Every arm, regardless of patient population, duration of therapy, addition of ribavirin or liver condition, achieved SVR rates of over 95% (Figure 1). In our opinion, these data support the hypothesis that ribavirin will not be needed, making the Gilead combination a true once a day therapy. That being said, this is a single center study, and so we expect the data to be less impressive in a broader patient population. We note that Gilead will run a Phase 3 study in treatment naive noncirrohtic patients that will be eight weeks in duration with and without ribavirin, and 12 weeks without ribavirin, ION-3, which will be part of the registration package for the combination. We continue to believe that despite some emerging advantages for the Gilead regimen, feedback from physicians suggests room for more than one approach. We should get a better idea of the comparative efficacy of each regimen by year end when Phase 3 data begins to report. We anticipate 25% penetration at peak for the ABBV/ENTA combination.



# FIGURE 1. LONESTAR Data

	GT1 naïve	GT1 naïve	GT1 naïve	GT1 exp	GT1 exp
	SFS/LED	SFS/LED/RB	SFS/LED	SFS/LED	SFS/LED/RBV
	8wks	V 8 wks	12 wks	12 wks	12 wks
n	20	21	19	19	21
Compensated cirrohsis	0%	0%	0%	50%	
SVR4			100%	95%	95%
SVR8	95%	100%			
relapse	5%			5%	

Source: Company reports



## **Company Description**

Enanta is a Watertown, Massachusetts based biotechnology company focused on anti-infectives. The company has partnered a protease inhibitor program with AbbVie Pharmaceuticals and an NS5a program with Novartis, as well as developing fully owned assets. The lead protease inhibitor, ABT-450, partnered with AbbVie, is in Phase 3 development.

### **Investment Risks**

Clinical risk. Drug development is a risky and capital-intensive endeavor. The vast majority of drugs that enter clinical

development fail to reach the market. Enanta's Phase 3 program with AbbVie may experience development setbacks; we point specifically to safety as a source of risk. In addition, Enanta has many early stage assets that may or may not make it to development in humans.

Regulatory risk. Enanta is reliant on its pharmaceutical partners, AbbVie and Novartis, to move its drug candidates through registration with the FDA and EMA and it is dependent on the pace of these regulatory entities to approve new drugs. Enanta's early stage HCV assets are from classes that have been placed on clinical hold, leading to increased scrutiny.

Intellectual Property risk. Enanta's lead clinical assets are covered by approved patents; however, other assets have patents pending. Patent expirations can result in a negative impact to sales. Additionally, generic companies may file abbreviated new drug applications to challenge current products with patent protection.

Commercial risk. Enanta is reliant upon their pharmaceutical partners, AbbVie and Novartis, to successfully commercialize assets. The HCV space is very competitive and Enanta's assets may lose share as new competitors come to market.

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

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The research analyst(s) who prepared this report does/do hereby certify that the views presented in this report are in accordance with my/our personal views on the securities and issuers discussed in this report. As mandated by SEC Regulation AC no part of my/our compensation was, is or will be directly or indirectly related to the specific views or recommendations expressed herein. This certification is made under the obligations set forth in SEC Regulation AC. Any other person or entity may not use it for any other purpose. This certification is made based on my/our analysis on the date of this report's publication. I/We assume no obligation to update this certification to reflect any facts, circumstances or events that may subsequently come to my/our attention. Signed Liisa A. Bayko and Heather Behanna

#### JMP Securities Disclosure Definitions:

JMP Securities currently makes a market in the security of Enanta Pharmaceuticals, Inc.

JMP Securities was manager or co-manager of a public offering for Enanta Pharmaceuticals, Inc. in the past 12 months.

#### **JMP Securities Investment Opinion Definitions:**

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.

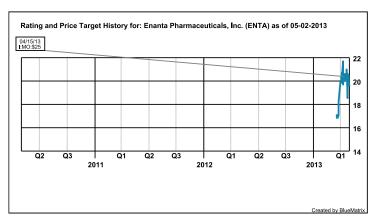
Market Underperform (MU): JMP Securities expects the stock price to underperform relevant market indices over the next 12 months.

JMP Securities Research Ratings and Investment Banking Services: (as of May 2, 2013)

							# Co's	
							Receiving	
							IB	
		# 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 OUTPERFORM	Buy	233	60.52%	Buy	233	60.52%	66	28.33%
MARKET PERFORM	Hold	145	37.66%	Hold	145	37.66%	16	11.03%
MARKET UNDERPERFORM	Sell	7	1.82%	Sell	7	1.82%	0	0%
TOTAL:		385	100%		385	100%	82	21.30%

#### **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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#### **Enanta Pharmaceuticals, Inc. (ENTA)**



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Jeffrey H. Spurr Director of Research (415) 835-3903

### **RESEARCH PROFESSIONALS**

### **FINANCIAL SERVICES**

		Medical Devices	
Asset Managers David Trone	(212) 906-3525	David Turkaly	(212) 906-3563
Chris Ross, CFA	(212) 906-3523	John Gillings	(212) 906-3564
Chins Ross, Cl A	(212) 900-3332	John Chings	(212) 900-3304
Commercial & Specialty Finance		Medical Devices & Molecular Diagnostics	
Christopher York	(415) 835-8965	J. T. Haresco, III, PhD	(415) 869-4477
Kevin Chen	(404) 848-7774	Ralph Fong	(415) 835-8916
TOVIII SHOT	(101) 010 1111	b 5g	( )
Consumer Finance			
David M. Scharf	(415) 835-8942	REAL ESTATE	
Kevane A. Wong	(415) 835-8976		
<b>.</b>	( -,	Housing & Land Development	
Financial Processing & Outsourcing		Peter L. Martin, CFA	(415) 835-8904
David M. Scharf	(415) 835-8942	Aaron Hecht	(415) 835-3963
Kevane A. Wong	(415) 835-8976		
rtovano / t. vvong	(110) 000 0010	Lodging & Property Services	
Insurance		William C. Marks	(415) 835-8944
Matthew J. Carletti	(312) 768-1784	Whitney Stevenson	(415) 835-8948
Christine Worley	(312) 768-1786		
Offitiguite Worley	(312) 700-1700	REITs: Healthcare	
Investment Banks & Brokers		Peter L. Martin, CFA	(415) 835-8904
David Trone	(212) 906-3525	Aaron Hecht	(415) 835-3963
Chris Ross, CFA	(212) 906-3532	Brian Nelson	(617) 235-8516
Ollis 1033, Ol A	(212) 900-3332		
Mortgage Finance		REITs: Office, Industrial, & Diversified	
		Mitch Germain	(212) 906-3546
Steven C. Del anev	(404) 848-7773	WILLOT Germain	(212) 900-3340
Steven C. DeLaney Trevor Cranston, CFA	(404) 848-7773 (415) 869-4431	Willer German	(212) 900-3340
Trevor Cranston, CFA	(415) 869-4431		(212) 900-3340
		TECHNOLOGY	(212) 900-3340
Trevor Cranston, ĆFA Charter Robinson	(415) 869-4431	TECHNOLOGY	, ,
Trevor Cranston, CFA	(415) 869-4431	TECHNOLOGY  Communications Equipment & Internet S	ecurity
Trevor Cranston, ĆFA Charter Robinson  HEALTHCARE	(415) 869-4431	TECHNOLOGY  Communications Equipment & Internet Serik Suppiger	ecurity (415) 835-3918
Trevor Cranston, ĆFA Charter Robinson  HEALTHCARE  Biotechnology	(415) 869-4431 (757) 613-8955	TECHNOLOGY  Communications Equipment & Internet S	ecurity
Trevor Cranston, ĆFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko	(415) 869-4431 (757) 613-8955 (312) 768-1785	TECHNOLOGY  Communications Equipment & Internet Security Erik Suppiger Christopher Slaymaker	ecurity (415) 835-3918
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D.	(312) 768-1785 (312) 768-1795	TECHNOLOGY  Communications Equipment & Internet Serik Suppiger Christopher Slaymaker  Internet & Digital Media	ecurity (415) 835-3918 (415) 835-3920
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD	(312) 768-1785 (312) 768-1795 (212) 906-3505	TECHNOLOGY  Communications Equipment & Internet Serik Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III	ecurity (415) 835-3918 (415) 835-3920 (212) 906-3528
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D.	(312) 768-1785 (312) 768-1795	TECHNOLOGY  Communications Equipment & Internet Serik Suppiger Christopher Slaymaker  Internet & Digital Media	ecurity (415) 835-3918 (415) 835-3920
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD	(312) 768-1785 (312) 768-1795 (212) 906-3505	TECHNOLOGY  Communications Equipment & Internet Start Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg	ecurity (415) 835-3918 (415) 835-3920 (212) 906-3528
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD Christopher T. Radom, PhD	(312) 768-1785 (312) 768-1795 (212) 906-3505 (212) 906-3519	TECHNOLOGY  Communications Equipment & Internet Serik Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg  Software	ecurity (415) 835-3918 (415) 835-3920 (212) 906-3528 (212) 906-3537
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD Christopher T. Radom, PhD Michael G. King, Jr.	(312) 768-1785 (312) 768-1795 (212) 906-3505 (212) 906-3520	TECHNOLOGY  Communications Equipment & Internet Serik Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg  Software Patrick Walravens	ecurity (415) 835-3918 (415) 835-3920 (212) 906-3528 (212) 906-3537 (415) 835-8943
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD Christopher T. Radom, PhD Michael G. King, Jr. Carter L. Gould Eric Joseph John L. Newman, PhD	(312) 768-1785 (312) 768-1785 (312) 768-1795 (212) 906-3505 (212) 906-3519 (212) 906-3520 (212) 906-3522	TECHNOLOGY  Communications Equipment & Internet Sterik Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg  Software Patrick Walravens Peter Lowry	ecurity (415) 835-3918 (415) 835-3920 (212) 906-3528 (212) 906-3537 (415) 835-8943 (415) 869-4418
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD Christopher T. Radom, PhD Michael G. King, Jr. Carter L. Gould Eric Joseph John L. Newman, PhD	(312) 768-1785 (312) 768-1785 (312) 768-1795 (212) 906-3505 (212) 906-3519 (212) 906-3520 (212) 906-3522 (212) 906-3514	TECHNOLOGY  Communications Equipment & Internet Serik Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg  Software Patrick Walravens	ecurity (415) 835-3918 (415) 835-3920 (212) 906-3528 (212) 906-3537 (415) 835-8943
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD Christopher T. Radom, PhD  Michael G. King, Jr. Carter L. Gould Eric Joseph John L. Newman, PhD Caroline Palomeque	(312) 768-1785 (312) 768-1785 (312) 768-1795 (212) 906-3505 (212) 906-3519 (212) 906-3520 (212) 906-3522 (212) 906-3514 (212) 906-3510	Communications Equipment & Internet Sterik Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg  Software Patrick Walravens Peter Lowry Greg McDowell	ecurity (415) 835-3918 (415) 835-3920 (212) 906-3528 (212) 906-3537 (415) 835-8943 (415) 869-4418 (415) 835-3934
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD Christopher T. Radom, PhD Michael G. King, Jr. Carter L. Gould Eric Joseph John L. Newman, PhD	(312) 768-1785 (312) 768-1785 (312) 768-1795 (212) 906-3505 (212) 906-3519 (212) 906-3520 (212) 906-3522 (212) 906-3514 (212) 906-3510	Communications Equipment & Internet Secrit Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg  Software Patrick Walravens Peter Lowry Greg McDowell  Wireless & Cloud Computing Technologie	(415) 835-3918 (415) 835-3920 (212) 906-3528 (212) 906-3537 (415) 835-8943 (415) 869-4418 (415) 835-3934
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD Christopher T. Radom, PhD  Michael G. King, Jr. Carter L. Gould Eric Joseph John L. Newman, PhD Caroline Palomeque	(312) 768-1785 (312) 768-1785 (312) 768-1795 (212) 906-3505 (212) 906-3519 (212) 906-3520 (212) 906-3522 (212) 906-3514 (212) 906-3510	Communications Equipment & Internet Sterik Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg  Software Patrick Walravens Peter Lowry Greg McDowell  Wireless & Cloud Computing Technological	(415) 835-3918 (415) 835-3920 (212) 906-3528 (212) 906-3537 (415) 835-8943 (415) 869-4418 (415) 835-3934 (415) 835-8998
Trevor Cranston, ČFA Charter Robinson  HEALTHCARE  Biotechnology Liisa A. Bayko Heather Behanna, Ph.D. Jason N. Butler, PhD Christopher T. Radom, PhD  Michael G. King, Jr. Carter L. Gould Eric Joseph John L. Newman, PhD Caroline Palomeque  Healthcare Services & Facilities	(312) 768-1785 (312) 768-1785 (312) 768-1795 (212) 906-3505 (212) 906-3519 (212) 906-3520 (212) 906-3522 (212) 906-3514 (212) 906-3510 (212) 906-3509	Communications Equipment & Internet Secrit Suppiger Christopher Slaymaker  Internet & Digital Media Ronald V. Josey III Peter Lunenburg  Software Patrick Walravens Peter Lowry Greg McDowell  Wireless & Cloud Computing Technologie	(415) 835-3918 (415) 835-3920 (212) 906-3528 (212) 906-3537 (415) 835-8943 (415) 869-4418 (415) 835-3934

## **ADDITIONAL CONTACTS**

Thomas R. Wright Director of Equities (212) 906-3599

Brian Nelson

Dan Wychulis Director of Institutional Sales (617) 235-8530

(617) 235-8516

**600 Montgomery Street, Suite 1100** San Francisco, CA 94111 www.jmpsecurities.com