Merrimack Pharmaceuticals March 15, 2013

(MACK/ NASDAQ)



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MACK: 4Q Results; ASCO Looks Lighter Than We Expected With More Data in 2H13

PT: \$14.00

Investment Summary

NAPOLI-1 Now to Readout in 2H13. Merrimack amended the NAPOLI-1 protocol to add a third arm, which has resulted in a readout being pushed back slightly from mid-2013 to 2H13.

Back Half of 2013 Will Be Data Heavy for MM-121. While we expect Phase 2 NSCLC data at AACR and possibly updated at ASCO in June, the majority of ongoing studies for MM-121 likely won't yield data until 2H13. There are four phase 2 trials being conducted for MM-121 in breast, ovarian, and non-small cell lung cancer, and we expect data from each of these studies to read out in 2H13. We do expect the mutant EGFR, NSCLC cohort to read out earlier, in 1H13.

Merrimack reported Earnings of \$(1.28) Per Share. The loss of \$(1.28) per share was in line but slightly below our estimate of \$(1.26) per share for FY12. Merrimack ended the quarter with ~\$110M in cash including a \$40M line of credit, which we believe will fund operations into 2014.

Discussion

Amendment to the NAPOLI-1 Study Cause Data to Be Pushed Back. NAPOLI-1 is Merrimack's only Phase 3 study and comparing MM-398 vs 5-FU/Leucovorin in 2nd line pancreatic cancer. The study was amended to add 5-FU/Leucovorin/MM-398 as a third arm due to the prevalence of FOLFIRI use, and adoption of the amended protocol has caused data to be delayed from mid-2013 to 2H13. MACK expects the MM-398 monotherapy arm to show a survival difference of approximately six-weeks compared to 5-FU Leucovorin alone and up to a three month difference when the combination of MM-398 and 5-FU Leucovorin is compared to 5-FU Leucovorin alone.

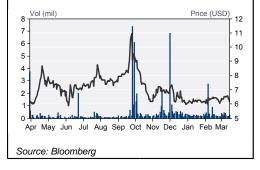
MM-121 is currently being evaluated in Phase II trials for ovarian, breast and lung cancer and these studies will read out in 2013. The Phase 2 study of MM-121/Tarceva for NSCLC has two cohorts, EGFR mutant and wild type, and we expect data from the mutant EGFR cohort to be presented at AACR and possibly an update at ASCO in June, but that we have wait until 2H13 for the EGFR WT cohort. In 2H13, we also expect the data from ER/PR+ HER2- metastatic breast cancer in patients treated with Exemestane w/wo MM-121. The phase 2 study of platinum resistant ovarian cancer patients treated with paciltaxel w/wo MM-121, which had been expected to read out in 2014 has recruited more guickly then expected and will read out also in 2H13.

Merrimack Reports FY12 Loss of \$(1.28) per Share. The loss suffered by Merrimack of \$(1.28) per share for FY12 was in line with \$(1.26) per share we modeled. Merrimack ended the quarter with ~\$110M in cash including a \$40M line of credit, which we believe will fund operations into 2014.

Valuation / Target Price

Our \$14 target price for MACK shares is based on the NPV of MM-398 global revenue (discounted 20%) in pancreatic cancer alone, which we forecast will peak at \$800M (see Exhibit 1). We assume orphan pricing for MM-398 peaking at \$80k per course in the US with conservative peak penetration in the US (35%) and ex-U.S. (25%). MACK's early stage pipeline, which is not included in our revenue forecasts, lends significant upside to our current valuation.

Price			\$6.15
52-Week High,	/Low		\$11.11-\$5.66
Shares Out (m	m)		94.2
Market Cap (m	ım)		\$579
Avg. Daily Vol	(000)		308,742
Short Interest			3.2%
Cash (mm)			NA
EV (mm)			NA
Book Value / S	hare		NA
EPS	FY12A	FY13E	FY14E
Mar	\$(2.14)	\$(0.25)	
June	\$(0.22)	\$(0.26)	
Sept	\$(0.25)	\$(0.26)	
Dec	\$(0.26)	\$(0.28)	
FY (Dec)	\$(1.28)	\$(1.05)	\$(0.90)
Prior:	\$(1.26)	\$(0.78)	
P/E (x)	NM	NM	NM
Revenue (\$M)			
FY (Dec)	\$48.9	\$50.0	\$81.7
Prior:	\$45.7		



116

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P/S (x)

Exhibit 1: MACK Milestones

MM-398 (encapsulated irinotecan)	Indication/Setting	Comment	Timing
Phase 3 (NAPOLI-1)	pancreatic (2nd line)	1°= OS; N= 405; Randomization= MM-398 vs. 5FU Leucovorin vs 5FU Leucovorin + MM-398	YE13
Phase 2	CRC	Investigator sponsored studies that will evaluate MM-398+5FU Leucovorin vs. FOLFIRI	Pending update
MM-121 (ErbB3 Mab)	Indication/Setting	Comment	Timing
Phase 2	NSCLC (EGFR naïve, EGFR refractory, EGFR responders)	1°= PFS; N= 260; Randomization= RMM-121 + Tarceva vs. Tarceva; Mutant EGFR; Phase 1 dose escalation into Phase 2	1H13
Phase 2	breast (mets HER-, ER+, PR+)	1°= PFS; N= 130; Randomization= MM-121 + Exemestane vs. Exmestane in anti-estrogen failures	2H13
Phase 2	breast (neoajvnt HER-, ER+)	1°= RR; N= 200; Randomization= MM-121 + paclitaxel vs. paclitexal in neo-adjuvant setting	2H13
Phase 2	Ovarian (platinum resistant)	1°= PFS (38 wks); N= 210; Randomization= 2:1, MM-121 + paclitaxel vs. paclitexal	2H13
Phase 2	NSCLC (EGFR naïve, EGFR refractory, EGFR responders)	1°= PFS; N= 260; Randomization= RMM-121 + Tarceva vs. Tarceva; WT EGFR; Phase 1 dose escalation into Phase 2	2H13
MM-111 (ErbB3, ErbB2 bi-specif)	Indication/Setting	Comment	Timing
Phase 2	HER2+ and FISH+/FISH- gastric cancer	1°= PFS, N= 180; Randomization= paclitax el +/- MM-111 +/- Herceptin	2014

Source: Company Reports and Brean Capital, LLC estimates

Exhibit 2: MACK Income Statement

Merrimack Pharmaceuticals INCOME STATEMENT Brean Murray, Carret & Co. Gene Mack, 212.702.6616																				
Fiscal Period: ends Dec. 31		2010		2011	1Q	2Q	3	Q	4Q	2012		1Q	2Q	3QE	4QE	201	BE	2014E		2015E
MM-398 US sales MM-398 ex-US sales Total WW sales MM-398	\$ \$ \$	1.1.1	\$ \$	-	\$ - \$ - \$ -	\$ -	\$ - \$ -	\$	- \$ - \$ - \$	-	\$ \$	- \$ - \$ - \$	- \$ - \$ - \$	- \$ - \$ - \$		\$ \$ \$	\$	26.7 - 26.7	\$ \$ \$	166.8 23.6 190.4
Product sales, net	-					<u> </u>				<u> </u>			<u>:</u>					26.7		190.4
Collaboration revenue Other Revenue		20.3		34.2	11.3	12.1	11.		14.2	48.9		11.5	12.0	12.5	14.0	50		55.0		60.0
Total Revenue	\$	20.3	\$	34.2	\$ 11.3	\$ 12.1	\$ 11.	.3 \$	14.2 \$	48.9	\$	11.5 \$	12.0 \$	12.5 \$	14.0	\$ 50	0 \$	81.7	\$	250.4
Cost of product sales	1	-		-					- '	-	ľ	- '	- '	- '	-		1	3.5		22.9
Gross Profit		20.3		34.2	11.3	12.1	11.		14.2	48.9		11.5	12.0	12.5	14.0	50		78.3		227.6
Research and development		58.3		100.6	31.7	28.8	30.	.9	34.6	125.9		31.7	33.0	34.0	36.3	135	0	145.0		150.0
Selling, general and administrative		11.4		14.5	3.7	3.6	4.	.3	4.2	15.8		5.0	5.2	5.5	6.3	22	0	26.6		76.6
Confingent consideration		(0.2)																		ļ
Amortization of intangible assets		-		-	-	-	-		-	-		-	-	-	-			-		-
Total operating expense (including COGS)		69.5		115.1	35.4	32.4	35.		38.7	141.7		36.7	38.2	39.5	42.6	157		175.0		249.5
Operating Income		(49.2)		(80.9)	(24.0)	(20.3)	(23.	.9) (24.5)	(92.7)		(25.2)	(26.2)	(27.0)	(28.6)	(107	0)	(93.3)		0.9
Interest Income		0.1		0.1	-	-	-	•	- [-		-	-	-	-			-		-
Interest expense		(3.7)		(0.0)						-		-	-	-	-			-		-
Other Income/ Expense		2.7		1.2	0.6	0.2	0.		(0.4)	1.0		-	-	-	-			-		-
net loss (gain) from non-controlling interest	<u> </u>	0.1		0.5 (79.2)	0.1	0.1	0.		0.1	0.5	_	- (05.0)	- (00.0)	- (07.0)	- (00.0)	//07		- (00.0)		-
Net Income Before Taxes Provision for Income Taxes	-	(50.1)		(19.2)	(23.3)	(20.0)	(23.	.2) (24.8)	(91.3)	_	(25.2)	(26.2)	(27.0)	(28.6)	(107	υ)	(93.3)		0.9 0.2
Net Income After Taxes	\$	(50.1)	\$	(79.2)	\$ (23.3)	\$ (20.0)	\$ (23.	.2) \$ (24.8) \$	(91.28)	\$	(25.2) \$	(26.2) \$	(27.0) \$	(28.6)	\$ (107	0) \$	(93.3)	\$	0.7
Basic Weighted Average Shares		11.0		11.3	11.8	90.6	93.	.7	94.7	72.8		100.7	101.7	102.7	103.7	102		103.2		104.2
Diluted Weighted Average Shares		11.0		11.3	11.8	90.6	93.	.7	94.7	72.8		100.7	101.7	102.7	103.7	102	2	103.2		104.2
Basic EPS as-reported (GAAP)	\$	(4.56)	\$	(6.98)	\$ (1.97)	\$ (0.22)	\$ (0.2	25) \$ (0.26) \$	(1.28)	\$	(0.25) \$	(0.26) \$	(0.26) \$	(0.28)	\$ (1.0	5) \$	(0.90)	\$	0.01
Diluted EPS as-reported (GAAP)	\$	(5.57)	\$	(7.67)	\$ (2.14)	\$ (0.22)	\$ (0.2	!5) \$ (0.26) \$	(1.28)	\$	(0.25) \$	(0.26) \$	(0.26) \$	(0.28)	\$ (1.0	5) \$	(0.90)	\$	0.01
Cash and Equivalents	\$	31	\$	50		\$ 106	\$ 8	7	\$	48						\$ 7	8 \$	21	\$	59
E=Estimate	٠	'-		٠					•						•		-			

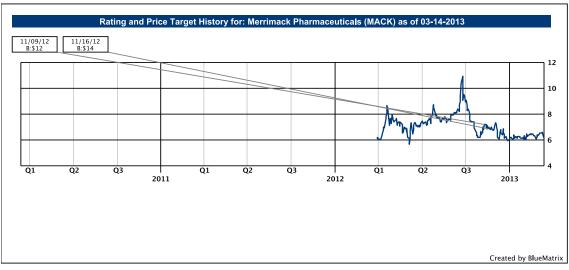
Source: Company Reports and Brean Capital, LLC estimates

Risks

Key risks to MACK shares include: (1) significant clinical failure risk of MM-398 or any of the company's other candidates in ongoing and planned clinical trials; (2) regulatory risk stemming from FDA perception of adequate data support for approval of MM-398 in any of the indications in which it is being developed or any of the other candidates in MACK's pipeline; and (3) financial risk from disappointing outcomes to MM-121 clinical studies should Sanofi choose to end the co-development arrangement; (4) risk from failure to successfully invalidate third party patent claims in an ongoing opposition proceeding in Europe that could otherwise limit the ability of MM-121 and MM-111 to be commercialized in Europe.

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			IB Serv./ Past 12Mos.		
Rating Category	Count	Percent	Count	Percent	
BUY	141	65.58%	8	5.67%	
HOLD	68	31.63%	2	2.94%	
SELL	6	2.79%	1	16.67%	
NOT RATED					

Note: Stock price volatility may cause temporary non-alignment of some ratings with some target prices.

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