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Pacira Pharmaceuticals, Inc. (PCRX - OUTPERFORM): New Data Shows Reduction in Opioid Burden with Exparel and Is Likely To Support Launch if Approved by Oct. 28th PDUFA, in our View. Reiterate OUTPERFORM

Price: \$9.25 Fair Value Estimate: \$20

- A retrospective analysis comparing Exparel to regular bupivacaine showed a longer time to first opioid use, reduction in opioid-related adverse events, reduction in opioid consumption, and better pain control with Exparel use. The data were generated from a pooled analysis of five active-control, double-blind, randomized, multicenter, parallel-group trials with over 700 patients across three surgical models (hemorrhoidectomy, total knee arthroplasty, and hernia repair). Results demonstrated that Exparel was statistically superior (P<0.0001) to bupivacaine on all four endpoints: 1) median time to first opioid (TTFO) was 3.5 times longer for Exparel patients versus bupivacaine patients (9.9 hrs vs. 2.7 hrs); 2) a 50% reduction in the total amount of morphine-equivalent opioids consumed for Exparel patients versus bupivacaine patients (7.9 mg vs. 15.8 mg); 3) a 46% reduction in the mean number of opioid-related adverse events (ORAEs) for Exparel patients versus bupivacaine (0.25 vs. 0.46); and 4) less pain over 72 hours for Exparel patients versus bupivacaine patients as measured by area under the curve (AUC) analysis (AUC0-72 315 vs. 427). (See page 2 for results table.) The data were presented at the 2011 Annual Meeting of the American College of Clinical Pharmacy (ACCP; October 16-19; Pittsburgh).
- We believe this retrospective data will be an important marketing tool for commercial launch and expect additional retrospective and prospective health outcome data to be presented over the coming months. In our view, the retrospective data presented at ACCP goes a long way in quantifying the reduction in opioid burden for patients using Exparel versus bupivacaine and should serve as an important marketing piece as most physicians are aware of the improved recovery experience by patients who use less opioids. Furthermore, the company is also generating data to support the pharmacoeconomic benefit of Exparel by quantifying the increase in cost related to treating an ORAE, including costs related to length of stay. We believe these retrospective and prospective data demonstrating both the clinical and economic benefit of Exparel will provide a compelling rationale for adoption by physicians. We also believe Pacira's comprehensive and detailed launch strategy highlights management's expertise in successful hospital-based commercial drug launches.
- The next major catalyst for Pacira is the PDUFA date of October 28, 2011 for Exparel treatment of postsurgical pain management. We estimate a 75% chance that Exparel is approved on its PDUFA date and think the stock could double on approval. With the recent pullback in PCRX shares, we see a favorable risk/reward scenario and remain confident in approval for three main reasons. First, the efficacy evidence for the NDA is based on two positive Phase 3 trials of Exparel which the FDA acknowledged at a pre-NDA meeting in February 2010 to be appropriately designed to evaluate safety and efficacy. Second, the safety profile of Exparel seems adequate for approval and while the FDA typically requires a 500-patient safety database for pain drugs, Pacira submitted data for over 1300 patients. Third, the FDA is already familiar with bupivacaine, the active ingredient in Exparel, as well as Pacira's DepoFoam technology. Furthermore, the hemorrhoidectomy (soft-tissue surgery) and bunionectomy (orthopedic surgery) indications were selected to support a broad label for use as a single dose local administration into the surgical wound to produce postsurgical analgesia. We believe Exparel is likely to receive this broad label given that other pain drugs such as OFIRMEV, Caldolor, and Zipsor were all recently approved with broad labels for pain.
- We reiterate our OUTPERFORM rating and fair value of \$20. Our fair value is calculated based on the sum-of-parts for each drug/indication combination using a 30% annual discount from our peak annual revenues projections and 1-10x multiple, depending on stage of development to reflect risk.
- Risks to our fair value include regulatory and commercial risk for Pacira's lead drug candidate Exparel. The PDUFA date for
 Exparel is October 28, 2011 and there is always risk that the FDA could delay approval. On the commercial front, Exparel
 will compete with generic forms of regular bupivacaine and, although we believe Exparel offers therapeutics advantages,
 some physicians may opt for the cheaper alternative.

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ACCP PRESENTATION

Poster Title: Bupivacaine Extended Release Liposome Injection (DepoFoam® bupivacaine) vs. Bupivacaine HCI: A Meta-analysis of Multimodal Trials of Doses Up to and Including 300 mg

Figure 1: Retrospective Analysis Comparing Exparel to Bupivacaine on Opioid Burden

	Exparel (75-300mg)	Bupivacaine (75-200mg)	P Value
AUC 0-72 hr	315	427	P<0.0001
Median TTFO	9.9 hours	2.7 hours	P<0.0001
Morphine	7.9 mg	15.8 mg	P<0.0001
ORAEs per patient	0.25	0.46	P<0.0001

^{*}TTFO=Time To First Opioid use; ORAE=Opioid-related Adverse Event

Source: Company data, Wedbush Securities, Inc.

POTENTIAL UPCOMING MILESTONES (*OUR ESTIMATES)

October 23-27, 2011 Participation at American College Surgeons (ACS; San Francisco, CA)

October 28, 2011 Exparel (infiltration) PDUFA date

2011* Select clinical candidate from DepoNSAID program
2011/2012* Potential partnership for ex-US rights of Exparel
2011/2012* Potential additional technology partnerships
Q1:12 US launch of Exparel (infiltration)

H1:12* Initiate pivotal trial of Exparel in nerve block H2:12* Pivotal trial data for Exparel in nerve block



Analyst Certification

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Wedbush Equity Research Disclosures as of October 19, 2011

Company	Disclosure
Pacira Pharmaceuticals, Inc.	1,3,4,5,7

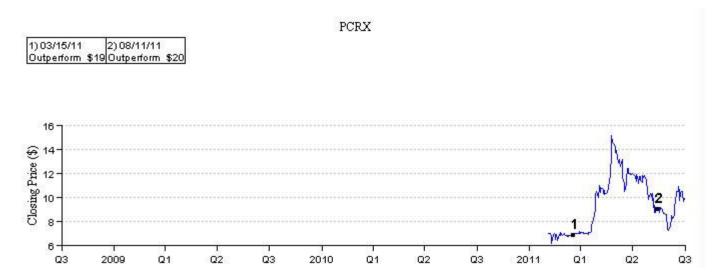
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