

Aratana Therapeutics, Inc. (PETX)

Productive 2013 Solidifies Leadership Position Ahead of Key Milestones; Reiterate Market Outperform

MARKET DATA

Price	\$17.60
52-Week Range:	\$6.56 - \$29.32
Shares Out. (M):	30.3
Market Cap (\$M):	\$533.3
Average Daily Vol. (000):	301.0
Cash (M):	\$46

Source: Thomson Reuters and JMP Securities LLC

MARKET OUTPERFORM | Price: \$17.60 | Target Price: \$38.00

INVESTMENT HIGHLIGHTS

We reiterate our Market Outperform rating and \$38 price target on shares of Aratana Therapeutics following 4Q13 results. For the quarter and the full year, Aratana reported \$123K in revenue related to its lymphoma franchise. The company reported EPS of \$0.33 for the quarter and (\$0.39) for the full year and ended the year with \$45.8M in cash and equivalents, which we believe should be sufficient to fund operations until 2015. The company had a very fruitful year as it signed two option agreements, moved all key products in the pipeline forward, and expanded its pipeline through the acquisition of Vet Therapeutics (and Okapi in 1Q14). We believe that both acquisitions cement Aratana as the go-to company for both large and small biopharma companies looking to leverage their discovery pipelines into the veterinary space. In our opinion, recent competitive concerns have created a buying opportunity ahead of some key milestones. Our \$38 price target is based on a blend our DCF analysis (\$37) and relative valuation (\$38) methodologies.

4Q13 results. In the quarter, R&D expenses totaled \$3.1M (vs. \$2.0M y/y), driven primarily by the development of ongoing programs, as well as an increase in the number of products in development as a result of the Vet Therapeutics acquisition. Currently, the company has 26 employees engaged in R&D efforts. G&A expenses totaled \$4.7M in the quarter (vs. \$0.8M y/y), driven primarily by one-time costs from the Vet Therapeutics acquisition as well as costs associated with its IPO and building of the commercial organization. Following the quarter, Aratana acquired Okapi Sciences. The upfront cash payment of \$13.9M was paid upon closing on January 6, 2014. Subsequently, the company was successful in raising net proceeds of ~\$90.5M through a secondary offering of common stock.

FY14 guidance. The company expects to spend \$35-\$40M (with a ~40/60 split between 1H14 and 2H14) in operating expenses, excluding one-time payments for the recent acquisitions and costs from further business development opportunities in 2014. The company anticipates \$25-\$30M of this spending to be on clinical development and post-licensure studies. On the commercial efforts side, the company is planning to begin building its oncology specialty sales force with ~10 people in FY14.

Upcoming milestones:

- AT001 (grapiprant, EP4 antagonist for osteoarthritis pain). Aratana announced positive top-line data from its pivotal dose-ranging study of AT-001 for dogs; it expects to begin a pivotal field effectiveness study with a once-daily dose in Q2. Aratana anticipates first FDA approval in 2016.

FY DEC	2013A	2014E	2015E
Revenue (\$M) 1Q	\$0.0	\$0.0	--
2Q	\$0.0	\$0.0	--
3Q	\$0.0	\$0.0	--
4Q	\$0.0	\$0.0	--
FY	\$0.1	\$0.0	\$0.0
EPS 1Q	(\$0.24)	(\$0.28)	--
2Q	(\$4.62)	(\$0.35)	--
3Q	(\$0.22)	(\$0.38)	--
4Q	\$0.33	(\$0.19)	--
FY	(\$0.39)	(\$1.27)	(\$1.52)
P/E	NM	NM	NM
Previous FY	(\$5.15)	(\$0.72)	(\$0.65)

Source: Company reports and JMP Securities LLC

STOCK PRICE PERFORMANCE



- AT-002 (capromorelin, ghrelin agonist for appetite stimulation). Aratana has initiated its pivotal field effectiveness trial in client-owned dogs and is actively enrolling patients. The company anticipates first FDA approval in 2016.
- AT-003 (bupivacaine liposome injection suspension for post-surgical pain). Aratana presented the AT-003 development program to the FDA's Center for Veterinary Medicine (CVM) and announced plans to initiate a pilot field study in client-owned dogs in the second quarter of 2014. Aratana anticipates first FDA approval in 2016.
- AT-004 B-cell Lymphoma (canine lymphoma monoclonal antibody against CD-20). AT-004 has received a conditional license from the USDA, and in 2013, Aratana submitted what it believes to be data to support a full licensure, which it expects in the next 9-12 months.
- AT-005: T-cell Lymphoma (canine lymphoma monoclonal antibody against CD-52). In January 2014, Aratana announced it had received a conditional license from the USDA for the T-cell lymphoma product. During 2014, Aratana expects to complete the studies it believes will support full licensure in 2015. In addition, Aratana will initiate other studies to help develop the lymphoma market.
- AT-006: Feline Herpes Virus (ciprovir, antiviral for feline ocular herpes infection). As part of the Okapi Sciences acquisition, Aratana acquired AT-006. This antiviral product is currently in a pivotal field study in Europe and Aratana expects to file for EU regulatory review in 2014. Shortly after the Okapi Sciences acquisition, Aratana filed an Investigational New Animal Drug (INAD) application with the CVM and is preparing to meet with the FDA to present the development plan.
- AT-007: Feline Immunodeficiency Virus. AT-007 was also part of the Okapi Sciences acquisition and is currently in a pilot trial in Europe. Shortly after the end of the fourth quarter, Aratana filed an INAD with the CVM and is preparing to meet with the FDA to present a development plan.
- Option Programs: During 2013, Aratana entered into option agreements relating to three molecules. As disclosed recently, two of these molecules were in the same therapeutic class and after performing an analysis of both compounds, Aratana has decided to continue option diligence on only one of these molecules. For each of the remaining two options, we anticipate making opt-in/opt-out decisions in the first half of 2014.

Company Description

Founded in 2010, Aratana is a development-stage biopharmaceutical company focused on the licensing, development, and commercialization of prescription medications for companion animals (i.e., pet therapeutics). The companion animal market represents a sizable opportunity with a number of therapeutic and medical needs that have yet to be fully realized or met. Aratana has an active in-licensing effort focused on identifying human therapeutics for development and commercialization as pet therapeutics. This model enables human health-focused pharma and biotech companies to extend drug candidates to the companion animal market. With a focus on both cats and dogs, a single, in-licensed drug candidate can offer two therapeutic programs, each of which can potentially offer its own arrangement with specific development milestones and royalties. Additionally, Aratana is developing its own commercial operations to potentially bring its current and future in-licensed drugs to market.

Investment Risks

Limited operating history and significant losses. The company is a development-stage company with a limited operating history and significant losses since its inception. Aratana is expected to continue to incur losses in the short- to medium-term, as it continues the development of product candidates. Previous losses, combined with expected future losses, will continue to have an adverse effect on stockholders' equity and working capital.

Dependence on the success of the three compounds currently in development. Aratana currently has no products approved for commercial distribution. To date, the company has invested much of its efforts and financial resources in the in-licensing, research, and development of AT-001, AT-002, and AT-003, currently the only product candidates that are still in development. If Aratana is not successful in commercializing one or more product candidates, operating results will be negatively impacted.

Regulatory environment. The denial or delay of regulatory approval (e.g., FDA, EMA) for Aratana's existing and future product candidates would delay commercialization efforts and adversely impact the potential to generate revenue and operating results.

Market acceptance/commercial success. Even if current or future product candidates obtain regulatory approval, they may fail to achieve market acceptance and commercial success, which would adversely affect the company's operating and financial results.

Financing risk. On June 27, 2013, Aratana completed an initial public offering, issuing 5.8 million shares of common stock at a price of \$6.00/share, resulting in net proceeds of \$35 million. The company plans to use the net proceeds of the offering to: (i) in-license and develop additional product candidates; (ii) commercialize its current and future product candidates; (iii) establish a direct sales organization in the U.S.; and (iv) for general corporate and working capital purposes. Cash on hand should be enough to fund clinical efforts for AT-001, AT-002, and AT-003 to completion. However, the company will need to raise additional capital in order to successfully commercialize these products and expand its product pipeline.

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

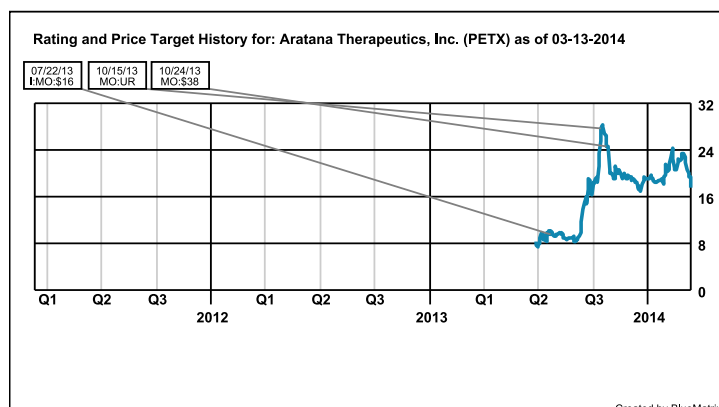
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JMP Rating	Regulatory Equivalent	# Co's Under Coverage	% of Total	Regulatory Equivalent	# Co's Under Coverage	% of Total	# Co's Receiving IB Services in Past 12 Months	% of Co's With This Rating
MARKET OUTPERFORM	Buy	248	56.88%	Buy	248	56.88%	99	39.92%
MARKET PERFORM	Hold	138	31.65%	Hold	138	31.65%	17	12.32%
MARKET UNDERPERFORM	Sell	7	1.61%	Sell	7	1.61%	0	0%
COVERAGE IN TRANSITION		43	9.86%		43	9.86%	0	0%
TOTAL:		436	100%		436	100%	116	26.61%

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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