

Pfenex Inc.

Second-Quarter Earnings Highlighted by Active Clinical Pipeline and Business Update

After the markets closed on Friday, August 29, Pfenex reported its first quarterly results since its initial public offering and provided an update on clinical programs. While the financial results were uneventful, the company reiterated the clinical development timeline, highlighted by a Phase III initiation and several new compounds entering clinical testing within the next 12-18 months. Exhibit 1 provides a detailed variance analysis of the company's operating results compared with our financial estimates.

- Pfenex's lead product candidate, PF582, is a biosimilar form of the \$4.3 billion drug Lucentis and has an expanding development pipeline. PF582 is the wholly owned, lead compound at Pfenex. We expect Phase Ib/IIa data later this year and project the company to advance the product candidate into a registration-enabling trial in 2015. We estimate that by 2018, roughly \$530 million in current sales volume of Lucentis will come off patent, and by 2020, we predict that figure will increase by an additional \$2.0 billion with patent expiry in the United States. In our view, the development timelines of the Phase III trial align nicely with the patent expiration schedule for Lucentis, which we believe is an attractive reference drug given the sizable market it addresses and challenging manufacturing profile.
- Behind PF582 is a portfolio of wholly owned and partnered development initiatives with three compounds entering clinical testing this year. We view the coming 12-18 months as extremely formative for Pfenex, with the potential for clinical data from multiple products and additional corporate and government collaborations, which could infuse nondilutive capital into the company as well as material increases in the level of awareness of the biosimilar commercial opportunity. We highlight PF530, a biosimilar form of Betaseron for the treatment of multiple sclerosis, which is expected to advance into clinical studies later this year, with Phase I data expected in the first half of 2015. We believe the activity surrounding the company's portfolio will present significant catalysts over the next two years, which we summarize in exhibit 3.
- Pfenex has roughly \$50 million in capital, which we believe will sustain the company for at least 18-24 months, based on our current burn rate estimates. With an estimated enterprise value of about \$75 million and numerous upcoming catalysts that stand to increase value, we believe that current price levels represent an attractive entry point for investors. We therefore maintain our Outperform rating on Pfenex shares.

Please see the following pages for more details.

Pfenex is a San Diego-based biotechnology company focused on biosimilars and difficult-to-manufacture protein-based therapeutics. The lead product candidate, PF582, a biosimilar to Lucentis (ranibizumab), is in Phase I/II study with data expected later this year.

John Sonnier +1 312 364 8224 jsonnier@williamblair.com

Andy T. Hsieh, Ph.D. +1 312 364 5051 ahsieh@williamblair.com September 03, 2014

Stock Rating: **Outperform**Company Profile: **Aggressive Growth**

Symbol: PFNX (NYSE)
Price: \$6.94 (52-Wk.: \$5-\$7)
Market Value (mil.): \$137
Fiscal Year End: December
Long-Term EPS Growth Rate: NA
Dividend/Yield: None

2013A	2014E	2015E
-\$3.76	-\$3.41	-\$1.01
	-\$3.41	-\$1.01
NM	NM	NM
	NM	NM
	-\$3.76	-\$3.76 -\$3.41 -\$3.41

Trading Data (FactSet)	
Shares Outstanding (mil.)	2
Float (mil.)	9
Average Daily Volume	134,590

Financial Data (FactSet)	
Long-Term Debt/Total Capital (MRQ)	0.0
Book Value Per Share (MRQ)	-62.9
Return on Equity (TTM)	-19.6

Two-Year Price Performance Chart



Sources: FactSet and William Blair estimates

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Portfolio Manager Summary

After the markets closed on Friday, August 29, Pfenex reported second-quarter financial results and provided an update on clinical programs. While the financial results were uneventful, the company reiterated the clinical development timeline, highlighted by a Phase III initiation and several new compounds entering clinical testing within the next 12-18 months. Exhibit 1 provides a detailed variance analysis of the company's operating results compared with our financial estimates. We updated our financial projections to come more in line with operating trends (exhibit 2). In addition, we expect an active news calendar for Pfenex in the next two years, which we highlight in exhibit 3.

Exhibit 1
Pfenex Inc.
Second Quarter 2014 Variance Analysis

(dollars in millions except EPS)

	PFNX Q2 2014A	WB Q2 2014E
Total Revenues	\$3	\$2
COGS	\$3	\$2
SG&A	\$2	\$2
R&D	\$1	\$1
Net Loss	(\$2)	(\$3)
EPS	(\$1.67)	(\$1.90)

Sources: Pfenex reports and William Blair & Company, L.L.C. estimates.

Exhibit 2
Pfnex Inc.
Guidance and Estimates
(dollars in millions except EPS)

	WB Previous 2014E	WB Revised 2014E	WB Previous 2015E	WB Revised 2015E	WB Previous 2016E	WB Revised 2016E
Total Revenues	\$8	\$9	\$8	\$8	\$8	\$8
COGS	\$7	\$8	\$8	\$8	\$9	\$9
SG&A	\$8	\$9	\$8	\$8	\$9	\$9
R&D	\$7	\$6	\$14	\$14	\$18	\$18
Net Loss	(\$13)	(\$13)	(\$20)	(\$21)	(\$26)	(\$28)
EPS	(\$3.31)	(\$3.41)	(\$0.93)	(\$1.01)	(\$0.94)	(\$1.01)

Sources: Pfenex reports and William Blair & Company, L.L.C. estimates.

Exhibit 3 Pfenex Inc. Timeline

Date	Product	Event
2014	PF530 (biosimilar Betaseron)	Phase I trial initiation in relapsing forms of MS (2H).
	PF582 (biosimilar Lucentis)	Phase lb/IIa trial completion in wet age-related macular degeneration (4Q).
	Recombinant Anthrax Vaccine	Phase la trial initiation in anthrax (2H).
	Recombinant Malaria Vaccine	Phase I trial initiation in malaria (2H).
2015	PF530 (biosimilar Betaseron)	Phase I trial results in relapsing forms of multiple sclerosis (1H).
	PF582 (biosimilar Lucentis)	Phase III equivalance trial in wet age-related macular degeneration (3Q).
	PF708 (generic Forteo)	ANDA-enabling pharmacokinetic bioequivalence trial initiation in osteoporosis (2H).
	Recombinant Malaria Vaccine	Phase I trial results in malaria.
2016	PF688 (biosimilar Cimzia)	Phase I trial initiation in Crohn's disease and rheumatoid arthritis.
	Recombinant Anthrax Vaccine	Phase la trial results in anthrax (3Q).
2017	PF582 (biosimilar Lucentis)	Phase III trial results in age-related macular degeneration.

ANDA = abbreviated new drug application.

Sources: Pfenex reports.

Valuation

Pfenex is trading at \$6.94 with a market cap of \$141 million, which we believe represents an attractive value at current price levels. Given the breadth of the clinical pipeline and an active news calendar in the next two years, we see significant room for upside potential in Pfenex shares.

Risks

While we view Pfenex as a well-capitalized company with a modest clinical risk profile, numerous risks remain given that the company's product candidates are biosimilars of commercialized drugs. The major clinical risk is the emergence of unexpected adverse events from compounds generated from Pfenex's proprietary manufacturing platform. The major capital risk is that additional infusions of funding are needed before the company can reach profitability, which might include further equity fundraising. The major regulatory risk is that PF582 could face scrutiny before approval. However, given the comprehensive bioanalytical testing completed by Pfenex, we believe the asset is de-risked, and we remain optimistic about the compound's path forward.

Exhibit 4 Pfenex Inc. Income Statement

(dollars in thousands except EPS and shares in thousands)

2013A Q4E 2014E 2018E 2019E 2021E Q1A Q2A Q3E 2015E 2016E 2017E 2020E Revenue 11,914 2,558 3,266 1,800 1,850 9,474 8,200 8,200 8,200 8,200 8,200 9,150 9,550 PF582 0 6,327 46,812 140,838 373,953 0 0 0 0 0 0 0 0 Total revenues \$11,914 \$2,558 \$3,266 \$1,850 \$9,474 \$8,200 \$8,200 \$8,200 \$14,527 \$55,012 \$149,988 \$383,503 \$1,800 1,908 2,544 1,500 1,550 10,000 10,000 25,285 220,000 Cost of revenue 6,423 7,502 8,000 9,000 77,537 Gross profit 5,491 650 722 300 300 1,972 200 (800)(1,800)4,527 29,727 72,451 163,503 SG&A 1,495 2,024 3,050 2,150 8,719 8,000 14,500 20,300 41,000 6,698 9,000 10,000 48,000 R&D 678 2,350 6,338 13,500 18,550 19,650 5,490 860 2,450 17,800 18,135 20,575 21,100 12,188 2,173 2,884 28,135 33,050 39,950 61,575 Total operating expenses 5,400 4,600 15,057 21,500 26,800 69,100 Loss from operations (\$6,697) (\$1,523) (\$2,162) (\$5,100)(\$4,300) (\$13,085) (\$21,300) (\$27,600) (\$29,935) (\$28,523) (\$10,223) \$10,876 \$94,403 (24.0)Other expense, net (36.0)(18.0)(21.0)(8.5)(8.5)(56.0)(32.0)(30.0)(28.0)(26.0)(22.0)(20.0)Net loss before income taxes (\$1,541)(27,630)(28,549)(10,247)10,854 94,383 (6,733)(\$2,183)(\$5,109)(\$4,309)(13,141)(21,332)(29,963)2,671 Income tax benefit (1) 0 0 0 (1) 0 0 0 0 0 (2,538)(14,157)Net loss (\$4,062) (\$29,963)(\$10,247) \$80,226 (\$1,542)(\$2,183)(\$5,109)(\$4,309)(\$13,142)(\$21,332)(\$27,630)(\$28,549)\$8,315 Net loss attributable to common stockholders (5,757) (1,977) (2,630) (5,109) (4,309) (14,024) (21,332) (27,630) (29,963) (28,549) (10,247) 8,315 80,226 Net loss per common share basic and diluted \$0.25 \$2.40 (\$3.76) (\$1.28) (\$1.67) (\$0.25) (\$0.21) (\$3.41) (\$1.01) (\$1.01) (\$1.05) (\$0.96) (\$0.33) Weighted-average common shares basic and diluted 21.217 27.432 28.529 29,670 30,856 33,373 1,531 1,548 1,571 20,500 20,702 11,080 32,090

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Please consult the last page of this report for all disclosures.

PFNX

Sources: Roche and Pfenex reports.

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William Blair was a manager or co-manager of a public offering of equity securities for Pfenex Inc. within the prior 12 months.

William Blair is a market maker in the security of Pfenex Inc. and may have a long or short position.

William Blair intends to seek investment banking compensation in the next three months from Pfenex Inc.

Within the past 12 months William Blair has provided or is providing investment banking services to or has an investment services relationship with Pfenex Inc.

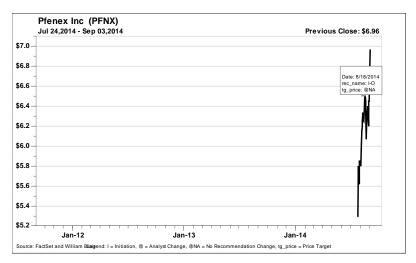
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DOW JONES: 17,067.56 S&P 500: 2,002.28 NASDAQ: 4,598.19



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