

RBC Capital Markets

January 28, 2014

Aerie Pharmaceuticals, Inc.

Channel checks indicate no reason for recent weakness

Our view: Weakness could be a buying opportunity as PG324 Phase IIb results are key near-term (mid-2014) and AR-13324 Phase III results are key longer-term (mid-2014).

Key points:

Tuesday's weakness resulted in several queries regarding AERI's drivers and expectations. Our checks indicate no fundamental reason for the stock to be down. Planning and timing for Phase IIb PG324 and Phase III AR-13324 studies is on track, results from the Phase IIb PG324 study are expected around mid-2014 and the Phase III AR-13324 study around mid-2015, balance sheet is strong with sufficient cash into 2Q:16 (well beyond the two catalysts). IMS prescription data underscores the opportunity for AERI and provides additional validation for its novel combination by showing both growth in the US glaucoma market, for both 1st and 2nd-line drugs, and Novartis' Simbrinza, a recently launched combination drug. We are buyers at current levels and especially on any pullbacks.

- Phase II PG324 study reads out mid-2014; provides visibility on blockbuster opportunity. AERI will evaluate PG324 (combination AR-13324 and latanoprost) in 300 patients and data from the 28 day study could be available around mid-2014/3Q:14. The primary efficacy endpoint is superiority of PG324 to AR-13324 and latanoprost.
- Phase III AR-13324 study could have data in mid-2015; another large drug by itself. Next up are interim 3-month data from animal ocular toxicity studies and end of Phase II meeting. Two Phase IIIs are expected around mid-2014 with 1,200 glaucoma patients in total (baseline IOPs: 21-26 mmHg) in the US with data around mid-2015. The primary efficacy endpoint is non-inferiority to timolol.
- Glaucoma prescriptions continue to increase. 1st-line glaucoma prescriptions totaled ~16M in 2013, up from 15.4M in 2012 (+3.9% Y/ Y). 2nd-line prescriptions totaled ~14.7M, up from 14.6M (+1.2% Y/Y). For second line drugs, combinations increased +4.1% Y/Y and carbonic anhydrase inhibitors increased +3.1% while beta blocker and alpha agonist prescriptions declined Y/Y.
- Novartis' Simbrinza growing albeit slowly. Simbrinza is a combination of Azopt (brinzolamide), a carbonic anhydrase inhibitor, and brimonidine (alpha agonist). Therefore, while efficacy looks solid, it includes all the side effect and inconvenience baggage associated with these 2nd line drug classes. Despite this prescriptions are increasing and during its analyst day update NVS highlighted Simbrinza and combination therapy as a growth opportunity in Glaucoma. We believe AERI's drugs with their novel, safer, differentiated and more convenience mechanisms and administration should do even better upon launch.

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Outperform

Speculative Risk

NASDAQ: AERI; USD 15.76

Price Target USD 25.00

WHAT'S INSIDE	
☐ Rating/Risk Change	☐ Price Target Change
☐ In-Depth Report	☐ Est. Change
Preview	✓ News Analysis

Scenario Analysis*

4	Downside Scenario	Current Price	Price Target	Upside Scenario	
	8.00	15.76	25.00	35.00	—
	↓ 49%		↑ 59%	† 122%	

*Implied Total Returns

Key Statistics

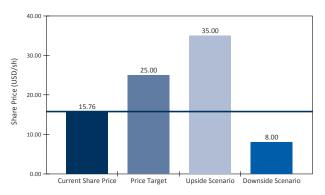
•			
Shares O/S (MM):	23.2	Market Cap (MM):	366
Dividend:	0.00	Yield:	0.0%

RBC Estimates

1			
2013E	2014E	2015E	
0.0	0.0	0.0	
(1.51)	(1.01)	(1.06)	
NM	NM	NM	
Q1	Q2	Q3	Q4
0.0A	0.0A	0.0A	0.0E
0.0E	0.0E	0.0E	0.0E
(0.41)A	(0.28)A	(10.81)A	(0.22)E
(0.24)E	(0.25)E	(0.26)E	(0.26)E
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	0.0 (1.51) NM Q1 0.0A 0.0E (0.41)A (0.24)E	2013E 2014E 0.0 0.0 (1.51) (1.01) NM NM Q1 Q2 0.0A 0.0A 0.0E (0.41)A (0.28)A	2013E 2014E 2015E 0.0 0.0 0.0 (1.51) (1.01) (1.06) NM NM NM Q1 Q2 Q3 0.0A 0.0A 0.0A 0.0E 0.0E 0.0E (0.41)A (0.28)A (10.81)A (0.24)E (0.25)E (0.26)E

Target/Upside/Downside Scenarios

Exhibit 1: Aerie Pharmaceuticals, Inc.



Source: RBC Capital Markets estimates

Target price/ base case

We value AERI at \$25 per share, which includes US and EU sales of AR-13324 and PG324. We assign a 65% probability of success (previously 60%) and a value of ~\$17 per share to the US and \$7 per share to the EU opportunity. We assume a US launch in 2017 and an EU launch in 2018. We forecast peak PG324 sales of \$800M and \$600M and AR-13324 sales of \$300 and \$200M the US and EU, respectively. Finally, we assume patent protection through 2030 and include a terminal value based on a discount rate of 15% and a growth rate of -50%.

Upside scenario

Our upside scenario includes ~\$25 per share in value for the US opportunity and ~\$10 per share in value for the EU opportunity. We forecast peak PG324 sales of \$1.2–1.3B in the US and \$900M–\$1B in the EU and AR-13324 sales of \$200–300MM in the US and \$200–300MM in the EU. We assign products in the pipeline a 60% probability of success, a discount rate of 15%, and a terminal growth rate of -50%.

Downside scenario

Our downside scenario assumes that PG324 will not be approved in the US or EU. We value the US opportunity for AR-13324 at ~\$5 per share and the EU opportunity at ~\$3 per share. We assume market share ramps up to roughly 15% of total second-line glaucoma prescriptions in the US and 10% in the EU. Under such a scenario, peak sales are forecast to be \$400–500M in the US and \$300–400MM in the EU. We assign AR-13324 a 60% probability of success, a discount rate of 15%, and a terminal growth rate of -50%.

Investment summary

We believe AERI shares offer the potential for significant upside as both products in development, AR-13324 and PG324, use a new mechanism of action for the treatment of glaucoma, a blockbuster potential market. AR-13324 will enter Phase III trials based on positive Phase IIb data and PG324 a Phase IIb study based on promising preclinical data in 2014. Results from these and additional studies are expected 2014–2016. Millions of patients worldwide suffer from glaucoma, most need multiple medications, and we forecast peak sales of AERI's products at ~\$1B.

AERI owns 100% of the rights to AR-13324 and PG324 worldwide and patent protection extends into 2030, which means the company is free to partner or be acquired. Given that ophthalmology remains an attractive therapeutic area and AERI's product candidates could have a convenient, one drop once per day efficacy and safety profile, progress through clinical and regulatory milestones, as well as a partnership, could all be upside catalysts.

Potential Catalysts for AERI Shares

- Phase IIb data for PG324 in 2014. Important catalyst as it could show differentiation in efficacy vs. latanoprost, the current market leader.
- Phase III data for AR-13324 in 2015. Important catalyst as positive data could lead to an NDA and MAA filing.
- Phase III data for PG324 in 2016. Key catalyst as clean safety and efficacy beyond latanoprost could make PG324 the firstline drug of choice.
- Potential partnership for AR-13324 and PG324. AERI owns worldwide rights to both product candidates and a partnership is likely after Phase III data.
- Potential approvals and launches in 2017 in the US and 2018 in the EU following regulatory filings in 2016.

Risks to Our Investment Thesis

- Pivotal Phase III and earlier-stage studies could fail.
 AR-13324 must show non-inferiority to a comparator over a longer period and PG324 must show a benefit in patients, which raises risk of failure.
- PG324 Phase IIb study could fail. Our assumption for success is based on pre-clinical data with PG324, and testing it in patients increases risk.
- AERI could fail to find a partner for AR-13324 and PG324 outside the US.
- Sales ramp of AR-13324 and PG324 could lag expectations
 as clinicians fail to take up AERI's drugs, payers put up
 hurdles for reimbursing branded drugs, and cheaper generic
 drugs with other mechanisms hamper market penetration.



Valuation

We value AERI at \$25 per share, which includes US and EU sales of AR-13324 and PG324. We assign a 65% probability of success and a value of ~\$17 per share to the US and \$7 per share to the EU opportunity. We assume a US launch in 2017 and an EU launch in 2018. We forecast peak PG324 sales of \$800M and \$600M and AR-13324 sales of \$300 and \$200M the US and EU, respectively. Finally, we assume patent protection through 2030 and include a terminal value based on a discount rate of 15% and a growth rate of -50%.

Price target impediments

Our price target is dependent solely on the clinical, regulatory and commercial success of AR-13324 and PG324. A Phase IIb study for PG324 and a Phase III study for AR-13324 are expected in 2014 and failure to demonstrate efficacy or safety in one or both of these studies would be a significant setback. Furthermore, any setbacks in regulatory approvals in the US or EU, delay in launch, failure to secure a partnership outside the US for AR-13324 and PG324, increased competition or other limitations to the market potential of these products either due to better efficacy and/or safety outcomes or pricing pressure due to the availability of generic drugs for glaucoma, could negatively impact our valuation.

Company description

Aerie Pharmaceuticals, Inc. is a biotechnology company targeting ophthalmic disorders specifically glaucoma, which is a blockbuster potential markets. Sales of products targeting glaucoma totaled \$4.5B globally and more than 30 million prescriptions for glaucoma drugs were written in the US alone. AERI's drug candidates work by inhibiting rho-kinase and the norepinephrine transporter, a new mechanism of action, something not seen for glaucoma in nearly two decades. AR-13324 could enter Phase III trials in 2014 and PG324 could enter Phase II trials in 2014 and Phase III trials in 2015. These drugs could be used as stand-alone agents for first- or second-line therapy or combined with existing agents.



Adnan Butt (415) 633-8588 Aerie Pharmaceuticals - Income Statement FYE December 31 (in MM: except per share REVENUES AR-13324 6.5 69.1 147.3 235.5 251.0 223.0 190.2 PG324 308.4 452.0 31.8 101.8 180.8 Product Sales 6.5 76.5 179.1 337.2 431.8 531.4 642.2 Rovalties 10.0 24.8 44.6 54.4 63.9 73.6 Total Revenues 6.5 86.6 203.9 381.8 486.2 595.2 715.8 **EXPENSES** COGS 17.9 33.7 43.2 53.1 64.2 R&D 3.2 3.2 2.4 3.4 12.1 3.8 4.0 4.3 4.5 16.5 20.0 22.5 20.0 10.0 5.0 5.0 5.0 5.0 5.0 SG&A 1.7 1.7 3.3 1.8 8.5 1.9 1.9 2.0 1.8 7.5 8.5 15.0 30.0 37.5 45.0 67.4 86.4 106.3 128.4 Other Total Expenses 1 Q 4.9 5.7 5.2 20.6 5.6 5.9 6.2 6.3 24.0 28.5 37.5 50.6 55.2 67.9 106.2 134.5 164.4 197.6 (4.9) (4.9) (5.7) (20.6) (5.9) (6.2)(6.3 (24.0) (28.5 (44.2) 31.4 136.0 275.6 351.7 430.8 518.1 Operating Income (Expense) OTHER Interest income 0.1 0.1 0.1 0.5 (0.2)(0.2)(0.1 (0.1)(0.1)(0.1)(0.2)(0.2)(0.2) (0.2)(0.2)(0.2 (0.2)(0.2)(0.2)Interest expense (0.4) (0.1)(0.2)Other (5.1)Total Other Income (Expense) (5.1)(23.8) (43.9 Income before Tax (5.1)(5.1)(10.9)(5.1)(20.9 (5.6)(5.9)(6.2)(6.3)(28.3 (37.2)31.7 136.3 275.9 352.0 431.1 518.4 10.8 46.3 119.7 146.6 176.3 Taxes Net income (loss) (5.1) (5.1) (10.9) (20.9 (5.6) (5.9) (6.2) (6.3 (23.8) (28.3 (37.2) (43.9 20.9 89.9 182.1 232.3 284.5 342.2 (5.1 EPS, Basic (GAAP) (\$0.41) (\$0.28) (\$10.81) (\$0.22) (\$1.51 (\$0.24) (\$0.25) (\$0.26) (\$0.26 (\$1.01) (\$1.06 (\$1.24) (\$1.44) \$0.67 \$2.84 \$5.63 \$7.04 \$8.45 \$9.97 EPS, Diluted (GAAP) (\$1.13) (\$0.17) (\$0.18) (\$0.19)(\$0.74) (\$0.81 (\$0.97) (\$1.13) \$2.24 \$4.46 \$6.75 \$7.99 (\$0.24)(\$0.19) (\$0.16 (\$0.94 (\$0.19 \$0.53 \$5.60 Shares outstanding, Basic 12.5 18.4 1.0 23.3 13.8 23.4 23.6 23.7 23.8 23.6 26.6 29.9 30.5 31.1 31.7 32.4 33.0 33.7 34.3 Shares outstanding, Diluted 26.9 9.6 31.8 22.3 31.9 32.1 32.2 32.3 32.1 35.1 38.4 39.0 39.6 40.2 40.9 41.5 42.2 42.8 2Q:13A 2013E 2Q:14E 2018 2023F Operating Ratio 2017 2021 10.0% 10.09 10.0% 10.0% 10.0% 10.0% 10.0% Gross Margin NA NA NA 90.0% 90.0% 90.0% 90.0% 90.0% 90.09 NA 90.09 R&D NA NA NA NA NA NA NA 308.7% 11.6% 2.5% 1.3% 1.0% 0.8% 0.7% SG&A NA NA NA NA NA NA NA 463.0% 43.3% 22.1% 17.7% 17.8% 17.9% 17.9% Operating Margin NA NΔ NA NΔ ΝΔ NA NA NA NΔ ΝΔ -681 7% 36.3% 66.7% 72 29 72.3% 72 4% 72 4% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% 34.0% -677.1% 24.2% 44.1% 47.7% 47.8% 47.8% 47.8% NA NA NA NA NA Net Margin NA NA NA Source: Company reports and RBC Capital Markets estimates **Balance Sheet - Select Items** 2Q:13A 2013E 2Q:14E 4Q:14E 2023E Cash and cash equivalents 2.4 4.6 69.5 69.5 80.1 75.8 71.2 66.5 66.5 134.4 101.4 61.7 75.0 140.1 280.8 492.4 755.0 1,072.1 Prepaid expenses and other current assets 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 71 9 71 9 82.5 140.8 110 3 100.8 197 5 386 3 626.4 918 8 1 269 0 35 7 1 78.2 73.6 68.9 68.9 73.2 Total current assets Property, plant and equipment, net 0.1 0.1 0.1 0.1 0.1 0.1 0.0 0.0 0.0 (0.1 (0.1) (0.2) (0.3) (0.4)(0.4) (0.5) (0.6 (0.7) 3.7 7.2 72.1 72.1 82.6 78.3 73.7 69.0 69.0 140.8 110.2 73.0 100.5 197.1 385.8 625.8 918.2 1,268.3 Total assets Current Liabilities **Total current liabilities** 11.8 18.1 18.1 18.1 18.1 18.6 20.1 22.2 11.5 **Total liabilities** 4.6 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 11.5 Share Capita 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 163.5 163.5 Share Premium 0.1 1.1 69.5 69.5 69.5 69.5 69.5 69.5 69.5 163.5 163.5 163.5 163.5 163.5 163.5 163.5 Accumulated deficit (74.0) (84.8)(88.4 (88.4 (92.4)(96.7)(101.3)(106.0 (106.0) (128.2 (159.3 (197.0 (170.0 (73.9) 114.3 352.7 643.4 991.7 Total stockholders' equity 42.5 42.5 38.5 34.1 24.8 24.8 96.6 65.5 27.8 54.9 150.9 339.2 577.6 868.2 1,216.5 (12.7)(22.4)29.5 Total liabilities and stockholders Equity 72.1 68.1 59.1 54.4 54.4 126.2 95.6 58.4 86.0 182.5 371.2 611.3 903.6 3.7 7.2 72.1 63.7 1.253.7 Cash Flow Statement - Select Item 2013E 20.9 89.9 232.3 284.5 342.2 Net Income (loss) (5.1)(20.9)(5.1 (20.9) (5.6)(5.9)(6.2)(6.3 (23.8)(28.3 (37.2) (43.9) 182.1 Depreciation and amortization 0.0 0.0 0.0 0.1 0.0 0.0 0.0 0.0 0.2 0.2 0.2 0.2 0.2 0.2 0.2 0.2 0.2 0.2 6.1 6.1 Stock based compensation 6.1 6.1 6.1 6.1 0.4 1.5 Net cash provided (used) by operating activities (5.1) (2.5) (11.6) (3.5) (7.5 (4.0) (4.3) (4.6) (4.7 (17.5) (26.0 (32.9) (39.6 13.4 65.2 140.8 211.7 262.7 317 2 Purchase of property and equipment and intangible assets (0.0)(0.0 (0.0)(0.0 (0.1 (0.1 (0.1 (0.1)(0.1) (0.1) Net cash used in investing activities (0.0) (0.0) (0.1)(0.1 (0.1) (0.1)(0.1 (0.1)(0.1 (0.1)(0.1 (0.1)(0.0)(0.0)(0.0 (0.1) (0.0)(0.0)Proceeds from issuances 68.4 68.4 94.0 7.0 13.3 68.4 88.7 Net cash provided by (used in) financing activities 94.0 Decrease in cash and cash equivalents (5.1)4.5 1.7 64.8 81.2 (4.0)(4.3)(4.6)(4.7 (17.6) 67.9 (33.0) (39.7 13.3 65.1 140.7 211.6 262.6 317.1 Cash and cash equivalents at the beginning of the year 2.9 (2.1)2.9 2.9 84.1 80.1 75.8 71.2 66.5 134.4 101.4 61.7 75.0 280.8 755.0

Source: Company reports and RBC Capital Markets estimates.

Cash and cash equivalents at the end of the year

75.8

66.5

71 2

66.5

1344

101 4

61.7

75.0

140 1

280.8

492.4

1.072.1

755.0

69 5

46

24

(2.1)

84 1

80.1



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Top Pick (TP): Represents analyst's best idea in the sector; expected to provide significant absolute total return over 12 months with a favorable risk-reward ratio.

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Underperform (U): Returns expected to be materially below sector average over 12 months.

Risk Rating

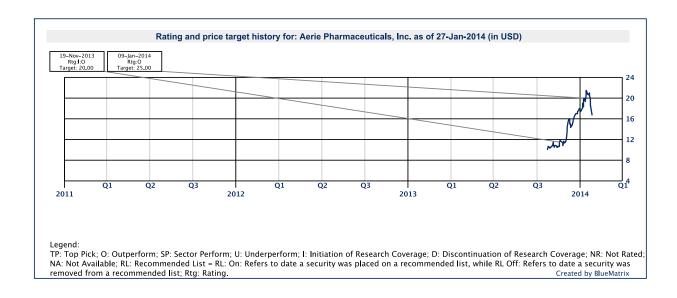
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	Distribution	of ratings		
	RBC Capital Market	s, Equity Research		
	As of 31-E	Dec-2013		
			Investment Banking	
			Serv./Past 12 Mo	os.
Rating	Count	Percent	Count	Percent
BUY [Top Pick & Outperform]	791	51.13	274	34.64
HOLD [Sector Perform]	666	43.05	179	26.88
SELL [Underperform]	90	5.82	14	15.56



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