

Zacks Small-Cap Research

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Elizabeth Senko, CFA

312-265-9484

esenko@zacks.com

scr.zacks.com

10 S. Riverside Plaza, Chicago, IL 60606

EyeGate Pharmaceuticals (EYEG-NASDAQ)

Moving towards approval

We are adjusting our valuation target to reflect progress in EYEG's PRK and dry eye pipeline and the announcement of Moxigel for bacterial conjunctivitis. Our valuation goes from \$12.00 to \$15.00 based on a 10-year DCF with terminal market shares of 15-20% in the US, 20% royalties, 10% discount rate, 20% terminal EBIT margin, 25% tax rate and \$6 million in net cash.

OUTLOOK

EyeGate Pharmaceuticals is a late-stage clinical company developing better solutions to current treatments for post-surgical eye care and moderate to severe dry eye using OBG, its proprietary cross-linked formulation of sodium hyaluronate (HA). The Company is preparing its IND filing for OBG as an alternative to a bandage contact lens following PRK surgery. EYEG is completing a follow-on pilot study for OBG in punctate epitheliopathies (PE), a condition arising from dry eye and other forms of eye irritation.

Current Price (09/25/20) **\$3.80**
Valuation \$15.00

SUMMARY DATA

52-Week High **\$12.20**
 52-Week Low **\$2.91**
 One-Year Return (%) **14.06**
 Beta **1.28**
 Average Daily Volume (sh) **26,268**

Shares Outstanding (mil) **5**
 Market Capitalization (\$mil) **\$17**
 Short Interest Ratio (days) **N/A**
 Institutional Ownership (%) **42**
 Insider Ownership (%) **57**

Annual Cash Dividend **\$0.00**
 Dividend Yield (%) **0.00**

5-Yr. Historical Growth Rates
 Sales (%) **N/A**
 Earnings Per Share (%) **N/A**
 Dividend (%) **N/A**

P/E using TTM EPS **N/A**
 P/E using 2020 Estimate **-2.3**
 P/E using 2021 Estimate **-2.7**

Zacks Rank **N/A**

Risk Level **High,**
 Type of Stock **Small-Blend**
 Industry **Med-Drugs**
 Zacks Rank in Industry **N/A**

ZACKS ESTIMATES

Revenue

(in millions of \$)

	Q1	Q2	Q3	Q4	Year
	(Mar)	(Jun)	(Sep)	(Dec)	(Dec)
2018	1.1 A	0.2 A	0.3 A	0.1 A	1.7 A
2019	2.7 A	0.0 A	0.0 A	0.0 A	2.7 A
2020					0.1 E
2021					8.0 E

Earnings per share

	Q1	Q2	Q3	Q4	Year
	(Mar)	(Jun)	(Sep)	(Dec)	(Dec)
2018	-\$2.10 A	-\$1.05 A	-\$1.05 A	-\$0.75 A	-\$4.50 A
2019	\$0.15 A	-\$0.60 A	-\$1.15 A	-\$0.76 A	-\$2.23 A
2020	-\$0.43 A	-\$0.38 A			-\$2.67 E
2021					-\$1.36 E

Zacks Projected EPS Growth Rate - Next 5 Years % **N/A**

COMPANY UPDATE

We are raising our target valuation for EyeGate Pharmaceuticals from \$12.00 to \$15.00 based on solid progress in its existing pipeline and the introduction of a new candidate for treating bacterial conjunctivitis.

Since early 2019, the Company focused all its efforts on its OBG (Ocular Bandage Gel) platform technology. The Company has completed four clinical trials for OBG, three for PRK and one for PE (punctate epitheliopathies).

EyeGate filed its IDE for OBG in corneal wound repair post photorefractive keratectomy (PRK) surgery in May 2019 and began its pivotal trial in June that year. In November and December, the Company received top-line data showing superiority in its primary endpoint of superiority in wound healing over standard-of-care a bandage contact lens (BCL). In the study, by day 3, 80% of study eyes receiving OBG were completely healed, compared with 67% in the control group.

In June 2020, EYEG received positive feedback from the FDA on its proposed packaging (a unique multi-dose bottle) and asked for several additional tests on the bottle. The Company expects to complete the tests and file a *de novo* application for OBG in PRK by the end of this year, with a decision in mid-2021.

A month later, EYEG received the go-ahead from the FDA for its pivotal study for using OBG in the treatment of punctate epitheliopathies (PEs) in dry eye. The Company expects to enroll patients in the first half of 2021. Top-line data is anticipated in the second half of 2021 and will be part of the *de novo* filing planned for later that year.

Finally, following a meeting with the FDA in August, EYEG announced plans for an IND for OBG in the treatment of bacterial conjunctivitis. The product, Moxigel, combines EYEG's OBG technology with moxifloxacin, an antibiotic. The IND filing requires a short (28-day) toxicity study and the Company expects to file its IND in the first half of 2021. Once the IND is approved, the Company is expected to proceed with two clinical studies, beginning in the second half of 2021.

Based on the Company's positive progress, we are raising our valuation from \$12.00 to \$15.00 based on a 10-year DCF with terminal market shares of 15-20% in the US for both its PRK and PE products, 20% royalties, 10% discount rate, 20% terminal EBIT margin, 25% tax rate and \$6 million in net cash.

HISTORICAL STOCK PRICE



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