

Eyenovia Reports Third Quarter 2018 Financial Results

NEW YORK, Nov. 13, 2018 (GLOBE NEWSWIRE) -- Eyenovia, Inc. (NASDAQ: EYEN), a biopharmaceutical company developing a pipeline of ophthalmology products utilizing its patented piezo-print technology to deliver microdosed medications topically to the eye, today announced financial results for the third quarter ended September 30, 2018.

Q3 2018 and Recent Business Highlights

- Received investigational new drug (IND) acceptance to enter Phase III with MicroStat for mydriasis
- Results from PG21 study demonstrating a robust IOP lowering effect from a microdosed prostaglandin and high ocular delivery success rate have been accepted in the peer-reviewed journal *Clinical Ophthalmology* and are currently *In Press*
- Submitted Optejet™ as the trademark for the Company's proprietary container closure system

Dr. Sean Ianchulev, Eyenovia's Chief Executive Officer and Chief Medical Officer commented, "With the acceptance of our IND application for MicroStat for mydriasis, we expect to initiate our first Phase III study this month, followed by two additional Phase III studies for myopia progression and chronic angle closer glaucoma over the next nine months. As we seek to further support these upcoming trials, our PG21 study demonstrated superior patient self-administration of 90 percent and similar intra ocular pressure lowering efficacy compared to traditional eyedrops and was recently selected for publication by *Clinical Ophthalmology*. We look forward to continuing to develop and validate our proprietary, high precision microdosing technology platform, and are very pleased to officially brand our technology, the Optejet."

Third Quarter 2018 Financial Review

For the third quarter of 2018, net loss was approximately \$(4.3) million, or \$(0.43) per share, compared to a net loss of approximately \$(0.9) million, or \$(0.10) per share for the third quarter of 2017.

Research and development expenses totaled approximately \$2.5 million for the third quarter of 2018, an increase of 336%, compared to approximately \$0.6 million for the same period in 2017.

For the third quarter of 2018, general and administrative expenses were approximately \$1.8 million, an increase of 482%, compared to approximately \$0.3 million for the third quarter of 2017.

Total operating expenses for the third quarter of 2018 were approximately \$4.3 million, an increase of 388%, compared to total operating expenses of approximately \$0.9 million for the same period in 2017.

As of September 30, 2018, the Company's cash balance was approximately \$21.0 million compared to \$24.6 million at June 30, 2018.

The Company reiterated the timeline for its 12-month key clinical milestones:

- Q4 2018: Initiate MicroStat Phase III trial
- H1 2019: Report MicroStat Phase III trial results
- H1 2019: Initiate MicroPine Phase III trial
- H1 2019: Initiate MicroProst Phase III trial
- H1 2019: MicroTears OTC registration

Conference Call and Webcast

The conference call is scheduled to begin at 8:30 am ET on Tuesday, November 13, 2018. Participants should dial 1-866-916-2921 (United States) or 1-210-874-7771 (International) with the conference code 1797567. A live webcast of the conference call will also be available on the investor relations page of the Company's corporate website at www.eyenoviabio.com.

After the live webcast, the event will be archived on Eyenovia's website for one year. In addition, a telephonic replay of the call will be available until November 20, 2018. The replay can be accessed by dialing 1-855-859-2056 (United States) or 1-404-537-3406 (International) with confirmation code 1797567.

About Eyenovia

Eyenovia, Inc. (NASDAQ: EYEN) is a specialty biopharmaceutical company building a portfolio of next generation topical eye treatments based on its proprietary delivery and formulation platform for microdosing. Eyenovia's pipeline is currently focused on the late-stage development of microdosed medications for myopia progression, glaucoma, mydriasis and other eye diseases.

Forward Looking Statements

Except for historical information, all of the statements, expectations, and assumptions contained in this press release are forward-looking statements. Forward-looking statements include, but are not limited to, statements that express our intentions, beliefs, expectations, strategies, predictions or any other statements relating to our future activities or other future events or conditions. These statements are based on current expectations, estimates and projections about our business based, in part, on assumptions made by management. These statements are not guarantees of future performance and involve risks, uncertainties and assumptions that are difficult to predict. Therefore, actual outcomes and results may, and are likely to, differ materially from what is expressed or forecasted in the forward-looking statements due to numerous factors discussed from time to time in documents which we file with the SEC. In addition, such statements could be affected by risks and uncertainties related to, among other things: risks involved in clinical trials, including, but not limited to, the initiation, timing, progress and results of such trials; the timing and our ability to submit applications for, obtain and maintain regulatory approvals for, our product candidates; our ability to develop and implement commercialization, marketing and manufacturing capabilities and strategies; the potential advantages of our product candidates; our ability to attract and retain key personnel; the rate and degree of market acceptance and clinical utility of our product candidates; our estimates regarding the potential market opportunity for our product candidates; intellectual property risks; our ability to identify additional products, product candidates or technologies with significant commercial potential that are consistent

with our commercial objectives; our expectations regarding our ability to fund our operating expenses and capital expenditure requirements; the impact of government laws and regulations; our competitive position; and general economic conditions. Any forward-looking statements speak only as of the date on which they are made, and except as may be required under applicable securities laws, we do not undertake any obligation to update any forward-looking statements.

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(Financial Statements to Follow)

EYENOVIA, INC.

Condensed Balance Sheets

	September 30, 2018 (unaudited)	December 31, 2017
Assets		
Current Assets:		
Cash	\$ 21,044,135	\$ 5,249,511
Prepaid expenses and other current assets	298,450	37,149
Total Current Assets	21,342,585	5,286,660
Property and equipment, net	11,152	27,960
Deferred offering costs	-	328,700
Security deposit	117,800	-
Total Assets	<u>\$ 21,471,537</u>	<u>\$ 5,643,320</u>

Liabilities and Stockholders' Equity

Current Liabilities:

Accounts payable	\$ 840,230	\$ 246,384
Accrued expenses and other current liabilities	888,984	306,263
Total Current Liabilities	1,729,214	552,647
Deferred rent	2,332	-
Total Liabilities	1,731,546	552,647

Commitments and contingencies

Stockholders' Equity:

Preferred stock, \$0.0001 par value, 6,000,000 shares authorized;

Series A Convertible Preferred Stock, 0 and 20,000,000 shares designated

as of September 30, 2018 and December 31, 2017, respectively,

0 and 2,932,431 shares issued and outstanding

as of September 30, 2018 and December 31, 2017, respectively

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Series A-2 Convertible Preferred Stock, 0 and 5,714,286 shares designated

as of September 30, 2018 and December 31, 2017, respectively,

0 and 788,827 shares issued and outstanding

as of September 30, 2018 and December 31, 2017, respectively

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Series B Convertible Preferred Stock, 0 and 10,000,000 shares designated

as of September 30, 2018 and December 31, 2017, respectively,

0 and 918,983 shares issued and outstanding

as of September 30, 2018 and December 31, 2017, respectively

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Common stock, \$0.0001 par value, 90,000,000 shares authorized;

10,088,996 and 2,566,530 shares issued and outstanding as of September 30, 2018 and December 31, 2017, respectively

1,009 257

Additional paid-in capital

50,070,169 24,351,138

Accumulated deficit

(30,331,187) (19,261,186)

Total Stockholders' Equity

19,739,991 5,090,673

Total Liabilities and Stockholders' Equity

\$ 21,471,537 \$ 5,643,320

EYENOVIA, INC.

Condensed Statements of Operations (unaudited)

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2018	2017	2018	2017
Operating Expenses:				
Research and development	\$ 2,487,573	\$ 570,422	\$ 6,993,832	\$ 2,233,193
General and administrative	1,832,794	314,859	4,079,249	735,759
Total Operating Expenses	4,320,367	885,281	11,073,081	2,968,952
Loss From Operations	(4,320,367)	(885,281)	(11,073,081)	(2,968,952)
Other Income (Expense):				
Interest income (expense)	(964)	665	3,080	1,396
Net Loss	<u>\$ (4,321,331)</u>	<u>\$ (884,616)</u>	<u>\$ (11,070,001)</u>	<u>\$ (2,967,556)</u>
Net Loss Per Share				
- Basic and Diluted	<u>\$ (0.43)</u>	<u>\$ (0.10)</u>	<u>\$ (1.20)</u>	<u>\$ (0.35)</u>
Weighted Average Number of Common Shares Outstanding				
- Basic and Diluted	<u>10,030,296</u>	<u>8,514,906</u>	<u>9,219,818</u>	<u>8,514,906</u>



Source: Eyenovia, Inc.