



## **ObsEva Reports Third Quarter 2017 Financial Results and Provides Business Update**

### **- All Three Development Compounds Progressing with Key Clinical Milestones Over the Next 12 Months -**

**Geneva, Switzerland and Boston, MA – November 14, 2017** - ObsEva SA (NASDAQ: OBSV), a clinical-stage biopharmaceutical company focused on the development and commercialization of novel therapeutics for serious conditions that compromise a woman's reproductive health and pregnancy, today reported financial results for the third quarter ended September 30, 2017 and provided a business update outlining recent corporate progress and upcoming milestones.

*"We are very proud of the progress that we are making in each of our development programs, and continue to deliver upon our objectives and timelines that were outlined at our IPO in January. Our team has been able to navigate the complexities of executing on global trials and the robust patient recruitment in our IMPLANT 2 and EDELWEISS clinical trials demonstrates the severe unmet medical need that exists and demonstrates the potential of our therapeutics to greatly improve patient care."* said Ernest Loumaye, Chief Executive Officer of ObsEva. *"With our recent successful financing, we are now positioned to achieve several clinical milestones in 2018-19."*

#### **Pipeline Progress Achieved in Third Quarter of 2017**

- Patient recruitment was completed in August for the Phase 3 IMPLANT 2 trial of nolasiban, ObsEva's oral oxytocin receptor antagonist designed to improve clinical pregnancy and live birth rates in women undergoing in-vitro fertilization (IVF). Approximately 760 patients have now been randomized in this European trial.
- Patient recruitment was completed in November, 2017 for ObsEva's EDELWEISS Phase 2b clinical trial of OBE2109, its oral GnRH receptor antagonist for the treatment of endometriosis. Enrollment is targeted at 330 patients globally.
- The Phase 3 PRIMROSE 1 and 2 trials of OBE2109 for the treatment of uterine fibroids randomized its first patients to active therapy following screening that began earlier this year. Enrollment is targeted at approximately 1,000 women in total. The primary endpoint of these trials is the reduction of heavy menstrual bleeding.

- Clinical trial protocol was finalized and regulatory applications submitted to begin the PROLONG trial, a Phase 2a clinical trial of OBE022, ObsEva's oral prostaglandin F2 alpha receptor antagonist for the treatment of pre-term labor in pregnant women between 24 and 34 weeks of gestation.

### **Upcoming Milestones**

ObsEva expects to achieve the following clinical and pipeline milestones over the remainder of 2017 and 2018:

- Announcement of primary endpoint results from the 760 patient Phase 3 IMPLANT 2 clinical trial of nolasiban in assisted reproductive technology (ART) in the first quarter of 2018.
- Announcement of primary endpoint results from the 330 patient Phase 2b EDELWEISS trial of OBE2109 for the treatment of endometriosis in mid-2018.
- Commencement of the Phase 2a PROLONG clinical trial of OBE022 in pre-term labor in the fourth quarter of 2017, with preliminary results in approximately 60 patients expected in late 2018.
- Completion of patient enrollment of the Phase 3 PRIMROSE 1 and 2 trials of OBE2109 for the treatment of uterine fibroids in late 2018.

### **Third Quarter 2017 Financial Results**

Net loss for the third quarter of 2017 was \$17.0 million, or \$0.59 per basic and diluted share. Research and development expenses were \$13.9 million and general and administrative expenses were \$3.0 million for the quarter ended September 30, 2017. As of September 30, 2017, ObsEva had cash and cash equivalents of \$68.4 million, which does not include net proceeds of approximately \$56 million raised in the private equity financing that was announced on October 10, 2017.

### **Conference Call Information**

ObsEva will host a conference call and audio webcast today at 8:00 a.m. Eastern Time to provide a business update and discuss third quarter 2017 financial results. To participate in the conference call, please dial 844-419-1772 (domestic) or (213) 660-0921 (international) and refer to conference ID 2197575. The webcast can be accessed under the "Investors" section of ObsEva's website [www.obseva.com](http://www.obseva.com).

### **About ObsEva**

ObsEva is a clinical-stage biopharmaceutical company focused on the clinical development and commercialization of novel therapeutics for serious conditions that compromise a woman's reproductive health and pregnancy. Through strategic in-licensing and disciplined drug development, ObsEva has established a late-stage clinical pipeline with development programs focused on treating endometriosis, uterine fibroids, preterm labor and improving ART outcomes. ObsEva is listed on The NASDAQ Global Select Market and is trading under the ticker symbol "OBSV". For more information, please visit [www.obseva.com](http://www.obseva.com).

### **Cautionary Note Regarding Forward Looking Statements**

Any statements contained in this press release that do not describe historical facts may constitute forward-looking statements as that term is defined in the Private Securities Litigation Reform Act of 1995. These statements may be identified by words such as "believe", "expect", "may", "plan," "potential," "will," and similar expressions, and are based on ObsEva's current beliefs and expectations. These forward-looking statements include expectations regarding the clinical development of ObsEva's product candidates and the timing of enrollment in and data from clinical trials. These statements involve risks and uncertainties that could cause actual results to differ materially from those reflected in such statements. Risks and uncertainties that may cause actual results to differ materially include uncertainties inherent in the conduct of clinical trials, ObsEva's reliance on third parties over which it may not always have full control, and other risks and uncertainties that are described in the Risk Factors section of ObsEva's Annual Report on Form 20-F for the year ended December 31, 2016, and other filings ObsEva makes with the SEC. These documents are available on the Investors page of ObsEva's website at <http://www.obseva.com>. Any forward-looking statements speak only as of the date of this press release and are based on information available to ObsEva as of the date of this release, and ObsEva assumes no obligation to, and does not intend to, update any forward-looking statements, whether as a result of new information, future events or otherwise.

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## Consolidated Statements of Comprehensive Loss

(in USD '000, except per share data)	Three-month period ended September 30,		Nine-month period ended September 30,	
	2017	2016	2017	2016
	<i>unaudited</i>		<i>unaudited</i>	
<b>Other operating income</b> .....	3	7	11	37
<b>OPERATING EXPENSES</b>				
Research and development expenses .....	(13,910)	(5,978)	(40,983)	(15,544)
General and administrative expenses .....	(3,001)	(1,946)	(9,601)	(3,321)
<b>Total operating expenses</b> .....	<u>(16,911)</u>	<u>(7,924)</u>	<u>(50,584)</u>	<u>(18,865)</u>
<b>OPERATING LOSS</b> .....	<u>(16,908)</u>	<u>(7,917)</u>	<u>(50,573)</u>	<u>(18,828)</u>
Finance income .....	(106)	7	754	34
Finance expense .....	(1)	(36)	(1)	(267)
<b>NET LOSS BEFORE TAX</b> .....	<u>(17,015)</u>	<u>(7,946)</u>	<u>(49,820)</u>	<u>(19,061)</u>
Income tax benefit / (expense) .....	21	-	(36)	-
<b>NET LOSS FOR THE PERIOD</b> .....	<u>(16,994)</u>	<u>(7,946)</u>	<u>(49,856)</u>	<u>(19,061)</u>
<b>Net loss per share</b>				
Basic.....	(0.59)	(0.37)	(1.78)	(0.89)
Diluted.....	(0.59)	(0.37)	(1.78)	(0.89)
<b>OTHER COMPREHENSIVE INCOME</b>				
Currency translation differences .....	<u>-</u>	<u>554</u>	<u>-</u>	<u>1,956</u>
<b>TOTAL OTHER COMPREHENSIVE INCOME</b> .....	-	554	-	1,956
<b>TOTAL COMPREHENSIVE LOSS FOR THE PERIOD</b> .....	<u>(16,994)</u>	<u>(7,392)</u>	<u>(49,856)</u>	<u>(17,105)</u>

## Consolidated Balance Sheet

(in USD '000)	September 30, 2017	December 31, 2016
	<i>unaudited</i>	<i>audited</i>
<b>ASSETS</b>		
<b>Current assets</b>		
Cash and cash equivalents .....	68,358	25,508
Other receivables .....	683	783
Prepaid expenses and deferred costs.....	1,679	2,415
<b>Total current assets .....</b>	<b>70,720</b>	<b>28,706</b>
<b>Non-current assets</b>		
Furniture, fixtures and equipment.....	339	121
Intangible assets.....	21,608	16,608
Other long-term assets .....	190	90
<b>Total non-current assets.....</b>	<b>22,137</b>	<b>16,819</b>
<b>Total assets .....</b>	<b>92,857</b>	<b>45,525</b>
<b>LIABILITIES AND SHAREHOLDERS' EQUITY</b>		
<b>Current liabilities</b>		
Current tax liability.....	36	-
Other payables and current liabilities .....	2,662	2,383
Accrued expenses .....	5,752	4,269
<b>Total current liabilities.....</b>	<b>8,450</b>	<b>6,652</b>
<b>Non-current liabilities</b>		
Post-employment obligations .....	2,840	2,832
<b>Total non-current liabilities .....</b>	<b>2,840</b>	<b>2,832</b>
<b>Shareholders' equity</b>		
Share capital .....	2,265	1,740
Share premium.....	162,982	71,966
Reserves.....	5,775	1,934
Accumulated losses .....	(89,455)	(39,599)
<b>Total shareholders' equity .....</b>	<b>81,567</b>	<b>36,041</b>
<b>Total liabilities and shareholders' equity .....</b>	<b>92,857</b>	<b>45,525</b>

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