



Myovant Sciences to Host Webcast and Conference Call at 8:30 a.m. Eastern Time Wednesday, April 22 to Discuss Results from Phase 3 SPIRIT 2 Study Evaluating Once-Daily Relugolix Combination Therapy in Women with Endometriosis and from Ovulation Inhibition

April 21, 2020

BASEL, Switzerland, April 21, 2020 (GLOBE NEWSWIRE) -- Myovant Sciences (NYSE: MYOV), a healthcare company focused on redefining care for women's health and prostate cancer, today announced it will hold a webcast and conference call beginning at 8:30 a.m. Eastern Time / 5:30 a.m. Pacific Time on Wednesday, April 22, 2020, to discuss results from the Phase 3 SPIRIT 2 study of once-daily relugolix combination therapy (relugolix 40 mg plus estradiol 1.0 mg, and norethindrone acetate 0.5 mg) in women with pain associated with endometriosis. The company will also discuss results from a separate ovulation inhibition study with relugolix combination therapy.

Webcast/Teleconference Details

To participate in the live conference call, please dial 1-800-532-3746 for domestic callers and +1-470-495-9166 for international callers. A live webcast of the conference call will also be available on the investor relations page of Myovant's website at investors.myovant.com and will remain archived on Myovant's website for at least 30 days.

About Myovant Sciences

Myovant Sciences aspires to be the leading healthcare company focused on redefining care for women's health and prostate cancer. The company's lead product candidate is relugolix, a once-daily, oral GnRH receptor antagonist. The company has three late-stage clinical programs for relugolix in uterine fibroids, endometriosis, and prostate cancer. The company is also developing MVT-602, an oligopeptide kisspeptin-1 receptor agonist, that has completed a Phase 2a study for the treatment of female infertility as part of assisted reproduction. Takeda Pharmaceuticals International AG, a subsidiary of Takeda Pharmaceutical Company Limited, the originator of relugolix, previously granted the company a worldwide license to develop and commercialize relugolix (excluding Japan and certain other Asian countries) and an exclusive license to develop and commercialize MVT-602 in all countries worldwide. Sumitovant Biopharma, Ltd., a wholly owned subsidiary of Sumitomo Dainippon Pharma Co., Ltd., is the majority shareholder of Myovant. For more information, please visit the company's website at www.myovant.com. Follow [@Myovant](https://twitter.com/Myovant) on Twitter and [LinkedIn](https://www.linkedin.com/company/myovant).

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