



Myovant Sciences to Host Second Fiscal Quarter 2021 Earnings Conference Call at 8:30 a.m. Eastern Time on October 26, 2021

October 12, 2021

BASEL, Switzerland, Oct. 12, 2021 (GLOBE NEWSWIRE) -- Myovant Sciences (NYSE: MYOV), a healthcare company focused on redefining care for women and for men, today announced it will host a webcast and conference call to discuss corporate updates and financial results for its second fiscal quarter 2021, ended September 30, 2021. The webcast and conference call will be held at 8:30 a.m. Eastern Time / 5:30 a.m. Pacific Time on October 26, 2021.

Investors and the general public may access a live webcast of the call by visiting the investor relations page of Myovant's website at investors.myovant.com. Institutional investors and analysts may also participate in the conference call by dialing 1-800-532-3746 in the U.S. or +1-470-495-9166 from outside the U.S.

A replay of the webcast, along with the earnings press release and presentation materials, will be archived on Myovant's investor relations website.

About Myovant Sciences

Myovant Sciences aspires to redefine care for women and for men through purpose-driven science, empowering medicines, and transformative advocacy. Founded in 2016, we have two FDA-approved products. ORGOVYX[®] (relugolix) was approved by the U.S. Food and Drug Administration in 2020 as the first and only oral gonadotropin-releasing hormone (GnRH) receptor antagonist for the treatment of adult patients with advanced prostate cancer, and relugolix is also under regulatory review in Europe for men with advanced prostate cancer. Relugolix combination tablet (relugolix 40 mg, estradiol 1.0 mg, and norethindrone acetate 0.5 mg) was approved in 2021 in the U.S. as MYFEMBREE[®] as the first once-daily treatment for the management of heavy menstrual bleeding associated with uterine fibroids in premenopausal women, and by the European Commission as RYEQO[®] for the treatment of moderate to severe symptoms of uterine fibroids in adult women of reproductive age. We have filed a supplemental New Drug Application (sNDA) for the management of moderate to severe pain associated with endometriosis, which was accepted for review by the FDA in September 2021. The therapy is also being assessed for the prevention of pregnancy. We are also developing MVT-602, an oligopeptide kisspeptin-1 receptor agonist, which has completed a Phase 2a study for female infertility as part of assisted reproduction. Sumitovant Biopharma, Ltd., a wholly owned subsidiary of Sumitomo Dainippon Pharma Co., Ltd., is our majority shareholder. For more information, please visit our 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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Source: Myovant Sciences, Inc.