



## **Lumos Pharma to Report Second Quarter 2022 Financial Results and Host Conference Call on August 9, 2022**

July 27, 2022

AUSTIN, Texas, July 27, 2022 (GLOBE NEWSWIRE) -- [Lumos Pharma, Inc.](#) (NASDAQ:LUMO), a clinical-stage biopharmaceutical company, today announced it will report its second quarter 2022 financial results after market close on Tuesday, August 9, 2022. The company will host a conference call and webcast at 4:30 PM ET that day to discuss these financial results and provide an update on clinical and corporate activities. There will be a question-and-answer session following the prepared remarks.

Investors and the general public are invited to listen to a live audio webcast of the conference call, which may be accessed five minutes prior to the start of the call by dialing (833) 634-2295 (U.S.) or +1 (412) 902-4176 (international), or through the link, <https://edge.media-server.com/mmc/p/ofen9o6f>. The link to the live webcast may also be found in the "Investors & Media" section of the Lumos Pharma website, under "[Events & Presentations](#)." A replay of the call will be available for two weeks from the date of the call and may be accessed through the same link above or by dialing (877) 344-7529 (U.S.) or +1 (412) 317-0088 (international) and using the passcode: 1602592.

### **About Lumos Pharma**

Lumos Pharma, Inc. is a clinical stage biopharmaceutical company focused on the development and commercialization of therapeutics for rare diseases. Lumos Pharma was founded and is led by a management team with longstanding experience in rare disease drug development and received early funding from leading healthcare investors, including Deerfield Management, a fund managed by Blackstone Life Sciences, Roche Venture Fund, New Enterprise Associates (NEA), Santé Ventures, and UCB. Lumos Pharma's lead therapeutic candidate is LUM-201, an oral growth hormone stimulating small molecule, currently being evaluated in a Phase 2 clinical trial, the OraGrowthH210 Trial, and a PK/PD trial, the OraGrowthH212 Trial, for the treatment of Pediatric Growth Hormone Deficiency (PGHD). If approved by the FDA, LUM-201 would provide an orally administered alternative to recombinant growth hormone injections that PGHD patients otherwise endure for many years of treatment. LUM-201 has received Orphan Drug Designation in both the US and EU. For more information, please visit <https://lumos-pharma.com/>.

Source: Lumos Pharma, Inc.

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