

Fusion Pharmaceuticals To Present At The Oppenheimer 34th Annual Healthcare Life Sciences Conference

HAMILTON, ON and BOSTON, Feb. 7, 2024 /PRNewswire/ -- Fusion Pharmaceuticals Inc. (Nasdaq: FUSN), a clinical-stage oncology company focused on developing next-generation radiopharmaceuticals as precision medicines, today announced that the Company will present virtually at the Oppenheimer 34th Annual Healthcare Life Sciences Conference on Wednesday, February 14, 2024, at 10:00 a.m. ET. Presenting on behalf of Fusion will be Chief Executive Officer John Valliant, Ph.D.

A webcast of the event will be available on the "Events and Presentations" page in the "Investors & Media" section of the Company's website at <https://ir.fusionpharma.com/events-webcasts>. A replay of the webcast will be archived on the Company's website for 90 days following the presentation date.

About Fusion

Fusion Pharmaceuticals is a clinical-stage oncology company focused on developing next-generation radiopharmaceuticals as precision medicines. Fusion connects alpha particle emitting isotopes to various targeting molecules in order to selectively deliver the alpha emitting payloads to tumors. Fusion's clinical portfolio includes: FPI-2265 targeting prostate specific membrane antigen (PSMA) for metastatic castration resistant prostate cancer currently in a Phase 2 trial; FPI-1434 targeting insulin-like growth factor 1 receptor currently in a Phase 1 clinical trial; and FPI-2059, a small molecule targeting neurotensin receptor 1 (NTSR1), currently in a Phase 1 trial. In addition to a robust proprietary pipeline, Fusion has a collaboration with AstraZeneca to jointly develop novel targeted alpha therapies (TATs) and combination programs between Fusion's TATs and AstraZeneca's DNA Damage Response Inhibitors (DDRIs) and immuno-oncology agents. The Company received IND clearance for FPI-2068, the first novel TAT under the collaboration, which targets EGFR-cMET. Fusion has also entered into a collaboration with Merck to evaluate FPI-1434 in combination with Merck's KEYTRUDA® (pembrolizumab) in patients with solid tumors expressing IGF-1R. Fusion has a Good Manufacturing Practice (GMP) compliant radiopharmaceutical manufacturing facility designed to support manufacturing of the Company's growing pipeline of TATs. To support Fusion's growing pipeline of TATs, the Company has signed strategic actinium supply agreements with Niowave, Inc. and BWXT Medical.

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