



## **X4 Pharmaceuticals to Announce Second Quarter 2022 Financial Results and Host a Conference Call and Webcast on August 4, 2022**

July 26, 2022

**BOSTON, July 26, 2022 (GLOBE NEWSWIRE) -- [X4 Pharmaceuticals, Inc.](#)** (Nasdaq: XFOR), a leader in the discovery and development of novel CXCR4-targeted small molecule therapeutics to benefit patients with diseases of the immune system, today announced that it will report its financial results for the second quarter ended June 30, 2022, and provide an update on recent business highlights, on August 4, 2022.

The Company will host a conference call and webcast on the same day at 8:30 a.m. ET to discuss these financial results and business highlights. The conference call can be accessed by dialing 1-855-327-6837 from the United States or 1-631-891-4304 internationally, followed by the conference ID: 10019589. The live webcast can be accessed on the investor relations section of X4 Pharmaceuticals' website at [www.x4pharma.com](http://www.x4pharma.com). Following the completion of the call, a webcast replay of the conference call will be available on the website.

### **About X4 Pharmaceuticals**

X4 Pharmaceuticals is a late-stage clinical biopharmaceutical company leading the discovery and development of novel therapies for people with diseases of the immune system. Our lead clinical candidate is mavorixafor, a first-in-class, small molecule antagonist of chemokine receptor CXCR4 that is being developed as a once-daily oral therapy. Due to mavorixafor's ability to antagonize CXCR4 and improve the healthy maturation and mobilization of white blood cells, we believe that mavorixafor has the potential to provide therapeutic benefit across a wide variety of diseases, including a range of chronic neutropenic disorders and certain types of cancer. The efficacy and safety of mavorixafor are being evaluated in a global Phase 3 clinical trial in patients with WHIM syndrome, a rare, inherited, primary immunodeficiency disease caused by genetic mutations in the CXCR4 receptor gene. We are also studying mavorixafor in two Phase 1b clinical trials – one as a monotherapy in patients with chronic neutropenia disorders including congenital, idiopathic, and cyclic neutropenia, and one in combination with ibrutinib in patients with Waldenström's macroglobulinemia (WM), a rare B-cell lymphoma. Further clinical development of mavorixafor in WM will now be subject to completing a strategic partnership as we focus our resources on advancing mavorixafor for the benefit of patients with chronic neutropenic disorders. We continue to leverage our insights into CXCR4 biology at our corporate headquarters in Boston, Massachusetts and at our research facility in Vienna, Austria. For more information, please visit our website at [www.x4pharma.com](http://www.x4pharma.com).

### **Corporate:**

Daniel Ferry (investors)  
Managing Director, LifeSci Advisors  
[daniel@lifesciadvisors.com](mailto:daniel@lifesciadvisors.com)  
(617) 430-7576

Mónica Rouco Molina, Ph.D. (media)

Account Supervisor, LifeSci Communications  
[mroucomolina@lifescicomms.com](mailto:mroucomolina@lifescicomms.com)



Source: X4 Pharmaceuticals