

October 28, 2024



Iterum Therapeutics to Host Morning Conference Call on U.S. FDA Approval of ORLYNVAH™ (Oral Sulopenem) for the Treatment of Uncomplicated Urinary Tract Infections

Monday, October 28, 2024, at 8:30 a.m. EDT

DUBLIN and CHICAGO, Oct. 28, 2024 /PRNewswire/ --



WHO: [Iterum Therapeutics plc](#) (Nasdaq: ITRM) is focused on delivering differentiated anti-infectives aimed at combatting the global crisis of multi-drug resistant pathogens to significantly improve the lives of people affected by serious and life-threatening diseases around the world.

WHAT: Conference call to discuss U.S. Food and Drug Administration (FDA) approval of Iterum's ORLYNVAH™ (Oral Sulopenem) for the treatment of uncomplicated urinary tract infections (uUTIs).

Speakers include: Corey Fishman (CEO) and Steve Aronin (Senior Vice President and Head of Clinical Development)

WHY: ORLYNVAH™ (sulopenem etzadroxil and probenecid) is the first oral penem approved for use in the U.S. and the first FDA-approved product for Iterum. ORLYNVAH™ is approved for the treatment of uUTIs caused by the designated microorganisms *Escherichia coli*, *Klebsiella pneumoniae* or *Proteus mirabilis* in adult women who have limited or no alternative oral antibacterial treatment options. It is only the second FDA-approved treatment for uUTIs in the past two decades.

For more details, view the press release issued Friday [here](#).

WHEN: Monday, October 28, 2024
8:30 a.m. Eastern Daylight Time

Dial-in information:
United States: +1 833-470-1428 | International: +1 404-975-4839
Access code: 936149

The conference call replay will be available in the [Events & Presentations](#) page of Iterum's website following the call.

About Iterum Therapeutics plc

Iterum Therapeutics plc is focused on delivering differentiated anti-infectives aimed at combatting the global crisis of multi-drug resistant pathogens to significantly improve the lives of people affected by serious and life-threatening diseases around the world. Iterum is advancing the development of its first compound, sulopenem, a novel penem anti-infective compound, with an oral formulation and IV formulation. Sulopenem has demonstrated potent in vitro activity against a wide variety of gram-negative, gram-positive and anaerobic bacteria resistant to other antibiotics. Iterum has received approval of its NDA for ORLYNVAH™ (oral sulopenem) for the treatment of uncomplicated urinary tract infections caused by the designated microorganisms *Escherichia coli*, *Klebsiella pneumoniae*, or *Proteus mirabilis* in adult women with limited or no alternative oral antibacterial treatment options by the U.S. Food and Drug Administration and has received Qualified Infectious Disease Product (QIDP) and Fast Track designations for its oral and IV formulations of sulopenem in seven indications. For more information, please visit www.iterumtx.com.

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