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**UNITED STATES**  
**SECURITIES AND EXCHANGE COMMISSION**  
Washington, D.C. 20549

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**FORM 8-K**

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**CURRENT REPORT**  
**Pursuant to Section 13 or 15(d)**  
**of the Securities Exchange Act of 1934**

Date of Report (Date of earliest event reported): August 14, 2019

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**Iterum Therapeutics plc**

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(Exact name of registrant as specified in its charter)

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**Ireland**

(State or other jurisdiction  
of incorporation)

**001-38503**

(Commission File Number)

**98-1283148**

(IRS Employer  
Identification No.)

**Block 2 Floor 3, Harcourt Centre,  
Harcourt Street,  
Dublin 2, Ireland**

(Address of principal executive offices)

(Zip Code)

Registrant's telephone number, including area code: **+353 1 903 8920**

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Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- ☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- ☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- ☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- ☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ☒

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☒

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Ordinary Shares, \$0.01 par value per share	ITRM	The Nasdaq Stock Market LLC

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**Item 2.02. Results of Operations and Financial Condition.**

On August 14, 2019, Iterum Therapeutics plc issued a press release announcing certain financial results for its second quarter ended June 30, 2019. A copy of the press release is furnished herewith as Exhibit 99.1.

The information in this current report on Form 8-K, including the press release attached as Exhibit 99.1 hereto, is being furnished, but shall not be deemed to be “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that section or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended. The information contained herein and in the accompanying exhibit shall not be incorporated by reference into any filing with the U.S. Securities and Exchange Commission made by Iterum, whether made before or after the date hereof, regardless of any general incorporation language in such filing.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits

<b>Exhibit No.</b>	<b>Description</b>
<a href="#"><u>99.1</u></a>	<a href="#"><u>Press Release of Iterum Therapeutics plc, dated August 14, 2019</u></a>

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**SIGNATURE**

Pursuant to the requirements of the Securities Exchange Act of 1934, as amended, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

**Iterum Therapeutics plc**

Dated: August 14, 2019

By: /s/ Corey N. Fishman  
Corey N. Fishman  
Chief Executive Officer

**FOR IMMEDIATE RELEASE****Iterum Therapeutics Reports Second Quarter 2019 Financial Results**

*--Phase 3 Topline Data anticipated in the fourth quarter--*

*-- NDA filings expected in first quarter of 2020--*

**DUBLIN, Ireland and CHICAGO, August 14, 2019** -- Iterum Therapeutics plc (Nasdaq: ITRM), a clinical-stage pharmaceutical company focused on developing next generation oral and IV antibiotics to treat infections caused by multi-drug resistant pathogens in both community and hospital settings, today reported financial results for the second quarter ended June 30, 2019.

“The second quarter was marked by continued progress in our three pivotal Phase 3 clinical trials of sulopenem,” said Corey Fishman, Chief Executive Officer of Iterum Therapeutics plc. “Based on current enrollment rates, we anticipate topline data from the complicated intra-abdominal (cIAI) and the complicated urinary tract infection (cUTI) trials in the fourth quarter. This would allow us the flexibility to file our new drug applications (NDAs) in the first quarter of 2020. Our trial for uncomplicated urinary tract infections (uUTI) just passed two-thirds enrollment, which triggers a pre-planned review of the blinded data for potential sample size adjustment. The outcome of this analysis will determine the enrollment timeline, which will guide our decision on the optimal filing strategy for the uUTI data.”

**Recent Highlights**

- **Presented research at ASM Microbe that supports the need for new antibiotics to treat uUTIs:** In June, Iterum presented data that showed patients with uUTIs in the community treated with the most commonly prescribed antibiotic for that infection, ciprofloxacin, have a significantly greater rate of treatment failure when the organism is quinolone resistant. Based on our uUTI trial, Iterum estimates that resistance to quinolones in the community is greater than 25% in many geographies in the U.S., and greater than 30% outside of the U.S.

**Second Quarter 2019 Financial Results**

As of June 30, 2019, Iterum had cash and cash equivalents of \$51.2 million and approximately 14.4 million shares outstanding. Iterum expects that its cash and cash equivalents will be sufficient to fund operations into 2020.

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Research and development (R&D) expenses for the second quarter of 2019 were \$24.4 million compared to \$13.7 million for the same period in 2018. The increase was primarily due to higher clinical trial expenses associated with our three Phase 3 clinical trials initiated in the third quarter of 2018, partially offset by a reduction in chemistry, manufacturing and control-related expenses as a result of the completion of manufacturing of clinical trial materials for our Phase 3 clinical trials by our primary suppliers.

General and administrative (G&A) expenses for the second quarter of 2019 were \$2.9 million compared to \$1.9 million for the same period in 2018. The increase was primarily due to increased costs associated with operating as a public company, additional headcount to support business activities, and increased marketing expenses.

For the second quarter of 2019, Iterum reported a net loss of \$27.6 million compared to a net loss of \$15.7 million for the same period in 2018.

### **About Sulopenem**

Sulopenem, a novel penem anti-infective compound with oral and IV formulations, has demonstrated potent *in vitro* activity against a wide variety of gram-negative, gram-positive and anaerobic bacteria resistant to other antibiotics. If approved, sulopenem will help address the significant clinical and economic need for new oral antibiotics that enable the effective treatment of resistant pathogens in the community, make possible the avoidance of hospitalization, and facilitate early hospital discharge by providing continuity-of-care step-down therapy. The safety profile of IV sulopenem has been documented in a Phase 2 program. Oral and IV sulopenem are being evaluated in three pivotal Phase 3 clinical trials for uncomplicated urinary tract infections, complicated urinary tract infections, and complicated intra-abdominal infections.

Probenecid, which is being co-administered with sulopenem in a bilayer tablet, is approved as an adjuvant to therapy for elevation and prolongation of plasma levels of  $\beta$ -lactam compounds including penicillin, ampicillin, methicillin, oxacillin, cloxacillin, and nafcillin.

### **About Iterum Therapeutics plc**

Iterum Therapeutics plc is a clinical-stage pharmaceutical company dedicated to developing 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 Therapeutics is advancing its first compound, sulopenem, a novel penem anti-infective compound, in Phase 3 clinical development with oral and IV formulations. Sulopenem has demonstrated potent *in vitro* activity against a wide variety of gram-negative, gram-positive and anaerobic bacteria resistant to other antibiotics. Iterum Therapeutics has received QIDP and Fast Track designations for its oral and IV formulations of sulopenem in seven indications. For more information, please visit <http://www.iterumtx.com>.

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## **Forward Looking Statements**

This press release contains forward-looking statements. These forward-looking statements include, without limitation, statements regarding expectations about future revenue, expenses, cash flows and net income or loss, the sufficiency of cash resources, the development, therapeutic and market potential of sulopenem, and the timing, progress and results of clinical trials and regulatory submissions. In some cases, forward-looking statements can be identified by words such as “may,” “believes,” “intends,” “seeks,” “anticipates,” “plans,” “estimates,” “expects,” “should,” “assumes,” “continues,” “could,” “will,” “future,” “potential” or the negative of these or similar terms and phrases. Forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause Iterum Therapeutics’ actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. Forward-looking statements include all matters that are not historical facts. Actual future results may be materially different from what is expected due to factors largely outside Iterum Therapeutics’ control, including the uncertainties inherent in the initiation and conduct of clinical trials, clinical trial patient enrollment, availability and timing of data from clinical trials, changes in regulatory requirements or decisions of regulatory authorities, changes in public policy or legislation, commercialization plans and timelines, if approved, the actions of third-party clinical research organizations, suppliers and manufacturers and other factors discussed under the caption “Risk Factors” in its Quarterly Report on Form 10- Q filed with the Securities and Exchange Commission (the “SEC”) on August 14, 2019, and other documents filed with the SEC from time to time. Forward-looking statements represent Iterum Therapeutics’ beliefs and assumptions only as of the date of this press release. Except as required by law, Iterum Therapeutics assumes no obligation to update these forward-looking statements publicly, or to update the reasons actual results could differ materially from those anticipated in the forward-looking statements, even if new information becomes available in the future.

### **Investor Contact:**

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Chief Financial Officer  
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**ITERUM THERAPEUTICS PLC**  
**Condensed Consolidated Statement of Operations**  
(In thousands except share and per share data)  
(Unaudited)

	Three months ended June 30,		Six months ended June 30,	
	2019	2018	2019	2018
Revenue	\$ —	\$ 185	\$ 37	\$ 376
Operating expenses:				
Research and development	(24,439)	(13,725)	(41,826)	(24,604)
General and administrative	(2,939)	(1,886)	(6,055)	(3,401)
Total operating expenses	(27,378)	(15,611)	(47,881)	(28,005)
Operating loss	(27,378)	(15,426)	(47,844)	(27,629)
Interest (expense) / income, net	(135)	(76)	(239)	9
Other income / (expense), net	32	(177)	156	(116)
Income tax expense	(157)	(68)	(291)	(157)
Net loss attributable to ordinary shareholders	\$ (27,638)	\$ (15,747)	\$ (48,218)	\$ (27,893)
Net loss per share attributable to ordinary shareholders – basic and diluted	\$ (1.93)	\$ (2.22)	\$ (3.37)	\$ (6.72)
Weighted average ordinary shares outstanding – basic and diluted	14,340,231	7,085,655	14,316,497	4,148,535

**ITERUM THERAPEUTICS PLC**  
**Condensed Consolidated Balance Sheet Data**  
(In thousands)  
(Unaudited)

	As of June 30, 2019	As of December 31, 2018
Cash, cash equivalents and short-term investments	\$ 51,241	\$ 84,551
Other assets	20,052	13,320
<b>Total assets</b>	<b>\$ 71,293</b>	<b>\$ 97,871</b>
Long-term debt, less current portion	10,771	13,079
Other liabilities	35,946	13,170
Total liabilities	46,717	26,249
Total shareholders' equity	24,576	71,622
<b>Total liabilities and shareholders' equity</b>	<b>\$ 71,293</b>	<b>\$ 97,871</b>