

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

**FORM 8-K**

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

Date of Report (Date of earliest event reported): April 23, 2025

**Cocrystal Pharma, Inc.**

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation)	001-38418 (Commission File Number)	35-2528215 (IRS Employer Identification No.)
19805 N. Creek Parkway Bothell, WA (Address of principal executive offices)		98011 (Zip Code)

Registrant's telephone number, including area code: (877) 262-7123

N/A

(Former name or former address, if changed since last report.)

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 (see  
General Instruction A.2. below):

- 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))

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

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock	COCP	The Nasdaq Stock Market LLC (The Nasdaq Capital Market)

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.

**Item 7.01 Regulation FD Disclosure.**

On April 23, 2025, Cocrystal Pharma, Inc. (the "Company") is making a presentation to certain members of the investment community. A copy of the slide presentation is being furnished as Exhibit 99.1.

The information in this Item 7.01 (including Exhibit 99.1) shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934 (the "Exchange Act") or otherwise subject to the liabilities under such section, and shall not be deemed to be incorporated by reference into any filing of the Company under the Securities Act of 1933 or the Exchange Act.

**Item 9.01 Financial Statements and Exhibits.**

Exhibit No.	Description
99.1	<a href="#">Cocrystal Pharma, Inc. Corporate Presentation, dated April 23, 2025</a>
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURES

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

Dated: April 23, 2025

**Cocrystal Pharma, Inc.**

By: /s/James Martin

Name: James Martin

Title: Chief Financial Officer and Co-Chief Executive Officer

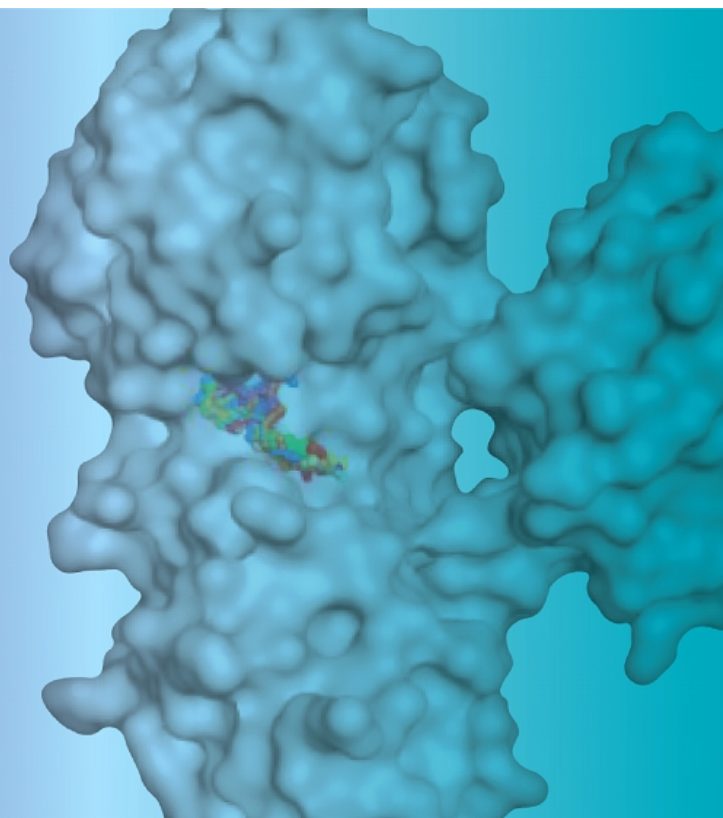
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Potent antivirals to combat some  
of the most serious diseases  
facing humanity

Life Science Innovation Northwest  
April 2025

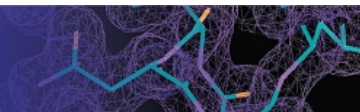
Nasdaq: COCP  
[www.cocrystalpharma.com](http://www.cocrystalpharma.com)



## Forward-Looking Statements

This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, including our ability to deliver significant growth from our multiple clinical assets, our plans for a human challenge study in 2025, the results of our ongoing Phase 2a study for oral influenza PB2 inhibitor, and the expected sufficiency of our cash balance to fund our planned operations.

Forward-looking statements are prefaced by words such as “anticipate,” “expect,” “plan,” “could,” “may,” “will,” “should,” “would,” “intend,” “seem,” “potential,” “appear,” “continue,” “future,” “believe,” “estimate,” “forecast,” “project,” and similar words. Forward-looking statements are based on our current expectations and assumptions regarding our business, the economy and other future conditions. Because forward-looking statements relate to the future, they are subject to inherent uncertainties, risks and changes in circumstances that are difficult to predict. We caution you, therefore, against relying on any of these forward-looking statements. Our actual results may differ materially from those contemplated by the forward-looking statements for a variety of reasons, including, without limitation, the risks arising from the tariff policy of the United States and the adverse effect on the financial markets the impact of possible cuts in federal spending on healthcare, the possibility of a recession, and the geopolitical conflicts in Israel and Ukraine on our Company, our collaboration partners, and on the U.S., UK, Australia and global economies, our ability to proceed with studies including recruiting volunteers for and procuring or manufacturing materials for such studies by our clinical research organizations and vendors, the results of our CRO’s studies referred to above, our and our collaboration partners’ technology and software performing as expected and maintenance and protection of related intellectual property rights, general risks arising from clinical trials, receipt of regulatory approvals, regulatory changes, development of effective treatments and/or vaccines by competitors, and potential mutations in the viruses we are targeting which may result in variants that are resistant to a product candidate we develop. Further information on our risk factors is contained in our filings with the Securities and Exchange Commission, including our Annual Report on Form 10-K for the year ended December 31, 2024. Any forward-looking statement made by us in this presentation speaks only as of the date on which it is made. Factors or events that could cause our actual results to differ may emerge from time to time, and it is not possible for us to predict all of them. We undertake no obligation to publicly update any forward-looking statement, whether as a result of new information, future developments or otherwise, except as may be required by law.



*Applying powerful, proprietary drug discovery platform technology to develop first- and best-in-class broad-spectrum antiviral drugs*

### Advancing programs in high-value antiviral drug targets

- Norovirus
- Influenza
- Coronavirus and respiratory viruses

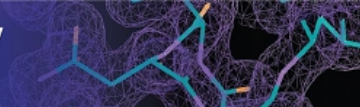
### Drug candidates with clinically validated mechanisms of action

- Effectively cure viral diseases
- Broad-spectrum and potent antiviral activity
- Designed to be effective for emerging variants and existing drug-resistant viruses
- Multiple routes of administration (oral, inhalation, and injectable)

### Proprietary drug discovery platform technology

- Unique drug discovery platform technology developed with Nobel Prize-winning technology

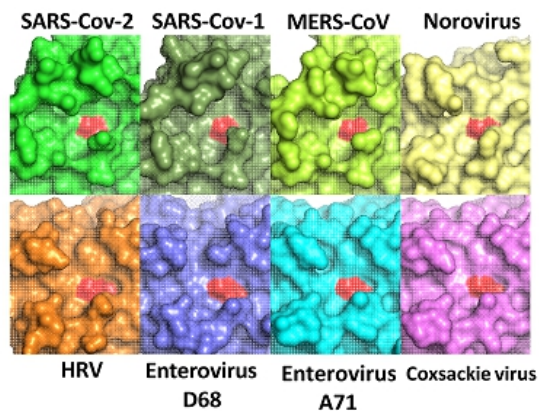
## Cocrystal's Structure-Based Drug Discovery Platform Technology For Pan-Viral Direct Acting Antiviral Development



### Cocrystal technology uniquely offers:

- 1 Systematic analysis of drug binding pockets
- 2 Rapid cocrystal structure determination
- 3 Structural insight into drug resistance
- 4 Novel structural hits and pockets
- 5 Multiple leads

### Cocrystal pan-viral inhibitors target highly conserved viral protease active site

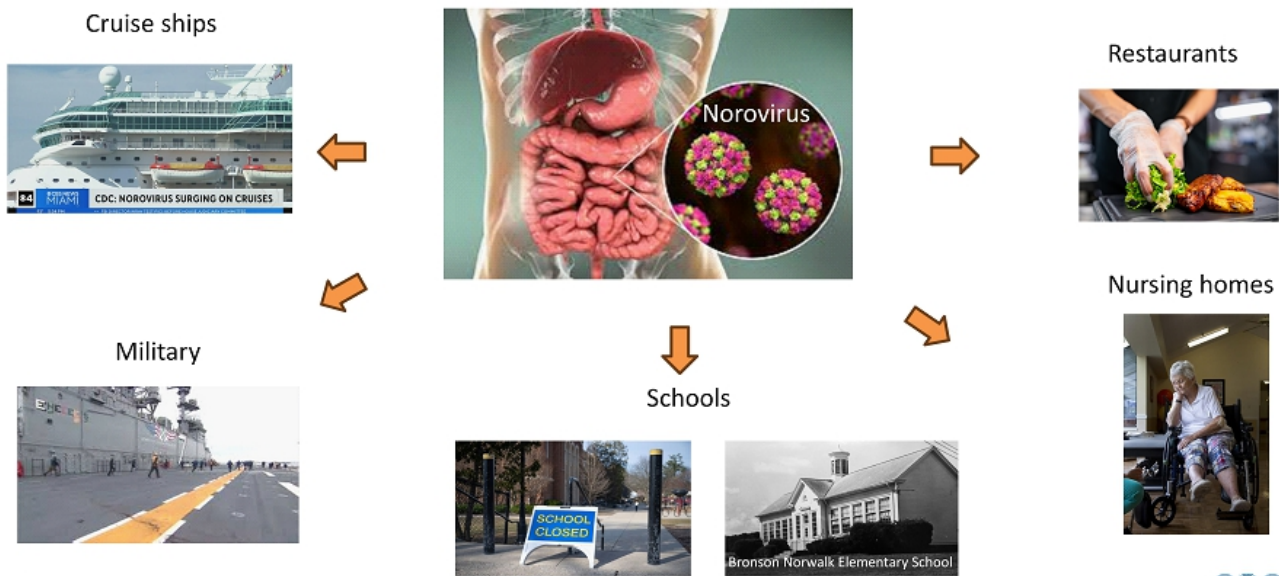


## Multiple clinical assets poised to deliver significant growth

Program	Candidate	Discovery	Preclinical	Phase 1	Phase 2	Phase 3
Norovirus & Coronavirus	Oral Pan-viral protease inhibitor CDI-988	[Progress bar: Discovery to Phase 1]				
	Replication inhibitors	[Progress bar: Discovery to Preclinical]				
Respiratory viruses	Pan-viral inhibitors	[Progress bar: Discovery to Preclinical]				
Influenza A	Oral PB2 inhibitor CC-42344	[Progress bar: Discovery to Phase 2]				
	Inhaled PB2 inhibitor CC-42344	[Progress bar: Discovery to Phase 1]				
Influenza A/B	Replication inhibitors	[Progress bar: Discovery to Preclinical]				

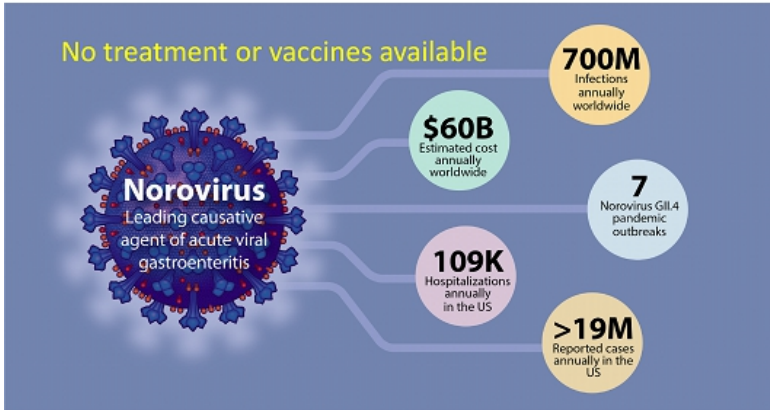
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## Norovirus Infection : No Approved Treatments or Vaccines Available



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# Norovirus Viral Gastroenteritis Represents a Large Global Market: \$72.7 Billion by 2032



Cocrystal's pan-viral protease inhibitor, CDI-988:

- First-in-class antiviral for norovirus
- Potential for both prevention and treatment
- Phase 1 SAD and MAD studies complete
- Human challenge study planned in 2025

\*Market Research Future, [Viral Gastroenteritis](#), 2023  
CDC: Norovirus Disease in the United States <https://www.cdc.gov/norovirus/burden.html>

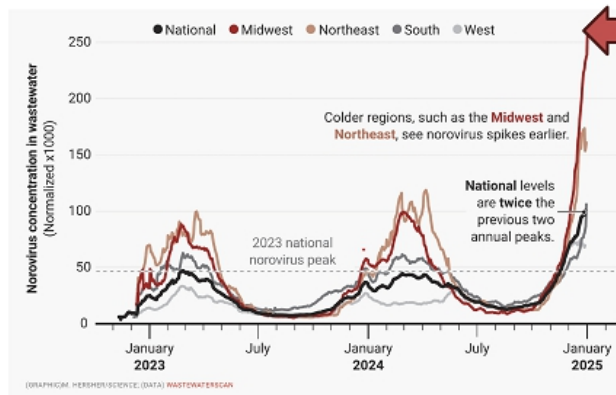
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## Big Surge of Norovirus Outbreaks in 2024-2025 After COVID-19 Pandemic

### Why the 'Ferrari of viruses' is surging through the Northern Hemisphere

Norovirus, which causes explosive diarrhea and vomiting, may be on the rise because of an antibody-dodging variant and post-COVID-19 socializing

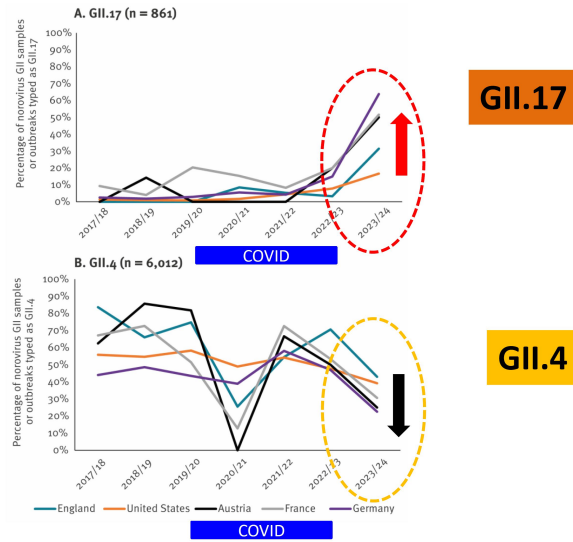
13 JAN 2025 · 6:00 PM ET · BY JON COHEN



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Reference: Science, 13 January 2025

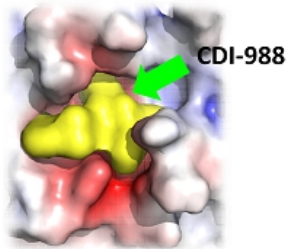
# Norovirus GII.17 Variants Become Dominant: >70% Responsible For Recent Norovirus Outbreaks



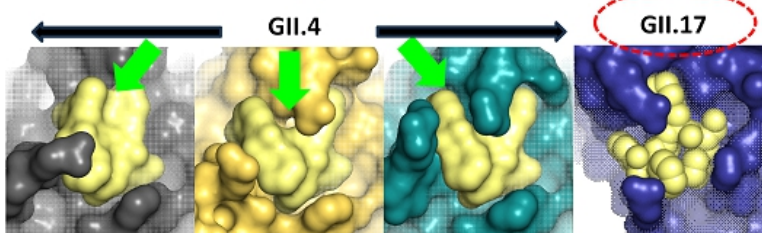
<https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2024.29.39.2400625>

# Pan-Viral Protease Inhibitor CDI-988 For Norovirus GII.4 and GII.17 and COVID

## (A) SARS-CoV-2



## (B) Norovirus



Cocrystal structures of norovirus proteases with CDI-988

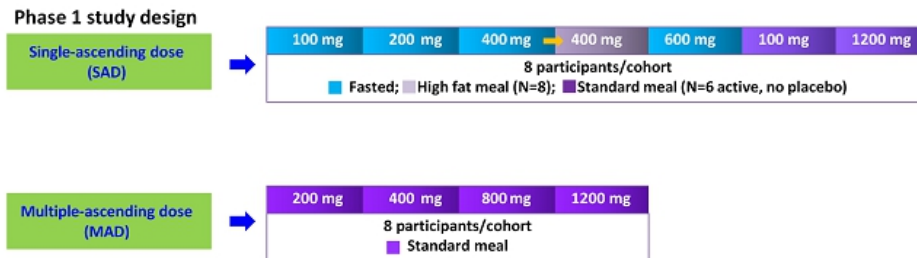
- First-in-class antiviral for norovirus
- Developed using Cocrystal's proprietary drug discovery platform technology
- Binds to a highly conserved region required for viral proteases
- Exhibits pan-viral activity against pandemic norovirus and SARS-CoV-2, SARS-CoV, and MERS-CoV strains
- Phase 1 complete
- One molecule, multiple indications

# Oral Pan-viral Protease Inhibitor CDI-988 Showed Favorable Safety and Tolerability

- Single-center, randomized, double-blind, placebo-controlled
- Single-ascending dose (SAD) and Multiple-ascending dose (MAD) cohorts
- Healthy adult volunteers (18 – 55 years old)

## Phase 1 study summary

- All dose cohorts well tolerated
- No serious adverse effects (SAEs)
- No treatment-related study discontinuations



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CRYSTAL  
PHARMA, INC.

## Investment Highlights

- Targeting multibillion-dollar, global markets for the treatment of acute and pandemic viral diseases
- Proprietary structure-based drug discovery platform technology provides opportunity for discovery and development of novel, broad-spectrum drug candidates
- Advancing multiple clinical programs
  - First-in-class dual oral norovirus and coronavirus protease inhibitor CDI-988
  - Oral influenza PB2 inhibitor CC-42344 – Phase 2a study continues in 2025
- Developing multiple discovery programs for respiratory viral diseases
  - Pan-viral protease inhibitors and influenza replication inhibitors
- Exploring pandemic preparedness collaboration opportunities
- Seasoned leadership includes experienced management, senior scientists and two Nobel laureates
- Cost-efficient operations and clean capital structure; cash sufficient to fund planned operations

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