# **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): January 13, 2025

# **Cocrystal Pharma, Inc.** (Exact name of registrant as specified in its charter)

	act name of registrant as specified in its cha	iiter)
Delaware	001-38418	35-2528215
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)
19805 N. Creek Parkway Bothell, WA		98011
(Address of principal executive offices)		(Zip Code)
Registrant's	telephone number, including area code: (8)	77) 262-7123
(Former	$\frac{N/A}{N/A}$ name or former address, if changed since la	ist report )
× ·		igation of the registrant under any of the following provisions <u>see</u>
□ Written communications pursuant to Rule 425 under the Secu	rities Act (17 CFR 230.425)	
□ Soliciting material pursuant to Rule 14a-12 under the Exchange	ge Act (17 CFR 240.14a-12)	
Pre-commencement communications pursuant to Rule 14d-2(	b) under the Exchange Act (17 CFR 240.14	d-2(b))
Pre-commencement communications pursuant to Rule 13e-4(	c) under the Exchange Act (17 CFR 240.13	e-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	СОСР	The Nasdaq Stock Market LLC (The Nasdaq Capital Market)
Indicate by check mark whether the registrant is an emerging grow the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).	wth company as defined in Rule 405 of the	Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of
		Emerging growth company $\Box$
If an emerging growth company indicate by check mark if the re-	gistrant 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 Exe		
Item 7.01 Regulation FD Disclosure.		
On January 13, 2025, Cocrystal Pharma, Inc. (the "Company") is being furnished as Exhibit 99.1.	making a presentation to certain members	s of the investment community. A copy of the slide presentation is
		n 18 of the Securities Exchange Act of 1934 (the "Exchange Act") erence into any filing of the Company under the Securities Act of

#### Item 9.01 Financial Statements and Exhibits.

Exhibit No.	Description
99.1	Cocrystal Pharma, Inc. Corporate Presentation, dated January 13, 2025
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: January 13, 2025

#### Cocrystal Pharma, Inc.

By: /s/ James Martin

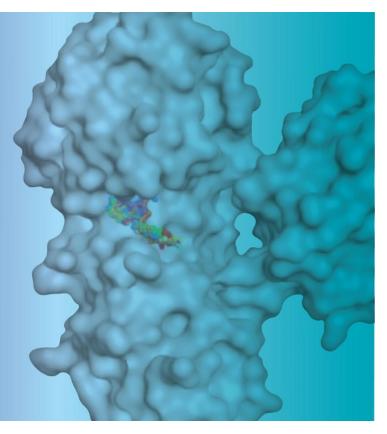
Name: James Martin Title: Chief Financia Chief Financial Officer and Co-Chief Executive Officer



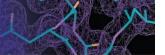
Potent antivirals to combat some of the most serious diseases facing humanity

Investor Presentation January 2025

Nasdaq: COCP 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 development pipeline; our technology platform's ability to produce viable drug candidates at reduced development timelines and costs; development efforts in our clinical programs, including our ongoing Phase 2a study for oral influenza PB2 inhibitor; our ongoing Phase 1 study with 3CL protease inhibitor for coronavirus and norovirus; 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," "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 any future interest rate increases in response to inflation, uncertainty in the financial markets, 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, financial difficulties experienced by certain partners and our ability to secure and maintain new collaboration partners, general risks arising from clinical trials, receipt of regulatory approvals, regulatory changes, development of effective treatments and/or vaccines by competitors, including as part of the programs financed by the U.S. government, 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, 2023. Any forwardlooking 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.

## C CRYSTAL

Applying powerful, proprietary

technology to develop first- and

best-in-class broad-spectrum

drug discovery platform

antiviral drugs

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## Advancing programs in high-value antiviral drug targets

- Influenza
- Norovirus
- · 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

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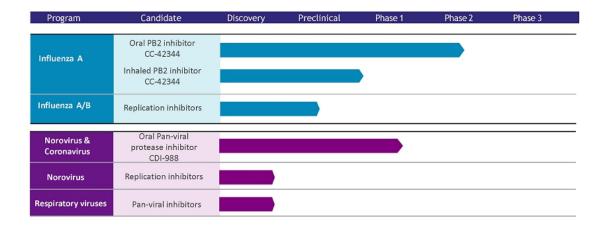
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## **Investment Highlights**

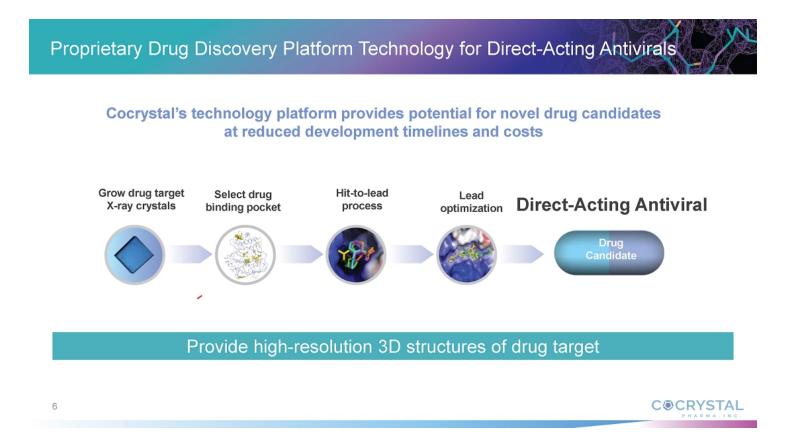
- Targeting multibillion-dollar, global markets for the treatment of acute and pandemic viral diseases
- Proprietary structure-based drug discovery platform technology provides opportunity for discovery and development of novel, broad-spectrum drug candidates
- Advancing multiple clinical programs
  - Oral influenza PB2 inhibitor CC-42344 Phase 2a study continues in 2025
  - · First-in-class dual oral norovirus and coronavirus protease inhibitor CDI-988 Phase 1 study interim data available
- 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



## Multiple clinical assets poised to deliver significant growth

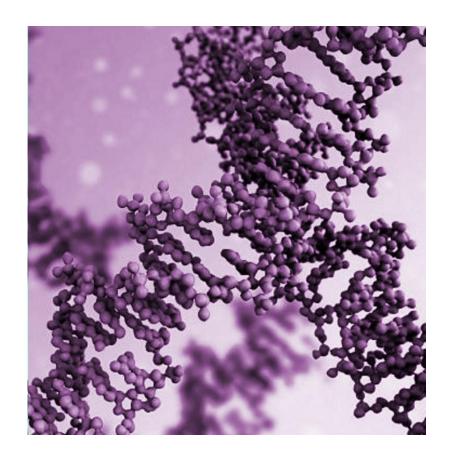


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- Urgent health risks with newly emerging pandemic viral outbreaks<sup>1,2</sup>
  - · Significant delay of effective new antiviral therapeutics and vaccine development
  - Challenging issues with current drug discovery approach one-target/one-drug paradigm
- Significant advantages of Cocrystal's pan-viral drug discovery approach
  - Cocrystal's proprietary structure-based drug design platform technology enables simultaneous drug design on the highly conserved regions of multiple viral drug targets
  - First pan-viral clinical drug candidate CDI-988 developed for the treatment of both norovirus and coronavirus infections
  - Facilitates the rapid development and may allow expedited regulatory pathways (fast track and/or breakthrough designation, and emergency use authorization)

<sup>1</sup> Accelerating antiviral drug discovery: lessons from COVID-19 <u>https://www.nature.com/articles/s41573-023-00692-8</u> <sup>2</sup> The urgent need for pan-antiviral agents: from multitarget discovery to multiscale design <u>https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7682558/</u>



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Pandemic and Seasonal Influenza Program

- 1 billion cases, 3-5 million severe illnesses and up to 650,000 deaths worldwide annually<sup>1</sup>
- Not well managed with currently approved vaccines having only 40-60% effectiveness<sup>2</sup>
- On average ~8% of the U.S. population contracts influenza each season<sup>3</sup>
- Influenza is responsible for ~\$10.4 billion in direct costs for hospitalizations and outpatient visits for adults in the U.S. annually
- Only influenza A causes pandemic flu and is responsible for majority of seasonal influenza infections<sup>1</sup>
- Potential emerging pandemic influenza A strains and drug-resistant strains against approved influenza antivirals, Tamiflu<sup>®</sup> and Xofluza <sup>®</sup>
  - Tamiflu has long history of drug resistance<sup>5</sup>
  - Xofluza has shown emergence of drug resistant mutations<sup>6</sup>

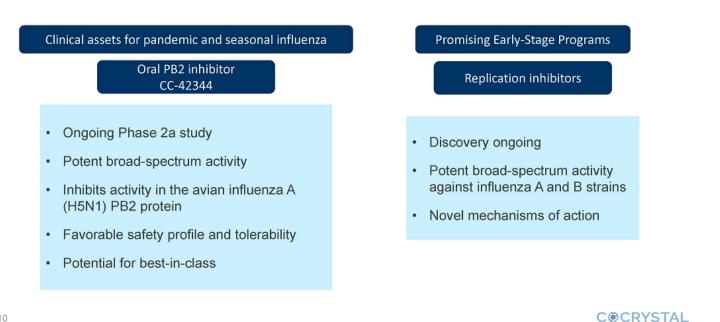
<sup>1</sup> World Health Organization (WHO) (March 2019): <u>https://www.who.int/news-room/fact-sheets/detail/influenza-(seasonal)</u>

<sup>2</sup> Center for Disease Control and Prevention (CDC): Vaccine Effectiveness: How Well Do Flu Vaccines Work?: <u>https://www.cdc.gov/flu/vaccines-work/vaccineeffect.htm</u> <u>3 CDC Seasonal Flu Microsite</u>

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- 4 CDC: Make It Your Business to Fight the Flu
- <sup>5</sup> ScienceDaily (March 2014) Tamiflu-resistant influenza related to mutations in genome: <u>https://www.sciencedaily.com/releases/2014/03/140331114237.htm</u>
- <sup>6</sup> NEJM Journal Watch (September 2018) A Promising Drug for Influenza?: <u>https://www.jwatch.org/na47413/2018/09/12/promising-drug-influenza</u>





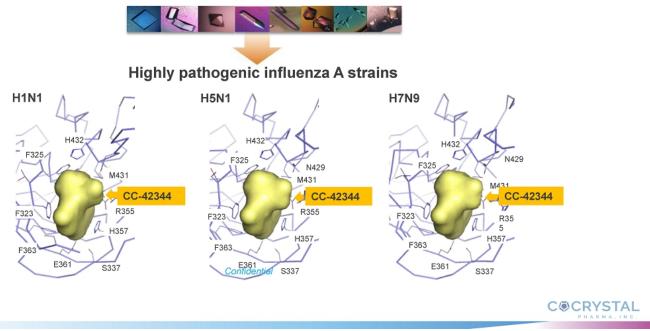
# CC-42344 Shows Broad-Spectrum Antiviral Activity Against Pandemic and Seasonal Influenza A Strains

Influenza serotype	Strain	CC-42344, EC <sub>50</sub> nM
H1N1	A/PR/8/34	1
Pandemic H1N1	California/04/2009	0.5
H1N1	A1/Denver/1/57	3
H1N1	A/Fort Monmouth/1/47	2
H1N1	A/NY/18/09	5
H3N2	A/AICHI/2/68	0.2
Highly pathogenic Avian H5N1	Duck/MN/1524/81	<3.2
Highly pathogenic Avian H5N1	Hong Kong/213/2003	4.5
Highly pathogenic Avian H5N1	Thailand/16/2004	<3.2
Highly pathogenic Avian H7N7	Netherlands/219/2013	5.6
Highly pathogenic Avian H7N9	Anhui/1/2013	<3.2
H1N1- Oseltamivir resistant	A/HK/2369/09 H274Y	9
H3N2-Oseltamivir resistant	A/Wuhan/395/95	0.5
H1N1- Baloxavir resistant (I38T)	A/PR/8/34 I38T	0.5

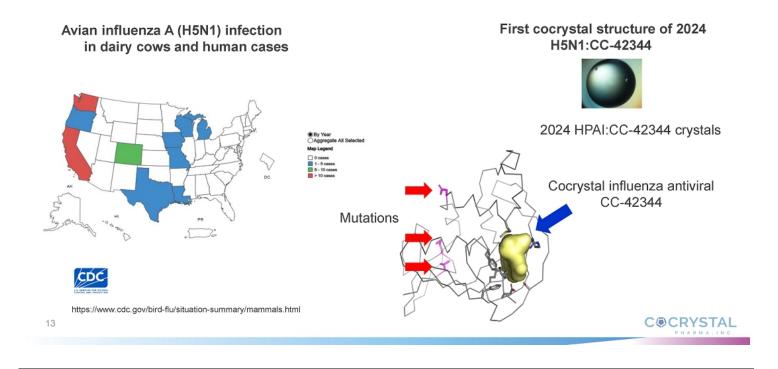
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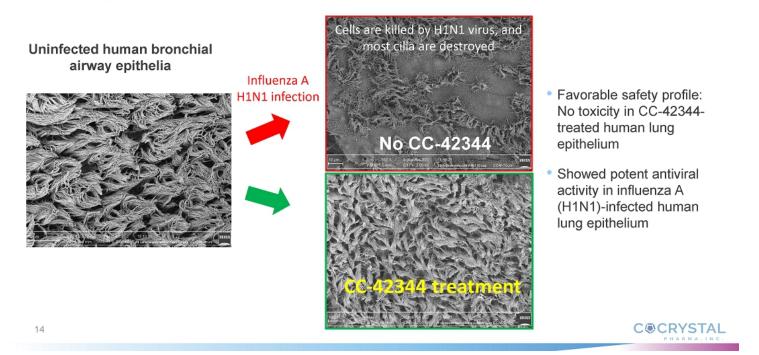
## Cocrystal proprietary drug discovery platform technology



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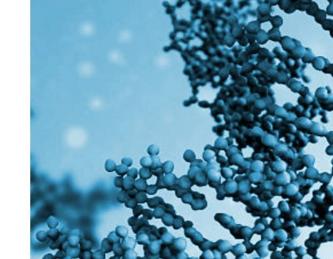


## CC-42344 Shows Potent Antiviral Activity in Influenza-Infected Human Lung Epithilium



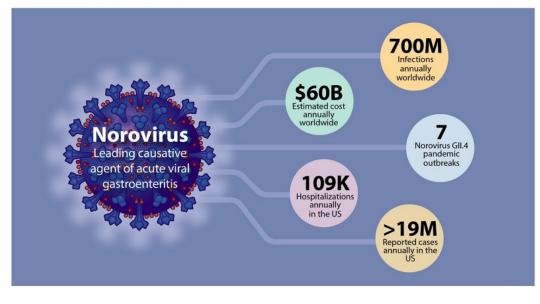
- Favorable safety profile and tolerability
- Potent, broad-spectrum activity against pandemic and seasonal strains
- High barrier to resistance
- Oral CC-42344: Human challenge Phase 2a study ongoing in UK
- Oral CC-42344: FDA feedback provides improved clarity on regulatory path and requirements for oral CC-42344 Phase 2b trial

Norovirus and Coronavirus Program Overview



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Norovirus Infection: No Approved Treatments or Vaccines Available



CDC: Norovirus Disease in the United States https://www.cdc.gov/norovirus/burden.html

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# Significant Unmet Need for New Antivirals to Combat Norovirus Infections

- Highly contagious virus that causes symptoms of acute gastroenteritis including nausea, vomiting, stomach pain and diarrhea
  - Transmissible by infected person, contaminated food or water, or touching contaminated surfaces
- Major cause of gastrointestinal illness in closed and crowded environments including hospitals, nursing homes, childcare facilities, military facilities and cruise ships
- Worldwide norovirus-related healthcare and lost productivity costs of ~\$60 billion dollars
- GII.4 noroviruses have caused the majority of recent norovirus outbreaks worldwide
- No approved antivirals or vaccines for norovirus

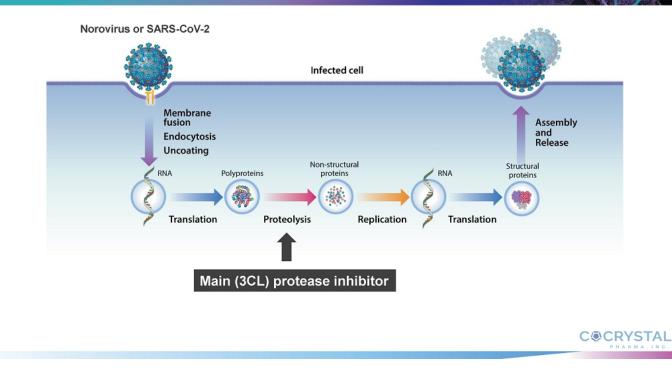
CDC: Norovirus Disease in the United States <u>https://www.cdc.gov/norovirus/burden.html</u> National Foundation for Infections Diseases <u>https://www.nfid.org/infectious-disease/norovirus/</u>





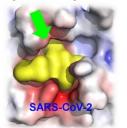
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Cocrystal Viral Protease Inhibitors Block the Essential Replication Process

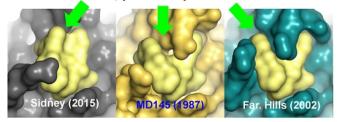


# First-in-Class Pan-Viral Protease Inhibitor CDI-988 for Treatment of Norovirus and COVID infections

## CDI-988, pan-viral protease inhibitor



CDI-988, pan-viral protease inhibitor



Cocrystal structures of Pandemic Norovirus and SARS-CoV-2 with CDI-988

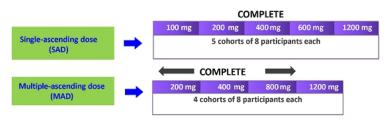
- Developed using Cocrystal's proprietary drug discovery platform technology
- Binds to a highly conserved region required for viral proteases
- Exhibits pan-viral antiviral activity against pandemic norovirus and coronavirus strains
- Demonstrated a favorable safety profile



# Accelerating Progress with First-in-Class Oral Pan-Viral Protease Inhibitor CDI-988: Phase 1 Near Completion

Phase 1 study in healthy participants to evaluate the safety, tolerability and PK of CDI-988

- · Single-center, randomized, double-blind, placebo-controlled
- · Single-ascending dose and multiple-ascending dose cohorts; 7-day nontreatment follow-up period
- Healthy adult volunteers
- Each cohort comprised 8 participants (6 on CDI-988; 2 on placebo)

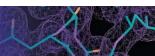


- All dose cohorts well tolerated; No treatment-related study discontinuations
- No serious adverse effects (SAEs)
- Few overall AEs: Most commonly occurring treatment-related AE was headache
- · Next step in clinical development: human challenge study in norovirus-infected subjects

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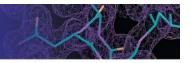
# **Experienced Board of Directors**

Roger Kornberg, Ph.D. Co-founder, Chairman of the Board & Chairman of the Scientific Advisory Board	<ul> <li>Nobel Laureate in Chemistry - the process by which genetic information from DNA is copied to RNA</li> <li>Welch Prize – highest award granted in the field of chemistry in the U.S.</li> <li>Leopald Mayer Prize – highest award granted in the field of biomedical sciences from the French Academy of Sciences</li> </ul>		
Steve Rubin Vice Chairman	<ul> <li>EVP-Administration &amp; Director of OPKO Health, Inc.</li> <li>Former SVP &amp; General Counsel of IVAX Corporation; SVP &amp; General Counsel of Telergy Inc.</li> </ul>		
Phillip Frost, M.D. Director	<ul> <li>Chairman &amp; CEO of OPKO Health, Inc.</li> <li>Former Chairman of Teva Pharmaceuticals; Chairman and CEO of IVAX Corporation – sold for \$7.4 billion</li> <li>Board of Regents of Smithsonian Institution; Board of Trustees of University of Miami; Trustee of Scripps Research Institutes</li> </ul>		
Fred Hassan Director	<ul> <li>Chairman of the investment firm Caret Group; Director of global private equity firm Warburg Pincus LLC</li> <li>Former Chairman &amp; CEO of Schering-Plough – acquired by Merck</li> <li>Former Chairman &amp; CEO of Pharmacia Corporation; senior positions at Wyeth &amp; Sandoz Pharmaceuticals</li> </ul>		
Anthony Japour, M.D. Director	<ul> <li>President, CEO &amp; Director of iTolerance</li> <li>Former CEO of AdvancedDx Biological Laboratories-USA; Medical Director of ICON plc</li> <li>Former with Elite Health Medical Group specializing in infectious diseases</li> </ul>		
Richard C. Pfenniger, Jr. Director	<ul> <li>Director of OPKO Health, GP Strategies Corporation &amp; Asensus Surgical, Inc.</li> <li>Former Chairman, CEO &amp; President of Continucare Corporation; CEO &amp; Vice Chairman of Whitman Education Group.</li> <li>Former COO, SVP-Legal Affairs &amp; General Counsel of IVAX Corporation</li> </ul>		
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# Seasoned Leadership



Management		Scientific Advisory Board	
Sam Lee, Ph.D. Co-Chief Executive Officer & President		Roger Kornberg, Ph.D. Chairman of the Board, Chairman of the Scientific Advisory Board	Professor Stanford University School of Medicine     Nobel Laureate
25+ years of anti-infective drug discovery research experience, including HCV and influenza antivirals; played key role in early development of phosphoinositide 3- kinase (PI3K) delta inhibitor, Zydelig	icos Zydelig	Michael Levitt, Ph.D. Member	Professor Stanford University School of Medicine     Nobel Laureate
		Baek Kim, Ph.D. Member	Director of Center for Drug Discovery Emory University
James J. Martin, MBA, CPA Co-Chief Executive Officer & Chief Financial Officer 25+ years of finance and management experience including providing financial leadership to commercial-stage, publicly traded health science companies	O VBI VACCINES	Bob Lehman, Ph.D. Member	<ul> <li>Professor (Emeritus)</li> <li>Stanford University School of Medicine</li> </ul>
	<i>⊘</i> мотиз <sup>а</sup> "	Gary Schoolnik, M.D.	Professor (Emeritus)     Stanford University School of Medicine
	SciVac	Roland Strong, Ph.D. Member	Professor     Fred Hutchinson Cancer Research Cente
	<u>nims</u>	Christophe Verlinde, Ph.D.	<ul> <li>Professor (Emeritus) University of Washington</li> </ul>

# Expanding Intellectual Property Portfolio



- Issued patents in U.S. and major countries
- Pending U.S. provisional applications

## Pandemic Influenza A

- PB2 (influenza A inhibitor)
  - Pending applications in PCT and Taiwan
  - Pending U.S. provisional applications

## Influenza A/B

- Influenza A/B inhibitor
- · Pending applications in U.S. and worldwide

### Norovirus

- Issued patents in U.S. and major countries
- Pending U.S. provisional applications

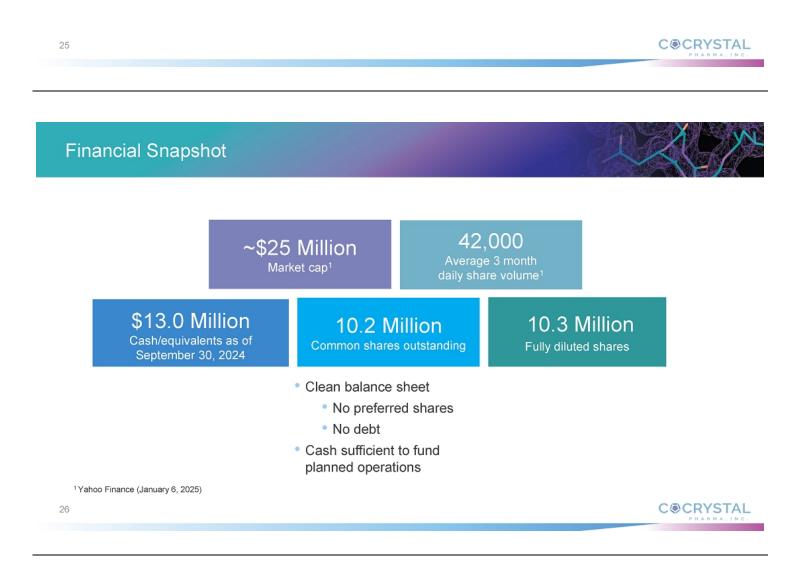
## HCV

NS5B (NNI)

- Issued patents in U.S.
- Pending applications in U.S. and worldwide
- Pending U.S. provisional application



- CDI-988 as oral treatment and prophylaxis for noroviruses, coronaviruses and other viral infections
  - ✓ Interim topline Phase 1 study results
  - Phase 1 results including higher-dose cohort
  - FDA pre-IND meeting
  - FDA IND submission
  - Enrollment initiation in norovirus challenge study
- CC-42344 as an oral treatment of pandemic and seasonal influenza A
  - ✓ Continuation of Phase 2a influenza challenge study



# **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
- Developing multiple discovery programs for respiratory viral diseases
- Exploring pandemic preparedness collaboration opportunities
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