

April 24, 2024



## Corbus Pharmaceuticals Announces Abstract Accepted for Presentation at 2024 ASCO Annual Meeting

NORWOOD, Mass., April 24, 2024 (GLOBE NEWSWIRE) -- Corbus Pharmaceuticals Holdings, Inc. (NASDAQ: CRBP) ("Corbus" or the "Company"), today announced that an abstract providing updated data from the Phase 1, first-in-human clinical data from a dose escalation study being carried out by its partner CSPC in China with CRB-701 (SYS6002) has been accepted for presentation at the [2024 American Society of Clinical Oncology](#) annual meeting, to be held May 31-June 4, 2024 in Chicago, IL. CRB-701 (SYS6002) is a next-generation antibody-drug-conjugate (ADC) targeting Nectin-4 with a third generation, site-specific cleavable linker and a homogenous drug antibody ratio of 2, using MMAE as the payload. Encouraging safety and efficacy data from this trial were presented at ASCO-GU 2024.

The abstract titled *Clinical Update Related to the First-In-Human Trial of SYS6002 (CRB-701), A Next-Generation Nectin-4 Targeting Antibody Drug Conjugate*, will be presented as a poster on June 1, 2024.

Nectin-4 is a clinically validated, tumor-associated antigen in urothelial cancer. The Nectin-4 ADC PADCEV<sup>®</sup> is approved for use in late metastatic urothelial cancer and recently received an expanded label under an accelerated approval from the Food and Drug Administration for use in combination with KEYTRUDA<sup>®</sup> for patients with locally advanced or metastatic urothelial carcinoma who are ineligible for cisplatin-containing chemotherapy.

### About CRB-701

CRB-701 (SYS6002) is a next-generation antibody-drug-conjugate (ADC) targeting Nectin-4, that contains a site-specific, cleavable linker and a homogenous drug antibody ratio of 2, using MMAE as the payload. Nectin-4 is a clinically validated, tumor-associated antigen in urothelial cancer. The Nectin-4 ADC PADCEV<sup>®</sup> (enfortumab vedotin-ejfv) is approved for use in late metastatic urothelial cancer and recently received an expanded label under an accelerated approval from the Food and Drug Administration for use in combination with KEYTRUDA<sup>®</sup> for patients with locally advanced or metastatic urothelial carcinoma who are ineligible for cisplatin-containing chemotherapy.

### About Corbus

Corbus Pharmaceuticals Holdings, Inc. is a precision oncology company with a diversified portfolio and is committed to helping people defeat serious illness by bringing innovative scientific approaches to well understood biological pathways. Corbus' pipeline includes CRB-701, a next generation antibody drug conjugate that targets the expression of Nectin-4

on cancer cells to release a cytotoxic payload, CRB-601, an anti-integrin monoclonal antibody which blocks the activation of TGF $\beta$  expressed on cancer cells, and CRB-913, a highly peripherally restricted CB1 inverse agonist for the treatment of obesity. Corbus is headquartered in Norwood, Massachusetts. For more information on Corbus, visit [corbuspharma.com](http://corbuspharma.com). Connect with us on [Twitter](#), [LinkedIn](#) and [Facebook](#).

### **Forward-Looking Statements**

This press release contains certain forward-looking statements within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934 and Private Securities Litigation Reform Act, as amended, including those relating to the Company's restructuring, trial results, product development, clinical and regulatory timelines, market opportunity, competitive position, possible or assumed future results of operations, business strategies, potential growth opportunities and other statement that are predictive in nature. These forward-looking statements are based on current expectations, estimates, forecasts and projections about the industry and markets in which we operate and management's current beliefs and assumptions.

These statements may be identified by the use of forward-looking expressions, including, but not limited to, "expect," "anticipate," "intend," "plan," "believe," "estimate," "potential," "predict," "project," "should," "would" and similar expressions and the negatives of those terms. These statements relate to future events or our financial performance and involve known and unknown risks, uncertainties, and other factors on our operations, clinical development plans and timelines, which may cause actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. Such factors include those set forth in the Company's filings with the Securities and Exchange Commission. Prospective investors are cautioned not to place undue reliance on such forward-looking statements, which speak only as of the date of this press release. The Company undertakes no obligation to publicly update any forward-looking statement, whether as a result of new information, future events or otherwise.

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### **INVESTOR CONTACT:**

#### **Sean Moran**

*Chief Financial Officer*

Corbus Pharmaceuticals

[smoran@corbuspharma.com](mailto:smoran@corbuspharma.com)

#### **Bruce Mackle**

*Managing Director*

LifeSci Advisors, LLC

[bmackle@lifesciadvisors.com](mailto:bmackle@lifesciadvisors.com)



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