Incyte To Present Initial Data for its TGFβR2×PD-1-directed Bispecific Antibody (INCA33890) and its Selective Inhibitor of G12D-mutated KRAS (INCB161734) at the European Society of Medical Oncology (ESMO) Congress 2025

July 24, 2025

- Mini oral presentation to highlight initial Phase 1 data of INCA33890, a promising TGFβR2×PD-1-directed bispecific antibody, in patients with advanced or metastatic solid tumors

- Oral presentation to feature initial data of INCB161734, a novel, selective and orally bioavailable inhibitor of G12D-mutated KRAS, in patients with advanced or metastatic solid tumors with KRASG12D mutations

WILMINGTON, Del.--(BUSINESS WIRE)--Jul. 24, 2025-- Incyte (Nasdaq:INCY) today announced that the Company will present key data from its oncology portfolio at the upcoming European Society of Medical Oncology (ESMO) Congress 2025, to be held October 17-21 in Berlin.

“We're looking forward to presenting the latest findings across our oncology portfolio at this year’s congress, including initial data for our TGFβR2×PD-1-directed bispecific antibody, INCA33890, and our novel, selective and orally bioavailable inhibitor of G12D-mutated KRAS, INCB161734,” said Pablo J. Cagnoni, M.D., President and Head of Research and Development, Incyte. “These data underscore the importance of establishing treatment options for advanced solid tumors and speak to the potential of INCA33890 and INCB161734 as personalized therapies to enhance outcomes for patients with cancer across a range of tumor types.”

Details on the abstracts accepted for oral presentation at ESMO include:

INCB161734 (KRAS G12D)

Preliminary Phase 1 Results Of INCB161734, A Novel Oral KRAS G12D Inhibitor, In Patients With Advanced Or Metastatic Solid Tumors
(Session Title: Proffered paper session: Developmental therapeutics. [October 19, 8:45 – 10:15 a.m. ET [2:45 – 4:15 p.m. CEST]. Abstract #916O.)

INCA33890 (PD-1/TGFβR2)

A Phase 1 Study Of INCA33890, A PD-1/Tgfβr2 Bispecific Antibody, For Advanced Solid Tumours
(Session Title: Mini oral session: Investigational immunotherapy. [October 17, 8:00 – 9:30 a.m. ET [2:00 – 3:30 p.m. CEST]. Abstract #1522.)

INCAGN2385 (LAG-3)

Retifanlimab (Anti–PD-1 Mab) Alone Or In Combination With Anti-LAG3 ± Anti-TIM3 Mabs In Previously Untreated, Recurrent And/Or Metastatic (R/M) PD-L1+ HNSCC: A Double-Blind Randomised Controlled Phase 2 Trial
(Session Title: Mini oral session: Head and neck cancer. [October 19, 10:30 a.m. – 12:00 p.m. ET [4:30 – 6:00 p.m. CEST]. Abstract #1325.)

All regular abstracts accepted for presentation at the ESMO Congress 2025 will be published online via the ESMO website on Sunday, October 12 at 6:05 p.m. ET (12:05 a.m. CEST). All accepted abstracts will be published online in the ESMO Congress 2025 Abstract Book, a supplement to the official ESMO journal, Annals of Oncology.

More information regarding the 2025 ESMO Congress can be found at: https://www.esmo.org/meeting-calendar/esmo-congress-2025.

About Incyte
A global biopharmaceutical company on a mission to *Solve On.*, Incyte follows the science to find solutions for patients with unmet medical needs. Through the discovery, development and commercialization of proprietary therapeutics, Incyte has established a portfolio of first-in-class medicines for patients and a strong pipeline of products in Oncology and Inflammation & Autoimmunity. Headquartered in Wilmington, Delaware, Incyte has operations in North America, Europe and Asia.

For additional information on Incyte, please visit Incyte.com or follow us on social media: LinkedIn, X, Instagram, Facebook, YouTube.

Incyte Forward-Looking Statements
Except for the historical information set forth herein, the matters set forth in this press release, including statements regarding the presentation of data from Incyte’s clinical development pipeline, the potential offered by INCA33890 and INCB161734, whether or when any development compounds or combinations will be approved or commercially available for use in humans anywhere in the world outside of the already approved indications in specific regions, and Incyte’s goal of improving the lives of patients, contain predictions, estimates and other forward-looking statements.

These forward-looking statements are based on Incyte’s current expectations and subject to risks and uncertainties that may cause actual results to differ materially, including unanticipated developments in and risks related to: unanticipated delays; further research and development and the results of clinical trials possibly being unsuccessful or insufficient to meet applicable regulatory standards or warrant continued development; the ability to enroll sufficient numbers of subjects in clinical trials; determinations made by the FDA, EMA, and other regulatory authorities; the efficacy or safety of Incyte and its partners’ products; the acceptance of Incyte and its partners’ products in the marketplace; market competition; sales, marketing, manufacturing and distribution requirements; and other risks detailed from time to time in our reports filed with the U.S. Securities and Exchange Commission, including our annual report on Form 10-K and our quarterly report on Form 10-K for the year ended December 31, 2024. Incyte disclaims any intent or obligation to update these forward-looking statements.

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