

Agenus to Present First Phase 2 Botensilimab Data in Advanced Cutaneous Melanoma at ASCO 2026

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- Four abstracts accepted across melanoma, colorectal cancer, and translational research

LEXINGTON, Mass.--(BUSINESS WIRE)-- **Agenus Inc.** (Nasdaq: AGEN), a leader in immuno-oncology innovation, today announced that four abstracts highlighting botensilimab (BOT), alone or in combination with balstilimab (BAL), have been accepted for presentation at the 2026 American Society of Clinical Oncology (ASCO) Annual Meeting, taking place May 29–June 2, 2026, in Chicago, Illinois.

The accepted abstracts reflect continued progress of Agenus' botensilimab and balstilimab immunotherapy program across colorectal cancer, melanoma, and translational research. They include the first presentation of Phase 2 clinical data for BOT with or without BAL in advanced cutaneous melanoma refractory or resistant to anti-PD-(L)1 therapy, with or without prior CTLA-4 inhibition, as well as a translational biomarker abstract evaluating an artificial intelligence foundation model in solid tumors and two trials-in-progress posters in colorectal cancer.

"Our presence at ASCO reflects strong momentum across the botensilimab and balstilimab program. From the first Phase 2 melanoma presentation to important colorectal cancer trial updates and innovative translational biomarker research, these presentations underscore our focus on delivering new immunotherapy options to patients who need them most," said Garo H. Armen, PhD, Chairman and CEO of Agenus.

Presentation Details:

1. Abstract Title: Botensilimab (BOT) ± balstilimab (BAL) in patients (pts) with advanced cutaneous melanoma (cMEL) refractory/resistant (R/R) to anti-PD-(L)1 ± CTLA-4: A phase 2 trial
Abstract No.: 9543

Presenter: Michael Atkins MD; Georgetown Lombardi Comprehensive Cancer Center, Georgetown University Medical Center

Session Title: Poster Session – Melanoma/Skin Cancers

Location: Hall A – Posters and Exhibits

Poster Board: 259

Date/Time: May 31, 2026, 9:00 AM–12:00 PM CDT

2. Abstract Title: Artificial intelligence (AI) foundation model as a predictor of efficacy of next-generation checkpoint inhibition with botensilimab (BOT) + balstilimab (BAL) in solid tumors using pretreatment H&E images

Abstract No.: 2535

Presenter: Ryan Dalton, Noetik

Session Title: Poster Session – Developmental Therapeutics—Immunotherapy

Location: Hall A – Posters and Exhibits

Poster Board: 325

Date/Time: May 30, 2026, 1:30 PM–4:30 PM CDT

3. Abstract Title: The CO.33/BATTMAN trial: A phase 3 randomized study of botensilimab + balstilimab versus best supportive care in chemo refractory unresectable colorectal adenocarcinoma that is not dMMR/MSI-H

Abstract No.: TPS3676

Presenter: Jonathan Loree MD; BC Cancer; Canadian Cancer Trials Group, Queen's University

Session Title: Poster Session – Gastrointestinal Cancer—Colorectal and Anal

Location: Hall A – Posters and Exhibits

Poster Board: 441a

Date/Time: May 30, 2026, 9:00 AM–12:00 PM CDT

4. Abstract Title: Phase 2 study of adjuvant botensilimab in combination with balstilimab in patients with microsatellite-stable colorectal cancer and persistent circulating tumor DNA following surgery and chemotherapy

Abstract No.: TPS3689

Presenter: Neil Segal, MD; Memorial Sloan Kettering Cancer Center

Session Title: Poster Session – Gastrointestinal Cancer—Colorectal and Anal

Location: Hall A – Posters and Exhibits

Poster Board: 447b

Date/Time: May 30, 2026, 9:00 AM–12:00 PM CDT

About Agenus

Agenus is a leading immuno-oncology company targeting cancer with a comprehensive pipeline of immunological

agents. The company was founded in 1994 with a mission to expand patient populations benefiting from cancer immunotherapy through combination approaches, using a broad repertoire of antibody therapeutics, adoptive cell therapies (through MiNK Therapeutics) and adjuvants. Agenus has robust end-to-end development capabilities, across commercial and clinical cGMP manufacturing facilities, research and discovery, and a global clinical operations footprint. Agenus is headquartered in Lexington, MA. For more information, visit www.agenusbio.com or @agenus_bio. Information that may be important to investors will be routinely posted on our website and social media channels.

About Botensilimab (BOT)

Botensilimab (BOT) is a human Fc enhanced multifunctional anti-CTLA-4 antibody designed to boost both innate and adaptive anti-tumor immune responses. Its novel design leverages mechanisms of action to extend immunotherapy benefits to “cold” tumors which generally respond poorly to standard of care or are refractory to conventional PD-1/CTLA-4 therapies and investigational therapies. Botensilimab augments immune responses across a wide range of tumor types by priming and activating T cells, downregulating intratumoral regulatory T cells, activating myeloid cells and inducing long-term memory responses.

Approximately 1,200 patients have been treated with botensilimab and/or balstilimab in phase 1 and phase 2 clinical trials. Botensilimab alone, or in combination with Agenus’ investigational PD-1 antibody, balstilimab, has shown clinical responses across nine metastatic, late-line cancers. For more information about botensilimab trials, visit www.clinicaltrials.gov.

About Balstilimab (BAL)

Balstilimab is a novel, fully human monoclonal immunoglobulin G4 (IgG4) designed to block PD-1 (programmed cell death protein 1) from interacting with its ligands PD-L1 and PD-L2. It has been evaluated in more than 900 patients to date and has demonstrated clinical activity and a favorable tolerability profile in several tumor types.

Forward-Looking Statements

This press release contains forward-looking statements that are made pursuant to the safe harbor provisions of the federal securities laws, including statements regarding its botensilimab and balstilimab programs, expected regulatory timelines and filings, and any other statements containing the words “may,” “believes,” “expects,” “anticipates,” “hopes,” “intends,” “plans,” “forecasts,” “estimates,” “will,” “establish,” “potential,” “superiority,” “best in class,” and similar expressions are intended to identify forward-looking statements. These forward-looking statements are subject to risks and uncertainties that could cause actual results to differ materially. These risks and uncertainties include, among others, the factors described under the Risk Factors section of our most recent

Annual Report on Form 10-K for 2025, and subsequent Quarterly Reports on Form 10-Q filed with the Securities and Exchange Commission. Agenus cautions investors not to place considerable reliance on the forward-looking statements contained in this release. These statements speak only as of the date of this press release, and Agenus undertakes no obligation to update or revise the statements, other than to the extent required by law. All forward-looking statements are expressly qualified in their entirety by this cautionary statement.

Investors

917-362-1370 | investor@agenusbio.com

Media

781-674-4422 | communications@agenusbio.com

Source: Agenus Inc.