Erasca Announces Three Presentations at the 2024 ASCO Annual Meeting

04.24.2024

Oral presentation will feature updated HERKULES-3 results for ERK1/2 inhibitor ERAS-007 in combination with encorafenib and cetuximab in BRAFm colorectal cancer

Poster presentations will feature SEACRAFT-1 Phase 1 trial design for naporafenib plus trametinib in RAS Q61X mutations, and preliminary FLAGSHP-1 Phase 1 results for ERAS-601 and cetuximab in chordoma

SAN DIEGO, April 24, 2024 (GLOBE NEWSWIRE) -- Erasca, Inc. (Nasdaq: ERAS), a clinical-stage precision oncology company singularly focused on discovering, developing, and commercializing therapies for patients with RAS/MAPK pathway-driven cancers, today announced the company will present one oral presentation and two poster presentations at the upcoming American Society of Clinical Oncology (ASCO) Annual Meeting, taking place May 31-June 4, 2024, in Chicago, Illinois.

The abstracts will be available on the ASCO meeting website. The posters will be available online at Erasca.com/events/presentations following the presentations.

Presentation Details

- **Abstract 3517 - Updated results from ERAS-007 plus encorafenib and cetuximab (EC) in patients (pts) with EC-naïve metastatic BRAF V600E colorectal cancer (CRC) in the phase 1b/2 HERKULES-3 study**
  Date/Time: Monday, June 3, 2024, 1:15 PM-2:45 PM CDT
  Type: Rapid Oral Abstract
  Session: Gastrointestinal Cancer—Colorectal and Anal
  Presenter: Dr. Aparna Parikh, Massachusetts General Hospital
  Location: McCormick Place Convention Center, Arie Crown Theater

- **Abstract TPS3178 - An open-label study to assess the safety and efficacy of naporafenib (ERAS-254) administered with trametinib in previously treated patients with locally advanced unresectable or metastatic solid tumor malignancies with RAS Q61X mutations (SEACRAFT-1)**
  Date/Time: Saturday, June 1, 2024, 9:00 AM-12:00 PM CDT
  Type: Poster Session
  Session: Developmental Therapeutics—Molecularly Targeted Agents and Tumor Biology
  Presenter: Dr. Alex Spira, Next Oncology Virginia
  Location: McCormick Place Convention Center, Exhibit Hall/Hall A, Poster Board 314a

- **Abstract 11522 – Preliminary results from FLAGSHP-1: A phase 1 study of ERAS-601 as a monotherapy or in combination with cetuximab in patients (pts) previously treated for advanced chordoma**
  Date/Time: Saturday, June 1, 2024, 1:30 PM-4:30 PM CDT
  Type: Poster Session
  Session: Sarcoma
  Presenter: Dr. Mrinal Gounder, Memorial Sloan Kettering Cancer Center
  Location: McCormick Place Convention Center, Exhibit Hall/Hall A, Poster Board 448

About Erasca
At Erasca, our name is our mission: To erase cancer. We are a clinical-stage precision oncology company singularly focused on discovering, developing, and commercializing therapies for patients with RAS/MAPK pathway-driven cancers. Our company was co-founded by leading pioneers in precision oncology and RAS targeting to create novel therapies and combination regimens designed to comprehensively shut down the RAS/MAPK pathway for the treatment of cancer. We have assembled one of the deepest RAS/MAPK pathway-focused pipeline in the industry. We believe our team's capabilities and experience, further guided by our scientific advisory board which includes the world's leading experts in the RAS/MAPK pathway, uniquely position us to achieve our bold mission of erasing cancer.

Cautionary Note Regarding Forward-Looking Statements
Erasca cautions you that statements contained in this press release regarding matters that are not historical facts are forward-looking statements. The forward-looking statements are based on our current beliefs and expectations and include, but are not limited to: our expectations regarding the potential therapeutic benefits of our product candidates, including ERAS-007, naporafenib, and ERAS-601. Actual results may differ from those set forth in this press release due to the risks and uncertainties inherent in our business, including, without limitation: our approach to the discovery and development of product candidates based on our singular focus on shutting down the RAS/MAPK pathway, a novel and unproven approach; preliminary results of clinical trials are not necessarily indicative of final results and one or more of the clinical outcomes may materially change as patient enrollment continues, following more comprehensive reviews of the data and more patient data become available; our assumptions around

which programs may have a higher probability of success may not be accurate, and we may expend our limited resources to pursue a particular product candidate and/or indication and fail to capitalize on product candidates or indications with greater development or commercial potential; potential delays in the commencement, enrollment, and completion of clinical trials and preclinical studies; our dependence on third parties in connection with manufacturing, research, and preclinical and clinical testing; unexpected adverse side effects or inadequate efficacy of our product candidates that may limit their development, regulatory approval, and/or commercialization, or may result in recalls or product liability claims; unfavorable results from preclinical studies or clinical trials; the inability to realize any benefits from our current licenses, acquisitions, and collaborations, and any future licenses, acquisitions, or collaborations, and our ability to fulfill our obligations under such arrangements; regulatory developments in the United States and foreign countries; later developments with the FDA or EU health authorities may be inconsistent with the feedback received to date regarding our development plans and trial designs; FTD may not lead to a faster development or regulatory review or approval process, and does not increase the likelihood that our product candidates will receive marketing approval; our ability to fund our operating plans with our current cash, cash equivalents, and marketable securities; and other risks described in our prior filings with the Securities and Exchange Commission (SEC), including under the heading “Risk Factors” in our annual report on Form 10-K for the year ended December 31, 2023, and any subsequent filings with the SEC. You are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date hereof, and we undertake no obligation to update such statements to reflect events that occur or circumstances that exist after the date hereof. All forward-looking statements are qualified in their entirety by this cautionary statement, which is made under the safe harbor provisions of the Private Securities Litigation Reform Act of 1995.

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Source: Erasca, Inc.

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