## NantHealth's Eviti Connect Data Shows Recycling of Chemotherapy and Biologics for Advanced CRC Patients More Common Than Switching to Drug Regimens with Alternative Mechanism of Action

January 25, 2020

NantHealth Presented These Findings at the ASCO 2020 Gastrointestinal Cancers Symposium

SAN FRANCISCO--(BUSINESS WIRE)-- NantHealth, Inc. (NASDAQ: NH), a next-generation, evidence-based, personalized healthcare company, today announced new real-world data on treatment patterns for patients with advanced colorectal cancer (CRC) during a poster session at the 2020 Gastrointestinal Cancer Symposium sponsored by the American Society of Clinical Oncology (ASCO).

The Symposium, held at the Moscone West Building in San Francisco, CA from January 23-25, provides evidence-based teaching methods and cutting-edge learning science to a diverse audience of leaders in oncology education, doctors and care teams.

NantHealth's presentation examined therapeutic preferences and treatment patterns among advanced CRC patients using data from NantHealth's Eviti Connect, an evidence-based treatment intelligence and web-based oncology decision support platform. Detailed information from 6,325 treatment plans was analyzed to identify treatment patterns using regorafenib and trifluridine + tipiracil for advanced CRC patients as third-line of therapy. National Comprehensive Cancer Network (NCCN) guidelines state that regorafenib and trifluridine+tipiracil are both treatment options for patients who have progressed through all available regimens.

Across all 6,325 treatment plans submitted for this patient population, regorafenib (n=217) or trifluridine+tipiracil (n=144) was the submitted treatment in 361 (5.5%) of the treatment plans, making them the 9<sup>th</sup> and 13<sup>th</sup> most frequently requested drugs (excluding growth factors, antiemetics, and leucovorin) in this setting. While the total number of treatment plans for regorafenib was higher than that for trifluridine+tipiracil, the submission of trifluridine+tipiracil has increased over time, consistent with the latter drug's more recent introduction into the market.

"Our analysis shows that recycling of chemotherapy and biologics in the late line setting is common and occurs more frequently than switching to a drug regimen with an alternative mechanism of action," said William A. Flood, MD, MS, Chief Medical Officer for Eviti, NantHealth. "As results cannot be fully explained by clinical trial outcome differences, NCCN guidelines preferences, or HEOR measures, we must delve deeper into why these therapeutic patterns exist to further our mission of optimizing patient outcomes and enabling value-based care in oncology. The uncertainty of what constitutes the 'best' treatment for this patient population provides an excellent opportunity to employ available data to guide patient-centered decision making and value-based care initiatives."

**Title**: "Real world data on treatment patterns of advanced CRC in 3<sup>rd</sup> line and beyond"

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**Abstract** #56, **Poster Session and Number**: C – Anal and Colorectal Cancer

Location: Moscone West Building, Level 1, West Hall

Date and Time: January 25, 2020, 6:30-7:55 AM PT and 12:15-1:45 PM PT

"Our Eviti platform enables access to near-real time data on physician behavior which can provide unique and critical information to pharma, payers, and provider networks to optimize treatment strategies," said Sandeep "Bobby" Reddy, MD, Chief Medical Officer, NantHealth.

## **About NantHealth:**

NantHealth, a member of the NantWorks ecosystem of companies, provides leading solutions across the continuum of care for physicians, payors, patients and biopharmaceutical organizations. NantHealth enables the use of cutting-edge data and technology toward the goals of empowering clinical decision support and improving patient outcomes. NantHealth's comprehensive product portfolio combines the latest technology in payor/provider platforms that exchange information in near-real time (NaviNet and Eviti), connected care solutions that deliver medical device interoperability (DCX device connectivity platform and VCX patient vitals software) and molecular profiling services that combine comprehensive DNA & RNA tumor-normal profiling with pharmacogenomics analysis (GPS Cancer®). For more information, please visit www.nanthealth.com or follow us on Twitter, Facebook and LinkedIn.

## Forward-Looking Statements: NantHealth

This news release contains certain statements of a forward-looking nature relating to future events or future business performance. Forward-looking statements can be identified by the words "expects," "anticipates," "believes," "intends," "estimates," "plans," "will," "outlook" and similar expressions. Forward-looking statements are based on management's current plans, estimates, assumptions and projections, and speak only as of the date they are made. Risks and uncertainties include, but are not limited to: our ability to successfully integrate a complex learning system to address a wide range of healthcare issues; our ability to successfully amass the requisite data to achieve maximum network effects; appropriately allocating financial and human resources across a broad array of product and service offerings; raising additional capital as necessary to fund our operations; achieving significant commercial market acceptance for our sequencing and molecular analysis solutions; establish relationships with, key thought leaders or payers' key decision makers in order to establish GPS Cancer as a standard of care for patients with cancer; our ability to grow the market for our Systems Infrastructure, and applications; successfully enhancing our Systems Infrastructure and applications to achieve market acceptance and keep pace with technological developments; customer concentration; competition; security breaches; bandwidth limitations; our ability to continue our relationship with NantOmics; our ability to obtain regulatory approvals; dependence upon senior management; the need to comply with and meet applicable laws and regulations; unexpected adverse events; clinical adoption and market acceptance of GPS Cancer; and anticipated cost savings. We undertake no obligation to update any forward-looking statement in light of new information or future events, except as otherwise required by law. Forward-looking statements involve inherent risks and uncertainties, most of which are difficult to predict and are generally beyond our control. Actual results or outcomes may differ materially from those implied by the forward-looking statements as a result of the impact of a number of factors, many of which are discussed in more detail in our reports filed with the Securities and Exchange Commission.

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