ASCOrs TAPUR Study Now Has More Than One Hundred Participants Receiving Treatment Drugs and Twenty-Five New Clinical Trial Sites

Next Planned Study Expansion is to Pediatric Population through Lowered Eligibility Age

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ALEXANDRIA, Va. – More than 100 participants are now enrolled on study drug in the American Society of Clinical Oncology's (ASCO®) Targeted Agent and Profiling Utilization Registry (TAPUR) Study. The trial launched in March 2016 at 37 sites and this month welcomes its newest participating clinical sites: Cancer Treatment Centers of America®, Intermountain Healthcare, Sanford Health, Swedish Cancer Institute and Providence Health & Services, and University of Nebraska Medical Center for a total of 63 participating sites in 15 states.

“We are gratified with the high degree of interest and participation in the TAPUR Study. Clinical sites and patients are clearly eager to be a part of this study and are enrolling more quickly than we had anticipated,” said ASCO Chief Medical Officer Richard L. Schilsky, MD, FACP, FASCO. “There’s an urgent need to explore new therapeutic options for patients with advanced solid tumor cancers, multiple myeloma, or B-cell non-Hodgkin lymphoma. TAPUR provides a unique venue to enable multi-stakeholder collaboration to learn about additional uses of targeted drugs that already have Food and Drug Administration approval in one or more cancers. The rapid enrollment in TAPUR will allow us to identify new signals of drug activity as quickly as possible.”

In the eight months since the trial opened, 174 individuals have consented to participate, and
105 participants have begun receiving a TAPUR Study drug. The five centers that have most recently begun patient recruitment will offer TAPUR at multiple locations, for a total of 25 new site locations, which will be fully up and running over the next few months. They are:

- Cancer Treatment Centers of America: Atlanta, Chicago, Philadelphia, Phoenix, and Tulsa
- Intermountain Healthcare: Intermountain Precision Genomics Cancer Research Clinic, Dixie Regional Medical Center-400 East Campus, Dixie Regional Medical Center-River Road Campus, and Intermountain Cancer Center, Cedar City
- Sanford Health: Sioux Falls, Bismarck, and Fargo
- Swedish Cancer Institute and Providence Health & Services: Pacific Gynecology Specialists, First Hill, Issaquah, Ballard, Edmonds, Neuroscience Institute, Eastside, Westside, Southeast, Newberg, and Willamette Falls
- University of Nebraska Medical Center: Main Campus and Village Point

Seven pharmaceutical companies are currently participating—AstraZeneca, Bayer, Bristol-Myers Squibb, Eli Lilly and Company, Genentech, Merck, and Pfizer. Among them, they provide 17 drugs that yield 15 different targeted therapy options (since some of the drugs are used in combination). ASCO has designed the TAPUR Study to facilitate the inclusion of additional drugs and companies as the trial continues and expects to offer additional regimens in the near future.

In addition, study leaders are planning to decrease the enrollment age from 18 to 12 years to extend the opportunity for participation to adolescent patients with advanced cancer, where there is a defined adolescent dose for the study drugs.

The TAPUR Study is designed to evaluate molecularly targeted cancer drugs and collect data on clinical outcomes to learn about additional uses of these drugs outside of indications already approved by the Food and Drug Administration. ASCO is using Syapse Precision Medicine Platform to automate the matching of patients to treatments based on genomic and clinical profiles, facilitate study workflows, and integrate clinical, genomic, treatment, and outcomes data.

ASCO is working with the Research Advocacy Network to conduct a TAPUR sub-study that will provide insights to help the oncology community understand how tumor genomic testing is being used by clinical oncologists and how to provide assistance with provider and patient education. ASCO is also collaborating with the Netherlands Center for Personalized Cancer Treatment on its Drug Rediscovery Protocol (DRUP) trial, which is very similar to TAPUR and
also opened earlier this year. Other international centers and research networks have expressed interest in developing versions of the TAPUR and DRUP protocols in their regions of the world. Doing so has the potential to create a global network to discover new uses of already approved targeted cancer treatments.

The TAPUR Study is registered on ClinicalTrials.gov (NCT 02693535), which includes a full list of inclusion/exclusion criteria and other information. Patients can find study information such as general eligibility criteria, participating clinical sites, and contact information for the study team at www.TAPUR.org. Researchers and practices interested in participating can fill out the TAPUR Study Site Questionnaire or contact the study team at: http://www.tapur.org/contact-us.

Watch a video that provides an overview of the TAPUR study.

**About ASCO:**

Founded in 1964, the American Society of Clinical Oncology, Inc. (ASCO®) is committed to making a world of difference in cancer care. As the world's leading organization of its kind, ASCO represents more than 40,000 oncology professionals who care for people living with cancer. Through research, education, and promotion of the highest-quality patient care, ASCO works to conquer cancer and create a world where cancer is prevented or cured, and every survivor is healthy. ASCO is supported by its affiliate organization, the Conquer Cancer Foundation. Learn more at www.ASCO.org, explore patient education resources at www.Cancer.Net, and follow us on Facebook, Twitter, LinkedIn, and YouTube.