Provectus Biopharmaceuticals to Present at 2015 BIO International Convention in Philadelphia

CFO and COO Peter Culpepper Presents Tuesday, June 16, at 10:45 am Eastern Time
The 2015 BIO International Convention Runs June 15-18

Thursday June 11, 2015

KNOXVILLE, Tenn.--(BUSINESS WIRE)--Provectus Biopharmaceuticals, Inc. (NYSE MKT: PVCT, www.pvct.com), a clinical-stage oncology and dermatology biopharmaceutical company ("Provectus" or the "Company"), announced today that it will present at the 2015 BIO International Convention on Tuesday, June 16, at 10:45 am Eastern.

Peter Culpepper, CFO and COO of Provectus, is scheduled to present in Theater 4, "The Incredible Helix." He will discuss the latest developments at Provectus, including the recently begun phase 3 clinical trial of PV-10, the Company's novel investigational drug for cancer, as a treatment for melanoma.

The 2015 BIO International Convention runs from June 15-18, 2015, at the Pennsylvania Convention Center in Philadelphia, Pennsylvania.

2015 BIO International Convention

In 2015, the BIO International Convention is headed to the heart of the U.S. biopharma industry. In close proximity to New York's financial markets and Washington, D.C.'s regulatory center, the Philadelphia Metropolitan Area is home to more than 1,200 companies, ranging from the biopharma industry's largest multinational companies to its fastest growing firms. The region's vast array of leading universities and research institutions fosters a collaborative environment. In the past 10 years alone, the region has drawn nearly $4 billion in venture capital funding (according to the Jones Lang LaSalle Life Sciences Cluster Report 2012).

About Provectus Biopharmaceuticals, Inc.

Provectus Biopharmaceuticals, Inc. specializes in developing oncology and dermatology therapies. PV-10, its novel investigational drug for cancer, is designed for injection into solid tumors (intralesional administration), thereby reducing potential for systemic side effects. Its oncology focus is on melanoma, breast cancer and cancers of the liver. The Company has received orphan drug designations from the FDA for its melanoma and hepatocellular carcinoma indications. PH-10, its topical investigational drug for dermatology, is undergoing clinical testing for psoriasis and atopic dermatitis. Provectus has completed phase 2 trials of PV-10 as a therapy for metastatic melanoma, and of PH-10 as a topical treatment for atopic dermatitis and psoriasis. Information about these and the Company's other clinical trials, including its current phase 3 study in melanoma, can be found at the NIH registry, www.clinicaltrials.gov. For additional information about Provectus, please visit the Company's website at www.pvct.com or contact Porter, LeVay & Rose, Inc.

FORWARD-LOOKING STATEMENTS: This release contains "forward-looking statements" as defined under U.S. federal securities laws. These statements reflect management's current knowledge, assumptions, beliefs, estimates, and expectations and express management's current views of future performance, results, and trends and may be identified by their use of terms such as "anticipate," "believe," "could," "estimate," "expect," "intend," "may," "plan," "predict," "project," "will," and other similar terms. Forward-looking statements are subject to a number of risks and uncertainties that could cause our actual results to materially differ from those described in the forward-looking statements. Readers should not place undue reliance on forward-looking statements. Such statements are made as of the date hereof, and we undertake no obligation to update such statements after this date.

Risks and uncertainties that could cause our actual results to materially differ from those described in forward-looking statements include those discussed in our filings with the Securities and Exchange Commission (including those described in Item 1A of our Annual Report on Form 10-K for the year ended December 31, 2014) and the following:

- our determination, based on guidance from the FDA, whether to proceed with or without a partner with the fully enrolled phase 3 trial of PV-10 to treat locally advanced cutaneous melanoma and the costs associated with such a trial if it is necessary to complete (versus interim data alone);
- our determination whether to license PV-10, our melanoma drug product candidate, and other solid tumors such as cancers of the liver, if such licensure is appropriate considering the timing and structure of such a license, or to commercialize PV-10 on our own to treat melanoma and other solid tumors such as cancers of the liver;
- our ability to license our dermatology drug product candidate, PH-10, on the basis of our phase 2 atopic dermatitis and psoriasis results, which are in the process of being further developed in conjunction with mechanism of action studies; and
- our ability to raise additional capital if we determine to commercialize PV-10 and/or PH-10 on our own, although our expectation is to be acquired by a prospective pharmaceutical or biotech concern prior to commercialization.

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