



Contact:  
Provectus Biopharmaceuticals, Inc.  
Peter R. Culpepper, Interim CEO, COO  
Phone: 866-594-5999 #30

Porter, LeVay & Rose, Inc.  
Marlon Nurse, DM, SVP – Investor Relations  
Phone: 212-564-4700  
Allison + Partners  
Todd Aydelotte, Managing Director –  
Media Relations  
Phone: 646-428-0644

**FOR IMMEDIATE RELEASE**

**PROVECTUS BIOPHARMACEUTICALS WILL PRESENT AT 2016 BIO  
INTERNATIONAL CONVENTION IN SAN FRANCISCO, CA**

**Presentation Scheduled for 2:45 pm PDT, Wednesday, June 8, 2016,  
Theater 2 of Moscone Center**

**BIO International Convention Runs June 6-9, 2016**

KNOXVILLE, TN, June 7, 2016 --Provectus Biopharmaceuticals, Inc. (NYSE MKT: PVCT, [www.provectusbio.com](http://www.provectusbio.com)), a clinical-stage oncology and dermatology biopharmaceutical company ("Provectus" or "The Company"), today announced that it will present at the 2016 BIO International Convention in San Francisco, CA.

The Provectus presentation is currently scheduled to take place at 2:45 pm Pacific Daylight Time on Wednesday, June 8, 2016 in Theater 2 of the Moscone Center. The conference itself runs June 6-9, 2016.

The presentation will be available on the Company's website, [provectusbio.com](http://provectusbio.com), after the presentation ends.

**About the 2016 BIO International Convention**

The BIO International Convention (BIO) attracts over 15,000 biotechnology and pharma leaders who come together for one week of intensive networking to discover new opportunities and promising partnerships. This event covers a wide spectrum of life science and application areas including drug discovery, biomanufacturing, genomics, biofuels, nanotechnology and cell therapy. For more information, visit <http://convention.bio.org/2016/>

**About Provectus Biopharmaceuticals, Inc.**

Provectus Biopharmaceuticals, Inc. specializes in developing oncology and dermatology therapies. PV-10 is a novel investigational drug for cancer: 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 can be found at the NIH registry, [www.clinicaltrials.gov](http://www.clinicaltrials.gov). For additional information about Provectus, please visit the Company's website at [www.provectusbio.com](http://www.provectusbio.com) or contact Porter, LeVay & Rose, Inc.

-more-

*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, 2015) 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 investigational drug product for melanoma 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 PH-10, our investigational drug product for dermatology, 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.

###