



December 8, 2015

Spectrum Pharmaceuticals Highlights Four Abstracts on SPI-2012 and Poziotinib at the San Antonio Breast Cancer Symposium (SABCS)

HENDERSON, Nev.--(BUSINESS WIRE)-- Spectrum Pharmaceuticals (NasdaqGS: SPPI), a biotechnology company with fully integrated commercial and drug development operations with a primary focus in Hematology and Oncology, today announced key presentations of clinical and scientific data related to its products at the San Antonio Breast Cancer Symposium (SABCS), being held in San Antonio, Texas, from December 8-12, 2015.

For more information about the SABCS meeting and for a complete list of abstracts, please refer to the conference website at <http://www.abstracts2view.com/sabcs15/>.

The following are key SPI-2012 related abstracts being presented at the SABCS meeting:

Abstract #	Type	Title	First Author	Date/Time Location: Halls A-B
P1-10-05	Poster	Randomized phase 2, open-label, dose-ranging study of a novel, long-acting G-CSF (SPI-2012) or pegfilgrastim for the management of neutropenia in patients with breast cancer (BC) treated with (Neo) adjuvant chemotherapy with docetaxel + cyclophosphamide (TC)	Vacirca	Wednesday, Dec 9 5:00 PM
OT3-02-13	Poster	Randomized Phase 3 Study of a Novel, Long-acting G-CSF (SPI-2012) Versus Pegfilgrastim in the Management of Chemotherapy-Induced Neutropenia in Early-Stage Breast Cancer Patients Receiving Docetaxel and Cyclophosphamide (TC) (ADVANCE Study)	Schwartzberg	Friday, Dec 11 5:00 PM

The following are key Poziotinib related abstracts being presented at the SABCS meeting:

Abstract #	Type	Title	First Author	Date/Time Location: Halls A-B
P4-13-19	Poster	Poziotinib, an oral, irreversible pan-HER inhibitor, demonstrates promising clinical activity in metastatic HER2 positive breast cancer patients	Seock-Ah Im	Friday, Dec 11 7:30 AM
OT3-01-10	Poster	A prospective, open-label, single-arm, multi-center, phase II exploratory study to evaluate the efficacy and safety of poziotinib (NOV120101) in patients with HER2-positive metastatic breast cancer who have received at least two prior HER2-directed regimens	Park	Friday, Dec 11 5:00 PM

About Spectrum Pharmaceuticals, Inc.

Spectrum Pharmaceuticals is a leading biotechnology company focused on acquiring, developing, and commercializing drug products, with a primary focus in Oncology and Hematology. Spectrum and its affiliates market five oncology drugs— FUSILEV[®] (levoleucovorin) for Injection in the U.S.; FOLOTYN[®] (pralatrexate injection), also marketed in the U.S.; ZEVALIN[®] (ibrutinomab tiuxetan) Injection for intravenous use, for which the Company has worldwide marketing rights; MARQIBO[®] (vinCRISTine sulfate LIPOSOME injection) for intravenous infusion, for which the Company has worldwide marketing rights, and BELEODAQ[®] (belinostat) for Injection in the U.S. Additionally, Spectrum's pipeline includes three drugs targeting blockbuster markets in advanced stages of clinical development. Spectrum's strong track record in-licensing and acquiring differentiated drugs, expertise and proven track record in clinical development have generated a robust, diversified, and growing pipeline of product candidates in advanced-stage Phase 2 and Phase 3 studies. More information on Spectrum is available at www.sppirx.com.

Forward-looking statement - This press release may contain forward-looking statements regarding future events and the future performance of Spectrum Pharmaceuticals that involve risks and uncertainties that could cause actual results to differ materially. These statements are based on management's current beliefs and expectations. These statements include, but are not limited

to, statements that relate to our business and its future, including certain company milestones, Spectrum's ability to identify, acquire, develop and commercialize a broad and diverse pipeline of late-stage clinical and commercial products, leveraging the expertise of partners and employees around the world to assist us in the execution of our strategy, and any statements that relate to the intent, belief, plans or expectations of Spectrum or its management, or that are not a statement of historical fact. Risks that could cause actual results to differ include the possibility that our existing and new drug candidates may not prove safe or effective, the possibility that our existing and new applications to the FDA and other regulatory agencies may not receive approval in a timely manner or at all, the possibility that our existing and new drug candidates, if approved, may not be more effective, safer or more cost efficient than competing drugs, the possibility that our efforts to acquire or in-license and develop additional drug candidates may fail, our lack of sustained revenue history, our limited marketing experience, our dependence on third parties for clinical trials, manufacturing, distribution and quality control and other risks that are described in further detail in the Company's reports filed with the Securities and Exchange Commission. We do not plan to update any such forward-looking statements and expressly disclaim any duty to update the information contained in this press release except as required by law.

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