



## **Synta Announces Presentations at the 2013 Annual Meeting of the American Association of Cancer Research**

April 5, 2013

LEXINGTON, Mass.--(BUSINESS WIRE)--Apr. 5, 2013-- Synta Pharmaceuticals Corp. (NASDAQ: SNTA) announced today that poster presentations related to studies with ganetespib, a selective Hsp90 inhibitor in clinical development by the company, will be presented at the 2013 Annual Meeting of the American Association of Cancer Research in Washington, D.C. Presentations include:

### **Inhibition of mTOR enhances the activity of HSP90 inhibitors in part through cessation of heat shock protein synthesis**

Presentation: Sunday, April 7, 1:00 - 5:00 PM ET

Abstract number: 1038

Authors: He, et al.

### **Ultra-deep sequencing of circulating free DNA to identify predictive, mutated HSP90 clients in the GALAXY Trial (NCT01348126): a randomized phase IIB/III study of ganetespib (STA-9090) in combination with docetaxel versus docetaxel alone in subjects with stage IIIB/IV NSCLC**

Presentation: Monday, April 8, 1:00 - 5:00 PM ET

Abstract number: 2012

Authors: Fennell, et al.

### **The importance of dose schedule with HSP90 inhibitors: Results from a Phase II study in dogs with mast cell tumors**

Presentation: Tuesday, April 9, 8:00 AM - 12:00 PM ET

Abstract number: 3369

Authors: London, et al.

### **Differential sensitivities to heat shock protein 90 (HSP90) inhibitors in anaplastic lymphoma kinase (ALK)-positive non-small cell lung cancer (NSCLC) cells**

Presentation: Tuesday, April 9, 8:00 AM - 12:00 PM ET

Abstract number: 3272

Authors: Lee, et al.

### **Heat shock protein 90 functional inhibition regulates epithelial to mesenchymal transformation, invasion and migration via NF- $\kappa$ B and HIF-1 $\alpha$ signaling in colorectal cancer**

Presentation: Tuesday, April 9, 8:00 AM - 12:00 PM ET

Abstract number: 2707

Authors: Ganji, et al.

## **About Ganetespib**

Ganetespib is an inhibitor of heat shock protein 90 (Hsp90) that is structurally unrelated to first-generation, ansamycin-related Hsp90 inhibitors. In preclinical experiments, ganetespib has shown activity in multiple tumor models both as a single agent and in combination with certain widely used cancer agents. Company-sponsored clinical studies with ganetespib include 1) the randomized GALAXY-1 and GALAXY-2 trials evaluating ganetespib in combination with docetaxel as second-line treatment of non-small cell lung cancer (NSCLC), 2) the CHIARA Phase 2 trial evaluating ganetespib monotherapy in ALK+ NSCLC, and 3) the ENCHANT Phase 2 trial evaluating ganetespib as first-line treatment for HER2+ and triple-negative metastatic breast cancer. In addition, ganetespib is being evaluated in investigator-sponsored trials for treatment of a number of solid tumor and hematologic cancer indications. Information on these trials can be found at [www.clinicaltrials.gov](http://www.clinicaltrials.gov).

The safety profile of ganetespib has been favorable in over 700 patients treated to date in more than 20 clinical trials. Transient, mild or moderate diarrhea has been the most commonly reported adverse event.

## **About Hsp90**

Hsp90 is a molecular chaperone required for the proper folding and activation of many cancer-promoting proteins, and is recognized as a key facilitator of cancer cell growth and survival. Many of the “client proteins” of Hsp90 – such as AKT, ALK, BCR-ABL, BRAF, EGFR, FLT3, HER2, HIF-1alpha, KIT, MET, PDGFRA, and VEGFR – are the targets of clinically validated cancer drugs. In preclinical studies, inhibiting Hsp90 causes the degradation of multiple client proteins and leads to cancer cell death.

## **About the GALAXY Trials**

The GALAXY (Ganetespib Assessment in Lung cAnCER with docetaXel) program consists of two randomized trials comparing the combination of ganetespib and docetaxel versus docetaxel alone in patients with Stage IIIB/IV NSCLC who have received one prior systemic therapy: a 300-patient Phase 2b/3 trial (GALAXY-1) to determine the patient population most likely to derive benefit from ganetespib, and a 500-patient confirmatory Phase 3 trial (GALAXY-2). More information about the GALAXY trials can be found at [www.clinicaltrials.gov](http://www.clinicaltrials.gov) (NCT01348126 and NCT01798485).

## **About Synta Pharmaceuticals**

Synta Pharmaceuticals Corp. is a biopharmaceutical company focused on discovering, developing, and commercializing small molecule drugs to extend and enhance the lives of patients with severe medical conditions, including cancer and chronic inflammatory diseases. Synta has a unique chemical compound library, an integrated discovery engine, and a diverse pipeline of clinical- and preclinical-stage drug candidates with distinct mechanisms of action and novel chemical structures. All Synta drug candidates were invented by Synta scientists using our compound library and discovery capabilities. For more information, please visit [www.syntapharma.com](http://www.syntapharma.com).

## **Safe Harbor Statement**

This media release may contain forward-looking statements about Synta Pharmaceuticals Corp. Such forward-looking statements can be identified by the use of forward-looking terminology such as "will", "would", "should", "expects", "anticipates", "intends", "plans", "believes", "may", "estimates", "predicts", "projects", or similar expressions intended to identify forward-looking statements. Such statements reflect our current views with respect to future events and are based on assumptions and subject to risks and uncertainties that could cause actual results to differ materially from those expressed or implied by such forward-looking statements, including those described in "Risk Factors" of our Form 10-K for the year ended December 31, 2012 as filed with the Securities and Exchange Commission. Synta undertakes no obligation to publicly update forward-looking statements, whether because of new information, future events or otherwise, except as required by law.

Source: Synta Pharmaceuticals Corp.

### **Investor Relations Contact:**

Synta Pharmaceuticals Corp.  
George Farmer, 781-541-7213  
[gfarmer@syntapharma.com](mailto:gfarmer@syntapharma.com)