Exelixis Provides Update on Timing of Key Cabozantinib Clinical Data Presentation at the ESMO 2016 Congress

September 20, 2016

- CABOSUN results now subject of an oral presentation in Presidential Symposium 3 on Monday, October 10 -

SOUTH SAN FRANCISCO--(BUSINESS WIRE)--Sep. 20, 2016-- Exelixis, Inc. (NASDAQ:EXEL) today provided an update on the timing of a key data presentation for cabozantinib at the European Society for Medical Oncology (ESMO) 2016 Congress, which is being held October 7-11, 2016 in Copenhagen, Denmark. Detailed results from CABOSUN, the randomized phase 2 clinical trial of cabozantinib compared with sunitinib in patients with previously untreated advanced renal cell carcinoma (RCC), has been selected for the Presidential Symposium 3 session on Monday, October 10, 2016, starting at 16:30 CEST (local Copenhagen time) / 10:30 a.m. EDT / 7:30 a.m. PDT.

The full logistical details for the CABOSUN data presentation are as follows:

Oral Presentation

[LBA30_PR] "CABOzantinib versus SUNitinib (CABOSUN) as initial targeted therapy for patients with metastatic renal cell carcinoma (mRCC) of poor and intermediate risk groups: Results from ALLIANCE A031203 Trial."

Dr. Toni Choueiri, Director, Lank Center for Genitourinary Oncology, Dana-Farber Cancer Institute, Boston, Massachusetts, USA

Session: Presidential Symposium 3, Monday, October 10, 2016 – 16:30-18:10 CEST, Copenhagen room

Note: This is a National Cancer Institute Cancer Therapy Evaluation Program (NCI-CTEP) study.

The late-breaking CABOSUN abstract was initially slated for an oral presentation at a Proffered Paper session on Saturday, October 8, 2016. Exelixis previously announced that data from the Exelixis-discovered compounds cabozantinib and cobimetinib would be the subject of fifteen presentations at the ESMO 2016 Congress. For full details on Exelixis' presence at the conference, please see the company's press release issued on August 31, 2016.

About Exelixis

Exelixis, Inc. (Nasdaq: EXEL) is a biopharmaceutical company committed to the discovery, development and commercialization of new medicines with the potential to improve care and outcomes for people with cancer. Since its founding in 1994, three medicines discovered at Exelixis have progressed through clinical development to receive regulatory approval. Currently, Exelixis is focused on advancing cabozantinib, an inhibitor of multiple tyrosine kinases including MET, AXL and VEGF receptors, which has shown clinical anti-tumor activity in more than 20 forms of cancer and is the subject of a broad clinical development program. Two separate formulations of cabozantinib have received regulatory approval to treat certain forms of kidney and thyroid cancer and are marketed for those purposes as CABOMETYX™ tablets (U.S. and EU) and COMETRIQ® capsules (U.S. and EU), respectively. Another Exelixis-discovered compound, COTELLIC® (cobimetinib), a selective inhibitor of MEK, has been approved in major territories including the United States and European Union, and is being evaluated for further potential indications by Roche and Genentech (a member of the Roche Group) under a collaboration with Exelixis. For more information on Exelixis, please visit www.exelixis.com or follow @ExelixisInc on Twitter.

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