



Exelixis to Release Second Quarter 2016 Financial Results on Wednesday, August 3, 2016

July 12, 2016

- Conference Call and Webcast to Follow at 5:00 p.m. EDT/ 2:00 p.m. PDT -

SOUTH SAN FRANCISCO, Calif.--(BUSINESS WIRE)--Jul. 12, 2016-- Exelixis, Inc. (NASDAQ: EXEL) announced today that its second quarter 2016 financial results will be released on Wednesday, August 3, 2016 after the markets close. At 5:00 p.m. EDT / 2:00 p.m. PDT, Exelixis management will host a conference call to discuss the results and provide a general business update. The conference call will be accessible via the Internet from the company's website.

To access the webcast link, log onto www.exelixis.com and proceed to the Event Calendar page under Investors & Media. Please connect to the company's website at least 15 minutes prior to the conference call to ensure adequate time for any software download that may be required to listen to the webcast. Alternatively, please call (855) 793-2457 (domestic) or (631) 485-4921 (international) and provide the conference call passcode 43700006 to join by phone.

A telephone replay will be available until 11:59 p.m. EDT on August 5, 2016. Access numbers for the telephone replay are: 855-859-2056 (domestic) and 404-537-3406 (international); the passcode is 43700006. A webcast replay will also be archived on www.exelixis.com for one year.

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 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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