May 6, 2013

pSivida Corp. Announces Third Quarter 2013 Financial Results Release Date and Conference Call Information

WATERTOWN, Mass.--(BUSINESS WIRE)-- pSivida Corp. (NASDAQ:PSDV) (ASX:PVA), a leader in developing sustained release, drug delivery products for treatment of back-of-the-eye diseases, today announced that its financial results for the third quarter of fiscal year 2013 will be released after the market close on Monday, May 13, 2013, followed the same day by a conference call and live webcast scheduled for 4:30 p.m. ET.

The conference call may be accessed by dialing (877) 303-9236 from the U.S. and Canada, or (760) 666-3569 from international locations. A live webcast will be available on the Investor Relations section of the corporate website at http://www.psivida.com.

A replay of the call will be available beginning May 13, 2013 at approximately 7:30 p.m. ET and ending on May 20, 2013. The replay may be accessed by dialing (855) 859-2056 within the U.S. and Canada or (404) 537-3406 from international locations, Conference ID Number: 68825398. A replay of the webcast will also be available on the corporate website during that time.

About pSivida Corp.

pSivida Corp., headquartered in Watertown, MA, develops tiny, sustained release, drug delivery products designed to deliver drugs at a controlled and steady rate for months or years. pSivida is currently focused on treatment of chronic diseases of the back of the eye utilizing its core technology systems, Durasert™ and BioSilicon™. The injectable, sustained release micro insert ILUVIEN® for the treatment of chronic DME, licensed to Alimera Sciences, Inc., has received marketing authorization in Austria, France, Germany, Portugal, Spain and the U.K. and is awaiting authorization in Italy. ILUVIEN for DME has not been approved in the U.S. pSivida plans to institute pivotal Phase III clinical trials for the treatment of posterior uveitis, a chronic back-of-the-eye disease, with the same micro-insert as ILUVIEN for DME. An investigator-sponsored clinical trial is ongoing for an injectable, bioerodible micro-insert to treat glaucoma and ocular hypertension. pSivida's FDA-approved Retisert® licensed to Bausch & Lomb Incorporated provides long-term, sustained drug delivery to posterior uveitis.

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The President's Blog: http://www.thechairmansblog.com/paul-ashton

For more information on pSivida, visit www.psivida.com.

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