



Ocular Therapeutix™ to Present Data at the Association for Research in Vision and Ophthalmology (ARVO) Annual Meeting

June 15, 2020

BEDFORD, Mass.--(BUSINESS WIRE)--Jun. 15, 2020-- Ocular Therapeutix, Inc. (NASDAQ:OCUL), a biopharmaceutical company focused on the formulation, development, and commercialization of innovative therapies for diseases and conditions of the eye, announced multiple scientific presentations at the Association for Research in Vision and Ophthalmology (ARVO) Annual Meeting. With the ARVO 2020 Annual Meeting being moved to a virtual format, video-recorded presentations became available online on [ARVOLearn](https://www.arvo.org/ARVO2020/ARVOLearn) as of June 12, 2020. Additionally, abstracts will be published in the journal, *Investigative Ophthalmology & Vision Science* (IOVS), no later than July 31, 2020.

"We are pleased to be presenting data on our pre-clinical and clinical programs at ARVO," commented Michael Goldstein, MD, MBA, Chief Medical Officer of Ocular Therapeutix. "At Ocular Therapeutix, we are excited with the progress we are making with advancing all of the programs in our pipeline. Our hydrogel platform continues to offer a patients and physicians a novel, hands free option to address both front- and back-of-the-eye diseases. We look forward to the continued progress of these programs while we continue expanding the capabilities of our diverse platform."

Presentations at ARVO:

- Evaluating Safety and Pharmacokinetics of OTX-CSI, a Sustained Release Intracanalicular Cyclosporine Insert in Beagles. *Vanslette A et al*
- Effect of Hydrogel Persistence on Pharmacodynamics and Tolerability of OTX-TIC, Travoprost Intracameral Implant in Beagles. *Langh J et al*
- Evaluating Tolerability and Pharmacokinetics of OTX-BDI, a Sustained-Release Intracanalicular Insert Co-Delivering Besifloxacin & Dexamethasone in Beagles. *Rosales C et al*
- Evaluation of the Safety and Efficacy of OTX-TP, an intracanalicular travoprost insert, for the treatment of patients with open-angle glaucoma or ocular hypertension: A Phase 3 Study. *Vantipalli S et al*

The online presentations will be hosted in the Media library catalog within ARVOLearn.

About Ocular Therapeutix, Inc.

Ocular Therapeutix, Inc. is a biopharmaceutical company focused on the formulation, development, and commercialization of innovative therapies for diseases and conditions of the eye using its proprietary bioresorbable hydrogel-based formulation technology. Ocular Therapeutix's first commercial drug product, DEXTENZA® (dexamethasone ophthalmic insert), is FDA-approved for the treatment of ocular inflammation and pain following ophthalmic surgery. Ocular Therapeutix recently completed a Phase 3 clinical trial evaluating DEXTENZA for the treatment of ocular itching associated with allergic conjunctivitis. OTX-TP (intracanalicular travoprost insert) is an intracanalicular insert in clinical development for the reduction of intraocular pressure in patients with primary open-angle glaucoma and ocular hypertension. The Company's earlier stage development assets currently in Phase 1 trials include OTX-TIC (travoprost intracameral implant) for the reduction of intraocular pressure in patients with primary open-angle glaucoma and ocular hypertension, OTX-CSI (cyclosporine intracanalicular insert) for the treatment of the signs and symptoms of dry eye disease and OTX-TKI (axitinib intravitreal implant) for the treatment of retinal diseases. Also, in collaboration with Regeneron, Ocular Therapeutix is currently developing OTX-IVT (aflibercept suprachoroidal injection), an extended-delivery formulation of aflibercept for the treatment of retinal diseases. Ocular Therapeutix's first product, ReSure® Sealant, is FDA-approved to seal corneal incisions following cataract surgery.

Forward Looking Statements

Any statements in this press release about future expectations, plans, and prospects for the Company, including the commercialization of DEXTENZA®, ReSure Sealant, or any of the Company's product candidates; the commercial launch of, and effectiveness of reimbursement codes for, DEXTENZA; the development and regulatory status of the Company's product candidates, such as the Company's development of and prospects for approvability of DEXTENZA for additional indications including allergic conjunctivitis, OTX-TP for the treatment of primary open-angle glaucoma and ocular hypertension, OTX-CSI for the treatment of dry eye disease, OTX-TIC for the treatment of primary open-angle glaucoma and ocular hypertension, OTX-TKI for the treatment of retinal diseases including wet AMD, and OTX-IVT as an extended-delivery formulation of the VEGF trap aflibercept for the treatment of retinal diseases including wet AMD; the presentation of data on the Company's products and product candidates at medical conferences; the ongoing development of the Company's extended-delivery hydrogel depot technology; the potential utility of any of the Company's product candidates; the potential benefits and future operation of the collaboration with Regeneron Pharmaceuticals, including any potential future payments thereunder; the expected impact of the COVID-19 pandemic on the Company and its operations; the sufficiency of the Company's cash resources and other statements containing the words "anticipate," "believe," "estimate," "expect," "intend," "goal," "may," "might," "plan," "predict," "project," "target," "potential," "will," "would," "could," "should," "continue," and similar expressions, constitute forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995. Actual results may differ materially from those indicated by such forward-looking statements as a result of various important factors. Such forward-looking statements involve substantial risks and uncertainties that could cause the Company's clinical development programs, future results, performance or achievements to differ significantly from those expressed or implied by the forward-looking statements. Such risks and uncertainties include, among others, the timing and costs involved in commercializing DEXTENZA, ReSure Sealant or any product candidate that receives regulatory approval, including the conduct of post-approval studies, the ability to retain regulatory approval of DEXTENZA, ReSure Sealant or any product candidate that receives regulatory approval, the ability to maintain reimbursement codes for DEXTENZA, the initiation, timing and conduct of clinical trials, availability of data from clinical trials and expectations for regulatory submissions and approvals, the Company's scientific approach and general development progress, the availability or

commercial potential of the Company's product candidates, the Company's ability to generate its projected net product revenue on the timeline expected, if at all, the sufficiency of cash resources, the Company's existing indebtedness, the ability of the Company's creditors to accelerate the maturity of such indebtedness upon the occurrence of certain events of default, the outcome of the Company's ongoing legal proceedings, the severity and duration of the COVID-19 pandemic including its effect on the Company's and relevant regulatory authorities' operations, the need for additional financing or other actions and other factors discussed in the "Risk Factors" section contained in the Company's quarterly and annual reports on file with the Securities and Exchange Commission. In addition, the forward-looking statements included in this press release represent the Company's views as of the date of this release. The Company anticipates that subsequent events and developments will cause the Company's views to change. However, while the Company may elect to update these forward-looking statements at some point in the future, the Company specifically disclaims any obligation to do so except as required by law. These forward-looking statements should not be relied upon as representing the Company's views as of any date subsequent to the date of this release.

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