

Ocuphire Pharma to Present at The H.C. Wainwright 24th Annual Global Investment Conference

FARMINGTON HILLS, Mich., Sept. 06, 2022 (GLOBE NEWSWIRE) -- Ocuphire Pharma, Inc. (Nasdaq: OCUP), a clinical-stage ophthalmic biopharmaceutical company focused on developing and commercializing therapies for the treatment of refractive and retinal eye disorders, today announced that Mina Sooch, MBA, Founder and CEO will present a corporate overview at the H.C. Wainwright 24th Annual Global Investment Conference being held September 12-14, 2022.

H.C. Wainwright 24th Annual Global Investment Conference – September 12-14, 2022

Title: Ocuphire Pharma (OCUP) Company Presentation

Date: Monday, September 12, 2022, available after 7 AM EST

Presenter: Mina Sooch, MBA, CEO and Founder, Ocuphire

Format: On-Demand Session

Conference

Regis.:

If you are interested in arranging a 1x1 meeting request, please contact your bank/conference representative or <u>ir@ocuphire.com</u>. For more details, please see the <u>Investors</u> and <u>Events</u> section of Ocuphire's corporate website.

About Ocuphire Pharma

Ocuphire is a publicly-traded (NASDAQ: OCUP), clinical-stage ophthalmic biopharmaceutical company focused on developing and commercializing small-molecule therapies for the treatment of refractive and retinal eye disorders. The Company's lead product candidate, Nyxol® eye drops (0.75% phentolamine ophthalmic solution), is a oncedaily, preservative-free eye drop formulation of phentolamine mesylate, a non-selective alpha-1 and alpha-2 adrenergic antagonist designed to reduce pupil size, and is being developed for several indications, including reversal of pharmacologically-induced mydriasis (RM), presbyopia and dim light or night vision disturbances (NVD), and has been studied in 12 completed clinical trials. Ocuphire has reported positive data from MIRA-2 and MIRA-3 registration trials and MIRA-4 pediatric safety trial for the treatment of RM. Ocuphire also reported positive top-line data from the VEGA-1 Phase 2 trial of Nyxol for treatment of presbyopia, both Nyxol as a single agent and Nyxol with low dose pilocarpine ("LDP") 0.4%

as adjunctive therapy. The Company recently reported positive top-line results from LYNX-1 Phase 3 trial of Nyxol for NVD. Ocuphire's second product candidate, APX3330, is an oral tablet designed to inhibit angiogenesis and inflammation pathways relevant to retinal and choroidal vascular diseases, such as diabetic retinopathy (DR) and diabetic macular edema (DME) and has been studied in 11 Phase 1 and 2 trials. The Company announced in March the completion of enrollment in the ZETA-1 Phase 2b clinical trial of APX3330 to treat DR/DME. Please visit www.clinicaltrials.gov to learn more about Ocuphire's ongoing APX3330 Phase 2b trial in DR/DME ZETA-1 (NCT04692688) and completed Nyxol trials: Phase 3 registration trial in NVD LYNX-1 (NCT04692688), Phase 3 registration trials in RM MIRA-2 (NCT04620213) and MIRA-3 (NCT04638660), Phase 3 registration trials in RM MIRA-2 (NCT04620213) and MIRA-3 (NCT04638660), Phase 3 registration trials in RM MIRA-2 (NCT04620213) and MIRA-3 (NCT05134974), MIRA-4 Phase 3 pediatric safety study (NCT04620213) and MIRA-3 (NCT05134974), MIRA-4 Phase 3 pediatric safety study (NCT04675151). As part of its strategy, Ocuphire will continue to explore opportunities to acquire additional ophthalmic assets and seek strategic partners for late-stage development, regulatory preparation, and commercialization of drugs in key global markets. For more information, visit www.ccuphire.com.

Contacts

Corporate

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Source: Ocuphire Pharma