

UNITED STATES PATENT AND TRADEMARK OFFICE

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BEFORE THE PATENT TRIAL AND APPEAL BOARD

SLAYBACK PHARMA LLC

Petitioner

v.

EYE THERAPIES, LLC

Patent Owner

Patent No. 8,293,742

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Case No. IPR2021-Unassigned

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**DECLARATION OF PAUL A. LASKAR, PH.D.**

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1. I, Paul A. Laskar, Ph.D., have been retained by counsel for Petitioner Slayback Pharma LLC (Petitioner or Slayback). I understand that Petitioner seeks *inter partes* review (“IPR”) of U.S. Patent 8,293,742 (EX-1001, ‘742 patent), assigned to Eye Therapies, LLC (Patent Owner), to request that the United States Patent and Trademark Office cancel claims 1-6 of the ‘742 patent as unpatentable. I submit this expert declaration in support of Petitioner’s IPR Petition for the ‘742 patent.

## **I. QUALIFICATIONS AND BACKGROUND**

### **A. Education and Experience**

2. I am currently President and Principal Consultant with Paul Laskar Associates, Inc.

3. In 1965 I received a B.A. in General Science (Chemistry, Biology) from University of Rochester.

4. In 1968 I received a B.S. in Pharmacy from University of Illinois – Medical Center.

5. In 1971 I received an M.S. in Pharmacy from University of Illinois – Medical Center.

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6. In 1974 I completed my Ph.D. in Pharmaceutical Sciences at Oregon State University with a minor in Biostatistics.

7. In 1988 I received an MBA from University of California at Irvine, (General Management, International Management, Marketing).

8. Following completion of my graduate work, I taught for about nine years at two colleges of pharmacy, University of Illinois-Chicago Campuses and Creighton University. In late 1982, I joined Allergan as a formulation scientist where I advanced ultimately to Director of Product Development in Allergan's dermatology business group, Herbert Laboratories. In this capacity, I was involved in CMC (chemistry, manufacturing, and controls) activities for Allergan's beta-blocker ophthalmic as well as other ophthalmic products as well as their first topical retinoid.

9. I then joined CoCensys where I led the CMC efforts for two successful INDs, one for a parenteral and the other for an oral liquid. In 1994, I joined Santen Inc, the U.S. subsidiary of a leading Japanese specialty company, Santen LTD. There I led the pharmaceutical development department which during my tenure resulted in four successful NDAs and formulation of Santen's first prostanoid product for glaucoma.

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