

**REPLY EXPERT REPORT OF
JOHN C. JAROSZ
ON OBJECTIVE INDICIA OF NON-OBVIOUSNESS**

TABLE OF CONTENTS

I.	INTRODUCTION	1
	A. Assignment	1
	B. Qualifications	1
	C. Evidence Considered	2
	D. Summary of Opinions	2
II.	FRAMEWORK.....	3
III.	MARKETPLACE SUCCESS OF PROLENSA®.....	4
	A. IMS Data	5
	B. Prolensa® Profitability	6
	C. Prolensa® Relative Performance	12
	D. Third-Party Perceptions	17
	E. Licensing	17
IV.	CAUSAL NEXUS	18
	A. “Life-Cycle Management”	19
	1. Benefits of What Mr. Hofmann Calls a “Life-Cycle Management” Strategy.....	19
	2. Bausch & Lomb’s Strategy	25
	B. Prolensa® Marketing and Promotion	29
	1. Prolensa® Marketing Expenditures	29
	2. Relative Marketing of Prolensa® and Ilevro®	36
	3. Prolensa® Marketing Message	38
	C. Pricing	40
	1. Discounts and Coupons.....	40
	2. Pricing Relative to Generic Ophthalmic NSAIDs	40
	3. Pricing Relative to Branded Ophthalmic NSAIDs	42
	D. Generic ANDA Filers	43
V.	CONCLUSION.....	46

I. INTRODUCTION

A. Assignment

1. I, John C. Jarosz, submit this reply expert report on behalf of Bausch & Lomb Incorporated, Bausch & Lomb Pharma Holdings Corp. (collectively, “Bausch & Lomb”) and Senju Pharmaceutical Co. Ltd. (“Senju”) (collectively, with Bausch & Lomb, “Patent Owners”) in connection with the above captioned cases. I have been retained to provide expert analysis and testimony, if necessary, regarding the commercial success of the inventions described in U.S. Patent Nos. 8,129,431 (“the ’431 patent”); 8,669,290 (“the ’290 patent”); 8,754,131 (“the ’131 patent”); 8,871,813 (“the ’813 patent”); and 8,927,606 (“the ’606 patent”) (collectively, the “Patents-in-Suit”). It is my understanding that the asserted claims of the Patents-in-Suit are embodied in Bausch & Lomb’s Prolensa® product.
2. On December 30, 2015, I submitted my opening expert report on objective indicia of non-obviousness in these cases.¹ Since then, I have received the responsive report of Ivan T. Hofmann.² I have been asked to provide my opinions regarding the analysis and conclusions set forth in the Hofmann Report. This report summarizes those opinions.
3. As with my initial report, I may modify or supplement my opinions, if necessary and allowed, based on the review and analysis of information provided to me subsequent to the filing of this report.

B. Qualifications

4. A complete description of my background and qualifications is provided in the Jarosz

¹ Opening Expert Report of John C. Jarosz on Objective Indicia of Non-Obviousness, December 30, 2015 (“Jarosz Report”). I submitted two versions of the Jarosz Report, one applicable to the Lupin Defendants and one applicable to the InnoPharma defendants. Page and Tab references in this report refer to the Lupin version of the Jarosz Report.

² Responsive Expert Report of Ivan T. Hofmann, CPA/CFF, CLP, February 1, 2016 (“Hofmann Report”).

Report. An updated copy of my curriculum vitae is provided as Reply Tab 1.

C. Evidence Considered

5. Since submitting the Jarosz Report, I have reviewed additional information from a variety of sources, including the Hofmann Report, the responsive report of Dr. Robert C. Cykiert,³ the reply report of Dr. William B. Trattler,⁴ materials produced by Patent Owners in this litigation, and information from publicly-available sources, such as academic journals and analyst reports. A complete list of additional materials that I have received and reviewed since the date of the Jarosz Report is attached as Reply Tab 2.

D. Summary of Opinions

6. In his report, Mr. Hofmann concluded that

Prolensa® is not a commercial success and the performance of Prolensa® is attributable to various extrinsic factors unrelated to the Patents-in-Suit. Specifically, the performance of Prolensa® is explained by the execution of a coordinated life-cycle management strategy for the bromfenac franchise which involved the following components: (1) the systematic migration to new bromfenac products and the discontinuation of legacy bromfenac products; (2) substantial marketing and promotional efforts; and (3) tactical pricing of Prolensa®. As a result, the performance of Prolensa® does not provide objective indicia of nonobviousness of the Patents-in-Suit.⁵

7. I disagree with Mr. Hofmann's conclusions for a number of reasons.
- Mr. Hofmann's conclusion that Prolensa® is not a commercial success is inconsistent with the evidence of Prolensa®'s marketplace performance over time. Prolensa® has achieved and maintained substantial acceptance, particularly in light of the array of competitive alternatives.

³ Responsive Expert Report of Robert C. Cykiert, M.D. on Objective Indicia of Non-Obviousness, February 1, 2016 ("Cykiert Report").

⁴ Reply Expert Report of William B. Trattler, MD, on Objective Indicia of Non-Obviousness, February 12, 2016 ("Trattler Reply Report").

⁵ Hofmann Report, at 15.

- Mr. Hofmann’s conclusion that the success of Prolensa® is due to factors *other than* the Patents-in-Suit is inconsistent with evidence showing that the patents have been motivating (important) factors in Prolensa®’s success.
- Mr. Hofmann’s characterization of any Prolensa® success as reflecting a “life-cycle management” strategy fails to acknowledge that the success of such a strategy requires that a new formulation must actually be deemed advantageous by the physician community before it will be prescribed. Here, physicians had a compelling reason to switch to Prolensa®, which is the improved side effect profile offered by Prolensa® relative to other available bromfenac formulations.
- Mr. Hofmann’s analysis of Prolensa® marketing expenditures fails to recognize that marketing is one of many factors that influence physician prescribing behavior, and its impact is modest. Physicians are informed by marketing efforts, but weigh heavily the quality and effectiveness of a drug, as well as patient requests, when deciding what to prescribe. Marketing spending alone is not sufficient if the drug does not offer clinical benefits to patients, as Prolensa® does.
- Mr. Hofmann’s conclusion that Prolensa® may be cheaper than generic bromfenac is inconsistent with the evidence and business realities. His analysis of Prolensa® net pricing relative to competing ophthalmic NSAIDs is incomplete.

II. FRAMEWORK

8. According to Mr. Hofmann, “any alleged commercial success must be *driven primarily* by and attributable to the purported merits of the claimed invention, and *not by other factors* unrelated to the allegedly novel features of the claimed invention. ... there must be a causal correlation, or ‘nexus,’ between the unique merit of the claimed invention and

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