

In the United States Patent and Trademark Office

Before the Patent Trial and Appeal Board

AMERIGEN PHARMACEUTICALS LIMITED,

Petitioner

v.

UCB PHARMA GMBH

Patent Owner

U.S. Patent No. 6,858,650

Filing Date: November 15, 2000

Issue Date: February 22, 2005

Title: STABLE SALTS OF NOVEL DERIVATIVES
OF 3,3-DIPHENYLPROPYLAMINES

Inter Partes Review No.: Unassigned

**PETITION FOR *INTER PARTES* REVIEW
UNDER 35 U.S.C. §§ 311-319 AND 37 C.F.R. §§ 42.100 ET SEQ.**

TABLE OF CONTENTS

I. INTRODUCTION 8

II. MANDATORY NOTICES (37 C.F.R. § 42.8(a)(1)) 8

 A. Real Party-In-Interest Under 37 C.F.R. § 42.8(b)(1)8

 B. Related Matters Under 37 C.F.R. § 42.8(b)(2)9

 C. Lead And Back-Up Counsel Under 37 C.F.R. § 42.8(b)(3).....9

 D. Service Information Under 37 C.F.R. § 42.8(b)(4).....10

III. GROUNDS FOR STANDING (37 C.F.R. §§ 42.101 and 42.104).....10

IV. PAYMENT OF FEES (37 C.F.R. § 42.103)11

V. STATEMENT OF THE PRECISE RELIEF REQUESTED AND THE REASONS THEREFOR (37 C.F.R. § 42.22(a))11

VI. SUMMARY OF THE ‘650 PATENT AND CHALLENGED CLAIMS13

VII. CLAIM CONSTRUCTION14

VIII. TECHNICAL BACKGROUND AND STATE OF THE ART14

 A. The Person of Ordinary Skill in the Art of the ‘650 Patent14

 B. Before the Invention, Antimuscarinic Compounds Were Used to Treat Overactive Bladder Conditions14

 C. Prodrugs Were Known to Solve Active Compound Difficulties17

 D. Numerous Salt Forms Were Known for Compounds Similar to the Most Effective Overactive Bladder Drugs21

IX. SCOPE AND CONTENT OF THE PRIOR ART22

 A. Skilled Artisans Had Ample Motivation to Focus on Optimizing 5-HMT to Obtain an Overactive Bladder Compound22

 1. *Postlind, the Detrol® Label, and Brynne 1998 Taught 5-HMT Was an Effective Compound for Overactive Bladder without Tolterodine.*22

 2. *Skilled Artisans Would Immediately Recognize the Benefit to Starting with their Knowledge of 5-HMT and Tolterodine and Not Other Compounds.*24

 B. Bundgaard Taught Predictable Modifications to Improve 5-HMT Delivery26

 C. Berge and Johansson Taught Fumarate Salts29

X.	DETAILED GROUNDS FOR UNPATENTABILITY.....	30
A.	Claims 1-5 are Obvious Over the Postlind and Bundgaard Publications in view of the Detrol® Label and Berge.....	30
1.	<i>A Person of Ordinary Skill Would Have Been Motivated to Look at Improved 5-HMT Administration in View of Tolterodine.....</i>	31
2.	<i>Postlind and Bundgaard Publications in View of the Detrol® Label and Berge Would Have Led to Prodrug Optimization and Fumarate Salt Forms.</i>	34
3.	<i>Summary of Proposed Rejection of Claims 1-5.....</i>	40
B.	Claims 21-24 are Obvious over the Postlind and Bundgaard Publications in view of the Detrol® Label and Berge.....	48
C.	Claims 1-5 and 21-24 Are Rendered Obvious by Brynne 1998, Bundgaard, and Johansson.....	53
1.	<i>A Person of Ordinary Skill Would Have Been Motivated to Look at Improved 5-HMT Administration in View of Tolterodine.....</i>	53
2.	<i>Brynne 1998 in View of Bundgaard and Johansson Would Have Led to Prodrug Optimization and Fumarate Salt Forms.....</i>	54
XI.	EVEN IF CONSIDERED, SECONDARY CONSIDERATIONS FAIL TO OVERCOME THE EVIDENCE OF OBVIOUSNESS.....	63
XII.	THE PROPOSED REJECTIONS RAISE NEW ISSUES IN WHICH PETITIONER WILL LIKELY PREVAIL.....	68

LIST OF EXHIBITS

- Ex. 1001: U.S.P.N. 6,858,650
- Ex. 1002: File History for U.S.P.N. 6,858,650
- Ex. 1003: Declaration of Dr. Steven Patterson, Ph.D.
- Ex. 1004: C.V. for Dr. Steven Patterson, Ph.D.
- Ex. 1005: “Johansson” – WO 94/11337 Filed 6 November 1992 – “Novel 3,3-Diphenylpropylamines, Their Use and Preparation”
- Ex. 1006: “Andersson Review” – BJU International (1999), 84, 923-947 – “The Pharmacological Treatment of Urinary Incontinence”; K-E Andersson, R. Appell, L.D. Cardozo, C. Chapple, H.P. Drutz, A.E. Finkbeiner, F. Haab, and R. Vela Navarrete
- Ex. 1007: “Brynne 1997” – International Journal of Clinical Pharmacology and Therapeutics (1997), 35, 287-295 – “Pharmacokinetics and pharmacodynamics of tolterodine in man: a new drug for the treatment of urinary bladder overactivity”; N. Brynne, M.M.S. Stahl, B. Hallen, P.O. Edlund, L. Palmer, P. Hoglund, and J. Gabrielsson
- Ex. 1008: “Thomas” – British Heart Journal (1995), 74, 53-56 – “Concentration dependent cardiotoxicity of terodine in patients treated for urinary incontinence”; S. Thomas, P. Higham, K Hartigan-Go, F. Kamali, P. Wood, R. Campbell, and G. Ford
- Ex. 1009: “Detrol® Label” – Pharmacia & Upjohn
- Ex. 1010: “Postlind” – Drug Metabolism and Disposition (1998), 26 (4), 289293 – “Tolterodine, A New Muscarinic Receptor Antagonist, Is Metabolized by Cytochromes P450 2D6 and 3A in Human Liver Microsomes”; H. Postlind, A. Danielson, A. Lindgren, and S. Andersson
- Ex. 1011: “Brynne 1998” – Clinical Pharmacology & Therapeutics (May 1998), 63(5), 529-539 – “Influence of CYP2D6 polymorphism

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- Ex. 1012: “Bundgaard” – Elsevier 1985 – “Design of Prodrugs”
- Ex. 1013: “Berge 1977” – Journal of Pharmaceutical Sciences (1977), 66 (1), 1-19 – “Pharmaceutical Salts”; S. Berge, L., Bighley, and D. Monkhouse
- Ex. 1014: “Andersson 1998” – Drug Metabolism and Disposition (1998), 26(6), 528-535 – “Biotransformation of tolterodine, a new muscarinic receptor antagonist, in mice, rats, and dogs”; S. Andersson, A. Lindgren, and H. Postlind
- Ex. 1015: “Nilvebrant” – Pharmacology and Toxicology (1997), 81, 169-172 – “Antimuscarinic Potency and Bladder Selectivity of PNU-200577, a Major Metabolite of Tolterodine”; L. Nilvebrant, P. Gillberg, and B. Sparf
- Ex. 1016: “DeMaagd” – P&T (2012), 37(6), 345-361 – “Management of Urinary Incontinence”; G. DeMaagd and T. Davenport
- Ex. 1017: “Appell” – Urology (1997), 50, 90-96 – “Clinical efficacy and safety of tolterodine in the treatment of overactive bladder: a pooled analysis”; R. Appell
- Ex. 1018: “Ashworth” – Home Care Provider (1997), 2(3), 117-120 – “Is My Antihistamine Safe?”; L. Ashworth
- Ex. 1019: “Lipinski” – Advanced Drug Delivery Reviews, 1997
- Ex. 1020: “Bundgaard PCT” – WO 92/08459 Filed 11 November 1991 – “Topical Compositions for Transdermal Delivery of Prodrug Derivatives of Morphine”
- Ex. 1021: “AUA Guideline” – American Urological Association Education and Research (2014) – “Diagnosis and Treatment of Overactive Bladder (Non-Neurogenic) in Adults: AUA/SUFU Guideline”; E. Gormley, et al

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