

Receipt Number

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IN THE UNITED STATES DISTRICT COURT  
FOR THE EASTERN DISTRICT OF MICHIGAN  
SOUTHERN DIVISION

Exhibit A

ELI LILLY AND COMPANY,

Plaintiff,

vs.

SUN PHARMACEUTICAL INDUSTRIES, LTD.

Defendant.

Case: 2:07-cv-13989  
Assigned To: Taylor, Anna Diggs  
Referral Judge: Pepe, Steven D  
Filed: 09-20-2007 At 03:27 PM  
CMP ELI LILLY AND CO V. SUN PHARMAC  
EUTICAL IND INC (TAM)

\_\_\_\_\_  
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**COMPLAINT**

Plaintiff Eli Lilly and Company, (hereinafter "Lilly") for its Complaint against Defendant Sun Pharmaceutical Industries, Ltd. (hereinafter "Sun"), hereby alleges as follows:

**NATURE OF THE ACTION**

1. This is a civil action for the infringement of United States Patent No. 5,658,590 ("the '590 patent"). This action relates to an Abbreviated New Drug Application ("ANDA") filed by Sun with the United States Food and Drug Administration ("FDA") for approval to market generic versions of Lilly's Strattera<sup>®</sup> drug products. This action arises under the patent laws of the United States, 35 U.S.C. § 100, *et seq.*

**PARTIES**

2. Plaintiff Eli Lilly and Company is an Indiana corporation having a principal place of business at Lilly Corporate Center, Indianapolis, Indiana 46285.

3. Upon information and belief, Defendant Sun is a corporation organized under the laws of India having a principal place of business at Acme Plaza, Andheri Kurla Road, Andheri (East), Mumbai 400-059, India.

**JURISDICTION AND VENUE**

4. This Court has jurisdiction over the subject matter of this action pursuant to 28 U.S.C. §§ 1331, 1338(a), 2201, and 2202.

5. Upon information and belief, this Court has personal jurisdiction over Sun. Upon information and belief, Sun is in the business of manufacturing, marketing, importing, and selling pharmaceutical drug products, including generic drug products. Upon information and belief, Sun directly, or through its subsidiaries Sun Pharmaceutical Industries, Inc., a Michigan corporation (hereinafter "Sun Michigan") and Caraco Pharmaceutical Laboratories, Ltd., a Michigan corporation with a principal place of business in Detroit, Michigan (hereinafter "Caraco"), manufactures, markets, and sells generic drugs throughout the United States and in this judicial district. Upon information and belief, Sun has purposefully conducted business, directly and through its subsidiaries, agents, and alter egos (including Sun Michigan and Caraco), in this judicial district, and this judicial district is a likely destination of Sun's ANDA product. Upon information and belief, Sun, through Sun Michigan and Caraco, owns and leases facilities in this judicial district. Upon information and belief, Sun has previously submitted to the jurisdiction of this Court and has further previously availed itself of this Court by asserting counterclaims in other civil actions initiated in this jurisdiction.

6. Venue is proper in this judicial district pursuant to, *inter alia*, 28 U.S.C. §§ 1391(b) and/or 1400(b).

**PLAINTIFF'S STRATTERA® PRODUCTS AND RELATED PATENT**

7. On August 19, 1997, the '590 patent, titled "Treatment of Attention-

Deficit/Hyperactivity Disorder,” was duly and legally issued to John H. Heiligenstein and Gary D. Tollefson and assigned to Eli Lilly and Company. A true and correct copy of the ‘590 patent is attached hereto as Exhibit A. The ‘590 patent claims methods of treating attention-deficit/hyperactivity disorder with tomoxetine. Tomoxetine is now known as atomoxetine. The claims of the ‘590 patent are valid and enforceable. The ‘590 patent expires on November 26, 2016.

8. Strattera<sup>®</sup> is the brand name for the commercial formulation of atomoxetine hydrochloride developed, manufactured, and sold by Lilly. Lilly submitted a New Drug Application to the FDA for Strattera<sup>®</sup> Capsules for the treatment of attention-deficit/hyperactivity disorder (NDA No. 21-411). NDA 21-411 was approved by the FDA on or about November 26, 2002, for Strattera<sup>®</sup> Capsules in strengths of Eq 10 mg, 18 mg, 25 mg, 40 mg, and 60 mg. Strattera<sup>®</sup> Capsules in strengths of Eq 80 mg and 100 mg were approved on or about February 14, 2005.

9. The Food And Drug Administration Center For Drug Evaluation And Research Approved Drug Products With Therapeutic Equivalence Evaluations (the “Orange Book”) lists the ‘590 patent for each of the strengths of Strattera<sup>®</sup> approved by the FDA under NDA No. 21-411.

10. Pursuant to 21 U.S.C. § 355a, Lilly is entitled to a six-month period of pediatric exclusivity for Strattera<sup>®</sup> beyond the date of expiration of the ‘590 patent.

#### **SUN’S ANDA FILING**

11. By letter dated August 6, 2007 (the “Sun Notice Letter”), Sun notified Lilly that Sun had submitted ANDA No. 79-020 to the FDA under § 505(j) of the Federal Food, Drug and Cosmetic Act, 21 U.S.C. § 355(j) (the “Sun ANDA”). On information and belief, the Sun ANDA seeks approval to engage in the commercial manufacture, use or sale of generic

Atomoxetine Hydrochloride Capsules, Eq 10 mg, 18 mg, 25 mg, 40 mg, 60 mg, 80 mg, and 100 mg Atomoxetine (collectively the “Sun Atomoxetine Capsules”) — generic versions of each of the FDA-approved Strattera<sup>®</sup> Capsule strengths — before the expiration date of the ‘590 patent.

12. By filing the Sun ANDA, Sun has necessarily represented to the FDA that the Sun Atomoxetine Capsules have the same active ingredient as Strattera<sup>®</sup>, have the same route of administration, dosage form, and strengths as Strattera<sup>®</sup>, are bioequivalent to Strattera<sup>®</sup>, and have the same or substantially the same proposed labeling and use as Strattera<sup>®</sup>.

13. In the Sun Notice Letter, Sun notified Lilly that the Sun ANDA contains a paragraph IV certification with respect to the ‘590 patent. Sun attached to the Sun Notice Letter a statement asserting its opinion that the ‘590 patent is invalid, unenforceable, or will not be infringed by the Sun Atomoxetine Capsules.

14. This action is being brought before the expiration of forty-five days from the date Lilly received the Sun Notice Letter, which Lilly received no earlier than August 7, 2007.

### COUNT I

#### INFRINGEMENT OF THE ‘590 PATENT

15. Lilly incorporates the preceding paragraphs as if fully set forth herein.

16. Sun’s submission of the Sun ANDA to obtain approval to engage in the commercial manufacture, use, offer to sell, or sale of the Sun Atomoxetine Capsules prior to the expiration of the ‘590 patent constitutes infringement of one or more of the valid claims of the ‘590 patent under 35 U.S.C. § 271(e)(2)(A).

17. Sun’s commercial manufacture, use, offer to sell, sale, or importation of the Sun Atomoxetine Capsules for the treatment of attention-deficit/hyperactivity disorder prior to the expiration of the ‘590 patent, and its inducement of or contribution to such conduct, would

further infringe the '590 patent under 35 U.S.C. §§ 271(a), (b) and/or (c). Sun's filing of the Sun ANDA and its intention to engage in the commercial manufacture, use, offer to sell, sale, or importation of the Sun Atomoxetine Capsules for the treatment of attention-deficit/hyperactivity disorder, and its intention to induce such conduct upon receiving FDA approval, create an actual case or controversy with respect to infringement of the '590 patent.

18. Upon FDA approval of the Sun ANDA, Sun will infringe the '590 patent by making, using, offering to sell, selling, or importing the Sun Atomoxetine Capsules in the United States for the treatment of attention-deficit/hyperactivity disorder, and by actively inducing and/or contributing to infringement by others, unless enjoined by this Court.

19. Lilly will be irreparably harmed if Sun's infringement is not enjoined. Lilly does not have an adequate remedy at law.

**PRAYER FOR RELIEF**

**WHEREFORE**, Lilly prays that this Court grant the following relief:

- A. A declaration that the '590 patent is valid and enforceable;
- B. A declaration that a claim or claims of the '590 patent are infringed by the manufacture, use, sale, offer for sale or importation of the Sun Atomoxetine Capsules, that Sun's submission of the Sun ANDA is an act of infringement of the '590 patent, that making, using, offering to sell, selling, or importing the Sun Atomoxetine Capsules for the treatment of attention-deficit/hyperactivity disorder by Sun, and its inducement of and/or contribution to such conduct by others, will infringe the '590 patent;
- C. An Order providing that the effective date of any approval of the Sun ANDA shall be a date which is not earlier than six months after the expiration of the '590 patent;
- D. An Order permanently enjoining Sun and its affiliates and subsidiaries, and each

of their officers, agents, servants, and employees, from making, using, offering to sell, selling, or importing the Sun Atomoxetine Capsules and from inducing or contributing to such conduct by others, until after six months after the expiration of the '590 patent;

E. An Order that damages or other monetary relief be awarded to Lilly if Sun engages in the commercial manufacture, use, offer to sell, sale, or importation of the Sun Atomoxetine Capsules, or in inducing or contributing to such conduct by others, prior to six months after the expiration of the '590 patent, and that any such damages or monetary relief be trebled and awarded to Lilly with prejudgment interest;

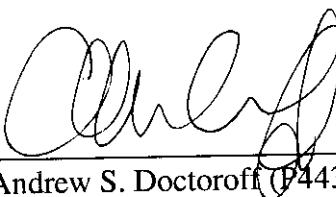
F. Reasonable attorneys fees, filing fees, and reasonable costs of suit incurred by Lilly in this action; and

G. Such further and other relief as this Court deems proper and just.

Respectfully Submitted,

**HONIGMAN MILLER SCHWARTZ  
AND COHN LLP**

BY: \_\_\_\_\_



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Dated: September 20, 2007

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