

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration Rockville MD 20857

MEMORANDUM

DATE:

September 12, 2006

TO:

Randall Lutter, Ph.D.

Associate Commissioner for

Policy and Planning

Food and Drug Administration

THROUGH: Jenny Slaughter

Director, Ethics and Integrity Staff

Office of Management Programs

Office of Management

FROM:

Igor Cerny, Pharm.D. /5/

Director, Advisors and Consultants Staff Center for Drug Evaluation and Research

SUBJECT: Conflict of Interest Waiver for Edmund Capparelli,

Pharm.D.

I am writing to request a waiver for Edmund Capparelli, Pharm.D., a consultant to the Center for Drug Evaluation and Research, from the conflict of interest prohibitions of 18 U.S.C. §208(a). The appointing official may grant waivers under section 208(b) (3) where "the need for the individual's services outweighs the potential for a conflict of interest created by the financial interest involved" and where the individual has made a disclosure of the financial interests at issue. We have determined that you are the appointing official for purposes of section 208. Therefore, you have the authority to grant Dr. Capparelli, a waiver under 18 U.S.C. §208(b) (3).

Section 208(a) prohibits Federal executive branch employees, including special Government employees, from participating personally and substantially in matters in which the employee, or any other person whose interests are imputed to the employee under 18 U.S.C. §208, has a financial interest. Since Dr. Capparelli is a special Government employee, he is under a statutory obligation to refrain from participating in an official capacity in any particular matter having a direct

and predictable effect on a financial interest attributable to him, his spouse, minor child, or general partner; an organization or entity for which he serves as an officer, director, trustee, general partner, or employee; and, a person with whom he is negotiating for, or has an arrangement concerning, prospective employment.

Dr. Capparelli has been asked to participate in the Clinical Pharmacology Subcommittee, of the Advisory Committee for Pharmaceutical Science, meeting where the subcommittee will: (1) hear an update on previous CPSC meeting recommendations and receive an introduction to the three new topics of this meeting; and, (2) discuss and provide comments on the evaluation of transporter-based drug interactions. These issues are particular matters of general applicability. Particular matters of general applicability focus on a discrete and identifiable class of persons, but do not involve specific parties.

In addition, Dr. Capparelli has been asked to participate in the subcommittee's discussion and to provide comments on the scope and strength of evidence to support the inclusion of pharmacogenetic information on CYP2D6 polymorphism in a revision of the label for tamoxifen to improve the benefit/risk of the drug. This issue is a particular matter involving specific parties.

The function of the Advisory Committee for Pharmaceutical Science, as stated in its charter, is to provide advice on scientific and technical issues concerning the safety, and effectiveness of human generic drug products for use in the treatment of a broad spectrum of human diseases, and as required, any other product for which the Food and Drug Administration has regulatory responsibility, and make appropriate recommendations to the Commissioner of Food and Drugs. The Committee may also review Agency sponsored intramural and extramural biomedical research programs in support of FDA's generic drug regulatory responsibilities.

Dr. Capparelli has advised the Food and Drug Administration that he has financial interests that could potentially be affected by his participation in in the matters to be discussed by the subcommittee.

Dr. Capparelli is a consultant to ______ regarding the pharmacokinetic analysis of intravenous ______ in preterm infants. He is also a member of ______ 's Data Safety Monitoring Board (DSMB) regarding a pharmacokinetic study of ______ in pregnancy. _____ (trade name ______) an anti-HIV drug. _____ and ____ could be affected by the subcommittee's discussions of (1) an update on the previous Clinical Pharmacology Subcommittee meeting recommendations and an introduction to the three new topics; and, (2) the evaluation of transporter-based drug interactions.

In addition, ____ makes _____, a competing product to tamoxifen.

As a consultant advising the Clinical Pharmacology Subcommittee of the Advisory Committee for Pharmaceutical Science, Dr. Capparelli potentially could become involved in matters that could affect his financial interests. Under 18 U.S.C. §208(a), he is prohibited from participating in such matters. However, as noted above, you have the authority under 18 U.S.C. §208(b)(3) to grant a waiver permitting Dr. Capparelli to participate in such matters as you deem appropriate.

For the following reasons, I believe that it would be appropriate for you to grant a waiver to Dr. Capparelli, which would permit him to participate in the matters previously described.

First, with respect to the subcommittee's discussions of (1) an update on the previous Clinical Pharmacology
Subcommittee meeting recommendations and an introduction to the three new topics; and, (2) the evaluation of transporter-based drug interactions, this waiver is justified, in part, because of the general nature of particular matters of general applicability. It is well recognized that particular matters of general applicability pose far less risk of a conflict of interest. Particular matters of general applicability include regulations, points-to-consider, guidelines, and policies governing classes of individuals, products, and organizations. Particular matters of general applicability do not include particular matters involving specific parties, such as recommendations regarding a specific product, or enforcement

matters involving known parties. Particular matters of general applicability will not have a special or distinct impact on any of Dr. Capparelli's financial interests in _____ or ____ other than as part of a class.

In addition, Dr. Capparelli's financial interests are not so substantial as to preclude his participation. He receives nominal compensation.

Further, the Federal Advisory Committee Act requires that committee memberships be fairly balanced in terms of the points of view represented and the functions to be performed by the various advisory committee members and Dr. Capparelli's participation will contribute to the balance of views represented and the diversity of opinions and expertise. The Committee's intended purpose would be significantly impaired if the agency could not call upon experts who have become eminent in their fields, notwithstanding the financial interests and affiliations they may have acquired as a result of their demonstrated abilities. Dr. Capparelli is Associate Clinical Professor of Pediatrics and Co-Director, Pediatric Pharmacology Research unit at University of California, San Diego. Dr. Capparelli's research focuses on pediatric pharmacology and pharmacokinetics. He has written over 100 publications and abstracts related to his field. Dr. Capparelli is a member of various professional societies, such as the American Society of Hospital Pharmacists, American College of Clinical Pharmacy, California Society of Hospital Pharmacists, American Society for Clinical Pharmacology and Therapeutics, and American College of Clinical Pharmacology. I believe that Dr. Capparelli's participation will help provide a foundation for developing advice and recommendations that are fair and comprehensive.

Page 5 - Randall Lutter, Ph.D.

Food and Drug Administration

Accordingly, I recommend that you grant Edmund Capparelli, Pharm.D., a waiver that will permit him to participate in the Clinical Pharmacology Subcommittee, of the Advisory Committee for Pharmaceutical Science, meeting where the committee will: (1) hear an update on previous CPSC meeting recommendations and receive an introduction to the three new topics of this meeting; (2) discuss and provide comments on the evaluation of transporter-based drug interactions; and, the subcommittee's discussion on the scope and strength of evidence to support the inclusion of pharmacogenetic information on CYP2D6 polymorphism in a revision of the label for tamoxifen to improve the benefit/risk of the drug. I believe that such a waiver is appropriate because in this case, the need for the services of Dr. Capparelli outweighs the potential for a conflict of interest created by the financial interests attributable to him.

CONCURRENCE:	<i>[s/</i>			9/22/06	
	Jenny Slaughter				Date
	Director, Ethi	ics and	Integrity	Staff	
	Office of Mana	agement	Programs		
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DECISION:					
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