RUTGERS UNIVERSITY
Office of Research and Sponsored Programs
ASB III, 3 Rutgers Plaza, Cook Campus
New Brunswick, NJ 08901

January 17, 2013

P.I. Name: Kaminski
Protocol #: E13-411

Kimberly Kaminski
Department of Pharmacy Practice & Administration
William Levine Pharmacy Bldg.
160 Frelinghuysen Road
Busch Campus

Dear Kimberly Kaminski:

Notice of Exemption from IRB Review

Protocol Title: "Resources Used for Obtaining Drug and Medical Information by Pharmacy Medical and Nursing Students"

The project identified above has been approved for exemption under one of the six categories noted in 45 CFR 46, and as noted below:

Exemption Date: 1/9/2013 **Exempt Category:** 2

This exemption is based on the following assumptions:

- **This Approval** - The research will be conducted according to the most recent version of the protocol that was submitted.
- **Reporting** – ORSP must be immediately informed of any injuries to subjects that occur and/or problems that arise, in the course of your research;
- **Modifications** – Any proposed changes MUST be submitted to the IRB as an amendment for review and approval prior to implementation;
- **Consent Form (s)** – Each person who signs a consent document will be given a copy of that document, if you are using such documents in your research. The Principal Investigator must retain all signed documents for at least three years after the conclusion of the research;

Additional Notes: **None**

Failure to comply with these conditions will result in withdrawal of this approval.

The Federalwide Assurance (FWA) number for Rutgers University IRB is FWA00003913; this number may be requested on funding applications or by collaborators.

Sincerely yours,

m/ Margaret Cooper
Acting for

Sheryl Goldberg
Director of Office of Research and Sponsored Programs
gibel@grants.rutgers.edu

cc: Michael Tuscani

ASSENT

You are invited to participate in a research study that is being conducted by Kimberly Kaminski and Alla Shatskov who are Post-Doctoral Pharm.D. Fellows in the Pharmacy Practice Department at Rutgers University. The purpose of this research is to determine the various clinical information resources pharmacy, medical and nursing students use for self-education and clinical decision making. The link below contains a brief seven question survey which will assess drug information collection preferences and resources when obtaining information regarding dosing, adverse events, drug interactions and disease state information.

This research is anonymous. Anonymous means that I will record no information about you that could identify you. This means that I will not record your name, address, phone number, date of birth, etc. If you agree to take part in the study, you will be assigned a random code number that will be used on each test and the questionnaire. Your name will appear only on a list of subjects, and will not be linked to the code number that is assigned to you. There will be no way to link your responses back to you. Therefore, data collection is anonymous.

The research team and the Institutional Review Board at Rutgers University are the only parties that will be allowed to see the data, except as may be required by law. If a report of this study is published, or the results are presented at a professional conference, only group results will be stated. All study data will be kept for three years.

There are no foreseeable risks to participation in this study. In addition, you may receive no direct benefit from taking part in this study.

Participation in this study is voluntary. You may choose not to participate, and you may withdraw at any time during the study procedures without any penalty to you. In addition, you may choose not to answer any questions with which you are not comfortable.

If you have any questions about the study or study procedures, you may contact:

Kimberly Kaminski, PharmD
Rutgers Pharmaceutical Industry Fellowship Program
160 Frelinghuysen Road, Room 405
Piscataway, NJ 08854
Phone: 609-897-3809
Kimberly.kaminski@rutgers.edu

If you have any questions about your rights as a research subject, you may contact the IRB Administrator at Rutgers University at:

Rutgers University, the State University of New Jersey
Institutional Review Board for the Protection of Human Subjects
Office of Research and Sponsored Programs
3 Rutgers Plaza
New Brunswick, NJ 08901-8559
Tel: 848-932-0150
Email: humansubjects@orsp.rutgers.edu

You will be given a copy of this assent form for your records.

If you are 18 years of age or older, understand the statements above, and will consent to participate in the study, click on the "I Agree" button to begin the survey. If not, please click on the "I Do Not Agree" button which you will exit this program.

I Agree | I Do Not Agree

APPROVED
Date: 1/9/13