POLICY

1. Purpose
Describe the composition of the Research Subjects Review Board (RSRB), as well as the roles and responsibilities of the RSRB membership.

2. Scope
This policy applies to all members, administrators, and consultants of the Research Subjects Review Board.

3. Definitions
3.1. Board Chair – An individual with relevant scientific/clinical background and expertise in research methods who is appointed by the Institutional Official to lead convened meetings and conduct administrative duties required for a successful operation of the Board.

3.2. Acting Board Chair – An individual with relevant scientific/clinical background and expertise in research methods who is named by the Chair and/or the RSRB Director, to lead convened meetings and conduct specific administrative duties in the absence of the Chair. This individual must be an experienced board member in order to fulfill this role.

3.3. Experienced Board Member – A member who has demonstrated during a period of active participation a broad understanding and competency with human subject protection ethics, board operations, and regulatory requirements, including expedited review procedures and is so designated by the Chair.

3.4. Alternate Board Member – A member formally appointed and listed in the membership roster who may substitute for a primary member with whom the alternate has similar qualifications. Experienced members may be asked to continue as alternates when their terms expire or when their workload prevents them from carrying out their duties.

3.5. Consultant – An individual who may be requested to provide additional scientific and/or specialty expertise to the board as necessary. Consultants are not permissible as voting members.

3.6. Prisoner Representative – A member formally appointed and listed in the membership roster who has the appropriate background, experience and working knowledge to provide an understanding and appreciation of prison conditions from the prisoner's perspective.
4. References

4.1. HHS 45 CFR 46.107; HHS 45 CFR 46.303(c); HHS 45 CFR 46.304; FDA 21 CFR 56.107

4.2. Policy 401 Functions of the RSRB Office
Policy 501 Levels of RSRB Review
Policy 802 Non-Compliance

4.3. Guideline for Board Member Terms of Appointment

5. Responsibilities

5.1. The RSRB is responsible for serving the research community at the University of Rochester in the processing, review, and approval of research activities. The determination of which board will be responsible for project review is based on the department of the submitting PI, the nature of the research, and the prospective subject population (Policy 401 Functions of the RSRB Office).

5.1.1. The RSRB has five internal review boards, including three biomedical boards, an HIV/AIDS and oncology board, and a behavioral and social sciences board. In addition, Western IRB (WIRB) reviews industry-initiated, industry-sponsored, research of FDA-regulated drugs and devices posing greater than minimal risk. Additional information can be found in Policy 401 Functions of the RSRB Office.

5.1.2. There is a designated RSRB staff member responsible for review and determination of exemption requests (Policy 501 Levels of RSRB Review).

5.2. The OHSP Director and RSRB Director are responsible for recommending to the Institutional Official (IO) individuals to serve as Board Chairs.

5.2.1. Because the successful operation of the Board is critical to the Human Research Protection Program, the Chair should possess:

5.2.1.1. A broad knowledge regarding regulations for the protection of human research subjects, and the ability to apply these regulations consistently and appropriately.

5.2.1.2. A commitment to devoting sufficient time to ensure an efficient RSRB review process.

5.2.1.3. The ability to maintain a constructive, productive atmosphere at RSRB meetings.

5.2.1.4. The ability to maintain productive working relations with the RSRB office staff.

5.2.1.5. A commitment to maintaining a current knowledge of the human subject protection process.
5.3. The OHSP Director and the RSRB Director, in consultation with the respective Board Chair, are responsible for identifying and recommending board member appointment and re-appointment to the IO.

5.3.1. The OHSP or the RSRB Director may request a representative from the Chair of a department that submits a high volume of studies.

5.3.2. An individual may express interest in board membership (self-nomination).

5.3.3. An individual may be referred by RSRB members, RSRB office staff, or OHSP staff.

5.3.4. Appointments of alternate members may be made in the same manner and for the same terms as primary members.

5.3.5. Selection of RSRB members by research investigators is not permitted.

5.4. The **Board Chair** is responsible for overseeing RSRB review of research, including but not limited to the following:

5.4.1. Apply the basic ethical principles of respect for persons, beneficence and justice as articulated in the Belmont Report.

5.4.2. Uphold federal, state, and local regulations, University policies and procedures and with regard to the protection of human research subjects.

5.4.3. Be knowledgeable about, and have understanding of, ethical issues, federal research regulations, applicable state law, and Institutional policies.

5.4.4. Involvement as needed in discussions with federal authorities.

5.4.5. Promote a culture consistent with the objectives of the University of Rochester’s Human Research Protection Program (HRPP), with special emphasis on the respect for and protection of individuals participating in research

5.4.6. Identify the expertise needed to comprise the RSRB and recommend potential members.

5.4.7. Direct the discussion and proceedings of the full-committee meetings, with the assistance of the RSRB Specialist, to keep discussion focused on the established agenda and ensure that guidelines for meeting procedures are followed.

5.4.8. Review protocols submitted for initial and continuing reviews as well as any amendments submitted.

5.4.9. Conduct review of research qualifying for exemption, as needed or as requested by RSRB staff.

5.4.10. Review projects that qualify as expedited (this task may be delegated to an experienced board member).

5.4.11. Review reportable events and take appropriate action as needed. This includes the authority of the Chair to take immediate action to suspend approval of a research project in order to protect research subjects from serious risks.

5.4.12. Ensure prompt reporting to the RSRB, institutional officials, and others, as needed, regarding research related events/problems, serious or continuing non-compliance with UR HRPP policy or the requirements or determinations of the RSRB, and any suspension or termination of RSRB approval.
5.4.13. Determine if a consultant is needed to supplement the expertise of the RSRB (this task may be delegated to another member or RSRB staff, based on expertise and experience).
5.4.14. Vote at full board meetings.
5.4.15. Serve as the official signatory for RSRB approval letters (or designate to an experienced board member).
5.4.16. Participate in the investigation of suspected non-compliance and the development of a plan of action to address the non-compliance.
5.4.17. Designate individuals who have the expertise and experience to review protocols as sole designee for expedited reviews based on their experience with human subject research activities, credentials, and their experience reviewing protocols.

5.5. **RSRB members** are responsible for determining that the rights and welfare of human subjects in research are adequately protected, including but not limited to the following actions:

5.5.1. Apply his or her particular expertise, as well as the basic ethical principles of respect for persons, beneficence and justice as articulated in the Belmont Report, to the review of research.


5.5.3. Conduct reviews as requested by the Chair/RSRB staff member and provide feedback on all assigned review materials.

5.5.3.1. A “primary reviewer” is responsible to review all the materials provided to the board and to lead the discussion and/or introduce the study at the convened meeting.

5.5.4. Communicate directly with investigators and meet with them and study team as needed to advise on research and human subject protections.

5.5.5. Prepare for and actively participate in board meetings.

5.5.6. Meet HRPP education requirements (initial or continuing, as appropriate).

5.5.7. Report to the RSRB Director any perceived allegations of undue influence on the actions of RSRB members (Policy 802 Non-Compliance).

5.6. **Alternate RSRB members** will apply all the responsibilities of regular members when called upon by the Chair to serve at a committee meeting. The alternate member receives and reviews the same material that the primary member would have received. Alternates may be designated as a replacement for a specific member with particular expertise or may be appointed for their general area of competency. Alternate members may attend any or all RSRB meetings, but may only vote and exercise the privileges of a primary member at such meetings when they are expressly substituting for an absent primary member. Any member of a board can be an alternate on any of the other
boards as long as he/she is an appropriate alternate based upon affiliation, scientific status, and specialty.

5.6.1. If both the designated alternate and the regular member attend the same meeting, only the regular member may vote.

5.6.2. When an alternate member is used, the minutes will reflect who the alternate member is replacing.

5.7. **Consultants** are responsible for providing additional expertise that may be needed for reviewing a specific study as may pertain to his/her qualifications, scientific knowledge and ability to evaluate potential ethical concerns inherent to the study, potential risks or benefits of the study procedures, or concerns relative to the study population particularly when vulnerable subjects are involved.

5.8. The IO, in consultation with the OHSP Director and RSRB Director, may act to remove a member of the RSRB, including a Board Chair, before the end of his/her term. This may occur if his/her participation in RSRB activities is deemed to be inadequate, inappropriate, or damaging to the reputation of the University and its research activities.

5.8.1. Removal of an RSRB member from service may occur under the concurrence of the RSRB Director and OHSP Director only.

5.8.2. Removal of an RSRB Chair requires the concurrence of the IO and the OHSP Director and RSRB Director.

5.8.3. Members cannot be removed from RSRB because of their voting record, or in an attempt to alter the RSRB’s membership for purposes of obtaining approval for a certain protocol or class of protocols.

6. **Requirements**

6.1. **Research Subject Review Board Membership Composition**

6.1.1. The membership of the RSRBs shall comply with federal regulations regarding membership composition [HHS 45 CFR 46.107; HHS 45 CFR 46.304; FDA 21 CFR 56.107] as follows:

6.1.1.1. Each RSRB shall be composed of no less than five members who are qualified through their experience and expertise to review research projects in terms of regulations, ethical principles, applicable laws, standards of professional conduct and practice, and Institutional commitment.

6.1.1.2. Each RSRB shall consist of members of various professions including at least one scientist, one non-scientist, and one member who is not otherwise affiliated with the Institution (e.g., a community member), and who is not part of the immediate family of a person who is affiliated with the institution.
6.1.1.3. Each RSRB shall reflect diversity in its membership in terms of experience and expertise and consider race, gender, cultural backgrounds, and sensitivity to such issues as community attitudes as factors in membership selection.

6.1.1.4. No RSRB shall consist entirely of men or entirely of women.

6.1.1.5. No RSRB shall consist entirely of members of one profession.

6.1.1.6. Each RSRB shall include representatives who are knowledgeable about and experienced in working with various vulnerable populations/groups such as children, prisoners, pregnant women or mentally or physically challenged individuals if research involving such populations is regularly reviewed by that board. This information is documented using the RSRB Board Member Representative Capacity form (Appendix 8).

6.1.1.7. Each RSRB may include representation from affiliated institutions as appropriate.

6.1.1.8. No RSRB shall consist of members (or alternates) who are responsible for business development of the organization, nor should these individuals carry out any day-to-day operations of the review process (e.g., Senior VP for Research).

6.1.2. New members are appointed for an initial term of one year at which time members are evaluated for continued service. Members will also be evaluated prior to the end of each subsequent term of service and Chairs will be evaluated on an annual basis to determine ongoing membership (see Guideline for Board Member Terms of Appointment). Members receive an appointment letter (see Appendix 3) to confirm membership on the RSRB and a re-appointment letter (see Appendix 4) to confirm continued service. Members are also acknowledged in writing at the end of their service as a board member (see Appendix 5).

6.1.3. Names and qualifications of board members, including changes in membership, are maintained in membership rosters as required by OHRP (Appendix 6).

6.1.3.1. Information will be maintained by the RSRB office to document member expertise with vulnerable populations and representation by community members.

6.1.3.2. The RSRB Director (or designee) will notify OHRP of changes in RSRB membership at least annually through on-line submission of an updated roster.

6.1.3.3. The RSRB will not disclose its membership to researchers or sponsors (Appendix 1: Sponsor Request for RSRB Membership Information).

6.1.3.4. Board members will be indemnified by the University for actions taken within the scope of duties as members of the RSRB. Indemnification is expressly extended to non-affiliated members of the RSRB while they are performing the duties of an RSRB member.
6.2. Consultants

6.2.1. The Director of OHSP, Director of RSRB, or the board may determine whether additional expertise from an outside reviewer is required for the review of a particular protocol. Consultants may be from within the University or outside the institution.

6.2.2. Prior to receipt of review materials, the consultant will review and sign the Confidentiality Agreement for Consultants (Appendix 7).

6.2.3. The consultant will be provided material relevant to the requested consultation and is expected to make recommendations regarding the study. Reviews provided in writing will be included in the meeting materials provided to board members.

6.2.4. If invited to attend the board meeting, consultant reviewer may participate in the deliberations, but may not vote.

6.2.5. Meeting minutes will document attendance of a consultant at board meetings as “guest”, indicate the specific study(ies) the consultant is reviewing, state any potential conflicting interests that may exist for the consultant, and capture any key information provided by the consultants.

6.3. Member Representation for Review of Vulnerable Populations

6.3.1. A prisoner representative with appropriate background and experience is assigned to each internal board to participate in the review of research involving incarcerated individuals. The prisoner representative will be appointed to the board in the same fashion as primary members.

6.3.2. A special board including the prisoner representative will meet to consider initial and continuing reviews of, as well as modifications to, research involving incarcerated individuals. The prisoner representative may assist with the review as an additional primary reviewer, and/or provide specific information related to the prisoner population. For federally funded research, these reviews will apply to either pre-planned or subsequent involvement of prisoners in the research. For all other research, the special board will review studies that intend to involve prisoners (per inclusion criteria), but will not be required to re-review previously approved studies that enroll a prisoner as an incidental subject (e.g., general population clinical trial for which a prisoner is eligible, or a general population study where an enrolled subject subsequently becomes incarcerated). The RSRB Chair and staff may refer such studies to either the special board or to the prisoner representative for review if circumstances warrant. When the special RSRB reviews research that involves prisoners, a majority of the board (exclusive of the prisoner representative) will have no association with the prison(s) involved, apart from their membership on the RSRB.
6.3.3. Research involving children will be reviewed by a board that contains members with appropriate background and experience relevant to the research.

6.4. Initial Training and Education of RSRB of Prospective and New Members

6.4.1. Prospective members are provided with reference materials (e.g., Institutional Review Board Member Handbook) and meet with the RSRB Director to discuss the expectations of board membership and to assess qualifications.

6.4.1.1. The prospective member then will undergo regulatory training with the OHSP Director, or designee with appropriate regulatory experience and knowledge. This training provides an overview of human subject protection issues, including laws, federal regulations and ethical principles.

6.4.1.2. Additional training is provided by the Chair and Specialist of the Board to which the new member is assigned. This process provides further preparation in RSRB policies and procedures, how the meetings are conducted and how protocols are reviewed with the option to attend a meeting as a non-voting member.

6.4.1.3. Prospective members must also successfully complete the University’s Human Subjects Protection Program (HSPP). The RSRB office keeps track of members’ HSPP number, expiration dates, and sends out HSPP renewal reminder notices in conjunction with OHSP.

6.4.1.4. Completion of initial training is documented (see Appendix 2) to verify that all elements have been covered prior to appointment.

6.4.2. New members may serve as primary reviewers at the Chair’s discretion. The Chair, or another experienced reviewer, will support the new member in his/her first review as needed.

6.5. Ongoing Training and Education of RSRB Members

6.5.1. The Board Member Resources on the OHSP website provides members an area to access articles of interest, presentations from previous meetings, board meeting materials.

6.5.2. Ongoing interactions at meetings and individual discussions with staff, Chairs and members, as well as email communications distributed by OHSP or RSRB staff.

6.5.3. Attendance at university-sanctioned events pertaining to human subject protection, such as presentations or audio conferences.

6.5.4. Brief educational presentations during board meetings.

6.5.5. Journal subscriptions provided by the RSRB office.

6.5.6. Attendance at the annual All Boards meeting or other training meetings provided through OHSP.

6.5.7. Attendance at the annual PRIM&R, or other national meetings, as selected.
Originator/Authors:
Kelley O'Donoghue, Director OHSP
Emily Flagg, Senior Regulatory Specialist

Appendices:
Appendix 1: Sponsor Request for IRB Membership Information
Appendix 2: Sample Board Member Initial Training Completion
Appendix 3: Sample Board Member Appointment Letter
Appendix 4: Sample Board Member Re-Appointment Letter
Appendix 5: Sample Board Member End of Service Letter
Appendix 6: RSRB Member and Alternate Member Roster Templates
Appendix 7: Confidentiality Agreement for Consultants Template
Appendix 8: RSRB Board Member Representative Capacity Form

Revision History:
Section 6.1.1.6 Add board member form; Appendix 6 update roster template; Add Appendix 8 (11/14)
Add Sect 6.1.1.8 per AAHRPP (01/15)

Supersedes Date:
11/03/2014

Approved By:
Robert Clark
Institutional Official and Senior VP for Research

Kelley A. O'Donoghue
Director, OHSP

Tiffany Gommel
Director, RSRB
Appendix 1: Sponsor Request for IRB Membership Information

RESEARCH SUBJECTS REVIEW BOARD

Sponsor Request for IRB Membership Information

Dear Principal Investigator/Sponsor:

This letter is in response to requests for information regarding the membership of the Research Subjects Review Board (RSRB), the IRB of record for the University of Rochester.

The University of Rochester has a Federalwide Assurance (#9386) with the Office for Human Research Protections (OHRA) in the Department of Health and Human Services (DHHS). The Federalwide Assurance is a binding agreement between DHHS and the University of Rochester that assures the federal government that the University will meet all requirements of Title 45, Part 46 of the Code of Federal Regulations (45 CFR 46) for all human subject research, regardless of sponsorship. As a requirement of the FWA, the RSRB meets both DHHS and FDA requirements for IRB membership as specified in Sections 45 CFR 46.107 and Section 21 CFR 56.107, respectively.

The Research Subjects Review Board complies with DHHS regulations regarding board member conflict of interest. Investigators do not participate in RSRB board meetings except by invitation, and then only to provide information requested by the board.

Additionally, RSRB board members leave the board room prior to discussion or voting on any study in which they may have a real or perceived conflict of interest.

By University policy, the RSRB membership list is not disclosed to sponsors or other external parties. Membership lists are filed as part of the University’s FWA.

If you have any other questions about board membership, please contact the RSRB at 585-275-2398.

Tiffany L. Gemmel, MS, CIP, CIP
Executive Director
Research Subjects Review Board

265 Crittenden Blvd • Suite 1.250 • Box #426628 • Rochester, NY 14642
585.273.2398

Revised: 03/13/2013
Appendix 2: Sample Member Initial Training Completion Form

RESEARCH SUBJECTS REVIEW BOARD

RSRB Member Initial Training Completion Form

In your role as a member of the RSRB, your primary responsibility is to protect the rights and welfare of research subjects. To ensure that you can effectively perform your responsibilities as a Board Member, you will need to be familiar with the following:

- Belmont Report and the ethical principles applicable to research with human subjects
- Health and Human Services (HHS) Regulations
- Food and Drug Administration (FDA) Regulations
- RSRB Procedures
- University Policies
- RSRB Online Submission System (ROSS)

In conjunction with your initial appointment to the Board, the RSRB office will provide you with the following training sessions. Please indicate the date the training was completed and initial each item confirming that you participated in that training. Continuing education, refresher training and opportunities to attend conferences or training sessions are also offered to Board Members.

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<th>Confirming Initials</th>
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<tr>
<td>RSRB Procedures (accessible online) and Board Operational Training</td>
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<td>ROSS Training</td>
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<tr>
<td>Human Subject Protection Certification (HSPP) Including Belmont Report and the ethical principles applicable to research with human subjects</td>
<td>3/31/2012</td>
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</tbody>
</table>

Your signature below verifies that you have received training in all of the above areas. You have also successfully completed the Human Subjects Protection Program (HSPP). This form will be kept in your RSRB member file with your other credential information.

Printed Name

Eligibility for Voting Membership

Kelley O’Conner, MHA
Executive Director
Research Subjects Review Board Office
Effective Date: 08/01/2010

Signature

Date: 5/19/2011

Paper copies of the Policy may not be the current version. The current version of this Policy is maintained and available on the OHSP shared network.
Appendix 3: Sample Board Member Appointment Letter

Office of the Senior Vice President for Research

March 11, 2013

[Name/Address]

Dear Ms. [Name],

It is my pleasure to appoint you as a scientific member of the Research Subjects Review Board for a one-year term effective January 22, 2013.

I appreciate your willingness to serve the University in this capacity and I thank you for your contributions.

Sincerely yours,

Robert L. Clark
Senior Vice President for Research
Appendix 4: Sample Board Member Re-Appointment Letter

September 25, 2012

[Name/Address]

Dear Mr. [Name],

It is my pleasure to reappoint you as a scientific member to Board 02 (Behavioral) of the Research Subjects Review Board for a three-year term effective August 18, 2012.

I appreciate your participation as a board member and want to thank you for your 13 years of service to the University of Rochester RSRB. I thank you for your continued contribution.

Sincerely yours,

[Signature]

Robert L. Clark
Interim Senior Vice President for Research
Appendix 5: Sample Board Member End of Service Letters

Office of the Provost
Ralph W. Kunel, PhD, MD
Provost and Executive Vice President

July 7, 2011

Eileen Blakely, RD, CDN
Pediatrics Genetics
Box 777

Dear Ms. Blakely:

In keeping with the policy of the Research Subjects Review Board (RSRB), I review the
status of individuals whose membership on this committee is due to expire.

Initial appointments are made to the board for a period of one year, with reappointments
to be made at my discretion every three years thereafter. According to records in the RSRB
Office, your third three-year appointment to this committee will expire on July 8, 2011. I am

because of the importance, to our faculty and the committee alike, of service on this vital, but
time-consuming committee. I view reappointment beyond the second three-year renewal to be highly
unusual and you have been dedicated enough to serve an additional 3-year term. Consistent with
that, at the conclusion of your current term, your participation on this committee is completed
with our thanks.

I would like to acknowledge your contributions to the RSRB, the University of
Rochester, and the rights of research participants for the last ten years. This is a very
important committee for our research subjects and our faculty, and on their behalf, I thank you.

Sincerely,

Ralph Kunel, PhD, MD
 Provost and Executive Vice President

cc: G. Chadwick
RSRB Administrative files

Office of the President
Joel Seligman
200 North W. Rd.
Rochester, NY 14627-5011
585.275.8018, 585.275.3873/4
seligman@rochester.edu

July 26, 2011

Eileen Blakely, RD, CDN, MS
Division of Pediatrics Genetics
Gelmano Children’s Hospital
Box 777

Dear Ms. Blakely:

Provost Kunel has informed me that you will soon be retiring off the University’s
Research Subjects Review Board after completing ten years of service. The Board is a vital and
time-consuming committee, and for you to have given service as a community member
representing the participant population for all of those years is commendable.

Your dedication to the rights of research participants by serving on this Board has been
critical to the clinical research at the University of Rochester. Without committed individuals
like you, this research would not be possible. I am deeply grateful for your dedicated service.

Sincerely,

Joel Seligman
President

cc: Ralph Kunel
Gary Chadwick
Steve Lambert

Paper copies of the Policy may not be the current version. The current version of this Policy is maintained and available on the OHSP shared network.
Appendix 6: RSRB Member and Alternate Member Roster Templates

**University of Rochester**

**IRB Registration**

**Board 0X - IRBREG#**

**Board 0XP - IRBREG#**

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<th>Earned Degree(s)</th>
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<th>Primary Specialty</th>
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**KEY**

* = Chair
** = Alternate Chair
*** = Prisoner Representative (note all other board members are included on Board 0X and Board 0XP)
^ = For representative capacity and experience description, see the RSRB Member Representation Form on file.

No Affiliation = Neither the member nor any of the member’s immediate family member are affiliated with the institution.

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**HHS Assurance No. FWA0000365**

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**Alternate Members for all University of Rochester RSRBs**

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<th>Gender (M/F)</th>
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**KEY**

** = Alternate Chair
^ = For representative capacity and experience description, see the RSRB Member Representation Form on file.
Appendix 7: Confidentiality Agreement for Consultants Template

Office for Human Subject Protection

Research Subjects Review Board (RSRB)
Confidentiality Agreement for Consultants

I, ______________________, the undersigned consultant to the RSRB, hereby acknowledge that I understand that certain information acquired by me, in connection with my consultation to the RSRB, is confidential and I agree to maintain the confidentiality of such information. Examples of such information include: identifying data or medical information about human subjects; proprietary information; trade secrets, medical or scientific data; research designs, concepts, discoveries or inventions; or any other information designated as, or which is reasonably understood to be, confidential. I agree that I will not disclose, or reproduce in any fashion, any confidential information that I obtain in connection with my consultation to the RSRB except as may be required by law.

I understand that one of the reasons for not disclosing information I obtain in connection with my consultation to the RSRB is to protect the privacy of research subjects and the confidentiality of their medical information. In addition, I understand that some information that is set forth in research protocols or other documents submitted to the RSRB may contain the proprietary or intellectual property of an investigator or a third party sponsor, and that disclosure of such information by me could result in irreparable harm to the investigator or the funding sponsor.

Signature ______________________  Date ______________

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Paper copies of the Policy may not be the current version. The current version of this Policy is maintained and available on the OHSP shared network.
Appendix 8: RSRB Board Member Representative Capacity Form

This form is intended to identify the range of experience and knowledge of Research Subjects Review Board members that may be applied during the review of research to ensure appropriate representation and expertise in review meetings.

Representative Capacity: Indicate below the group(s) or population(s) with whom you have knowledge or experiences to bring to the Board to address as appropriate.

- Children (under 18 years)
- Research subjects – i.e., subject advocates or personal experience (and or family)
- Ed 12 students
- Terminally ill (life expectancy ≤ 6 months)
- UR students (under 18 years)
- Nursing home residents
- UR students (18 years and over)
- Limited or uncooperative
- UR employees
- Developmentally impaired
- Pregnant women
- Economically disadvantaged
- Prisoners
- N/A (no additional representation to note)

Indication of Experience: Provide a brief description of all relevant experiences contributing to the representative group(s) noted above (e.g., profession, URI experience with research in vulnerable populations, certification, license).

[Signature]
Board Member Name

[Signature]
Board Member Signature

Date: 6/30/2014

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