

Maryann Davis, Ph.D.
Department of Psychiatry
UMass/Memorial Health Care
Docket# H-11048

RE: NOTICE OF REAPPROVAL

Title:

Network analysis of transition services.

This letter is to certify that the project identified above has been re-reviewed by the Committee for the Protection of Human Subjects in Research. The Committee is recognized by the following FWA # 00004009.

The Committee has considered the following:

1. The Adequacy of the protection of the rights and welfare of the subjects involved.
2. The risks and potential medical benefits to the subjects in relation to the importance of the knowledge gained.
3. The adequacy and appropriateness of the methods used to secure informed consent.

The study was reapproved by expedited review on: _____

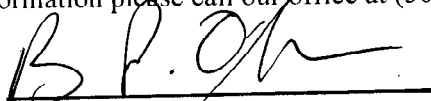
The study was reapproved at the Full IRB Committee meeting on: 8.16.05

The study approval expires on: 9.4.06

As the Principal Investigator for this study, you have the following obligations:

1. To report all serious and unexpected adverse reactions to the Human Subjects Committee within five working days. Any death of a subject, regardless of the cause, must be reported.
2. To use only an HSC-approved version of the informed consent. The approval is for no longer than a one year period and expires one year from the approval date listed above.
3. To give every subject a copy of the consent form and, if drugs are involved, to place a signed copy of the consent form in the patient's chart.
4. To obtain approval from the HSC before instituting any change in the protocol or the consent form.

Thank you for your cooperation with the Committee. If you have any questions on the above information please call our office at (508) 856-4261.


Brian P. O'Sullivan, M.D., Chair

Roger Luckmann, M.D., Vice Chair
Oren P. Schaefer, M.D., Vice Chair
Lucie Lajeunesse, BS, Alternate Vice Chair