From: Beers Block, Patricia
Sent: Thursday, June 06, 2002 1:42 PM
To: [purged]
Cc: OC GCP Questions
Subject: IRB and Informed Consent Regulations

Dear Dr. [purged],

As we discussed today, the regulations entitled "Protection of Human Subjects" (21 CFR 50) and "Institutional Review Boards" (21 CFR 56) can be found at the following web addresses respectively:

www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/showCFR.cfm?CFRPart=50

www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/showCFR.cfm?CFRPart=56

If you have any additional questions after reading these sections, please feel free to contact us either by email or by phone (301-827-3340)

Sincerely,

Pat Beers Block