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Questions, Comments, Suggestion?

If you have questions, comments or suggestions about how we can improve our human research protection program (HRPP) at UHCMC, send an email to: clinicalresearch@uhhs.com or contact Carol Fedor, Clinical Research Manager at (216) 844-5524

Education Updates!

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Contact Us

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FDA Requirements Regarding IRB Registration –

Both IRB committees for the UHCMC IRB (IRB00000684 and IRB00001691) are registered per the new Food and Drug Administration (FDA)'s registration requirements as reviewing FDA regulated research. Confirmation of this registration can be found by searching the UHCMC IRB Organizational Information number (IORG0004088) on the [HHS Office of Human Research Protections \(OHRP\) registration system](#) and clicking on the link for "U Hosps Case Med Ctr." Please note that both committees of the UHCMC IRB also review federally funded research and are also registered as such.



United States Department of Health & Human Services
THE OFFICE FOR HUMAN RESEARCH PROTECTIONS

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IRB Organization Information

IORG0004088 - U Hosps Case Med Ctr (Active)

Located at: Cleveland, OHIO
Expires: 09/23/2012

IRBs for this Organization: 2
[Agency Only Access](#)

| IRB# | IRB Name | City | State/Country | Status | IRB Type |
|-------------|---|-----------|---------------|--------|----------|
| IRB00000684 | U Hosps Case Med Ctr IRB #1 - (J Committee; Biomedical) - J Committee | Cleveland | OHIO | Active | OHRP/FDA |
| IRB00001691 | U Hosps Case Med Ctr IRB #2 - (M Committee; Biomedical) - M Committee | Cleveland | OHIO | Active | OHRP/FDA |

Human Subject Protection Program – Recertification Reminder

Individuals involved with human research need to be Human Subject Protection (HSP) certified. University Hospitals partners with Case Western Reserve University and utilizes the on-line Collaborative Institutional Training Initiative (CITI) Program. The training program certifies and enrolls individuals in the Human Subject Protection program.

As a reminder, many individuals entered the HSP program in 2004 when the CITI program was introduced at University Hospitals. You are responsible for ensuring you have earned enough CRECs (Continuing Research Education Credits) for recertification. **With recertification necessary every 3 years, a large number of individuals will need to recertify again around September 2010.** To view the IRB Policy, [Certification in Human Subjects Protections](#), [click here](#).

Helpful Tips

When recording information (data) in a research record, be certain to enter the "true value" (i.e., the actual data from the source). Down arrows and/or quotation marks should not be used to indicate repeat information.