

Correction to Biomedical Protocol Template

The FDA fax number in 11.7 of the *Adverse Event Reporting Requirements* section was changed to correct a typo. The correct fax number is 1-800-FDA-0178. The version date on the document header was also changed to 12/13/07.

This document is located on the Clinical Research Unit's section of the DF/HCC website under "Forms/Checklists/Templates." Note: You need to log in to this site with your DF/HCC user name and password.

For further information or questions, please contact:

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