**GHUCCTS Good Clinical Practice/Best Research Practices Seminar**

**October 26, 2017**

Please rate the extent to which you are now able to meet each of the training objectives by checking the box under the number that best reflects your opinion. (1 is lowest, 5 is highest)

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| --- | --- | --- | --- | --- | --- |
|  | **1** | **2** | **3** | **4** | **5** |
| Identify standards of Good Clinical Practice (GCP). |  |  |  |  |  |
| Understand the regulations/standards that form the framework of GCP. |  |  |  |  |  |
| Define source documentation and common pitfalls. |  |  |  |  |  |
| Identify the elements of a regulatory binder. |  |  |  |  |  |
| Understand the need for Delegation of Authority Logs and proper use. |  |  |  |  |  |
| Identify common errors in informed consent documentation. |  |  |  |  |  |
| Describe the elements of thorough adverse event assessment. |  |  |  |  |  |
| Describe when study SOPs should be developed. |  |  |  |  |  |
| Learn preliminary steps in preparing for a government inspection. |  |  |  |  |  |
| Identify three common audit findings and applicable corrective methods |  |  |  |  |  |

What new piece of information did you gain by attending this training? \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

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Will your current research practices be changed in any way after this training? \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Please list any topics for which you would like to receive additional training. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

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