Site Personnel Log	Protocol #/Title:

Investigator/Investigator #	Institution	Investigational Product Name/#	Center/Site #

List Individuals delegated significant study-related tasks (ICP GCP 4.1.5). Signatures/Initials required for all persons authorized to make entries and/or corrections to Case Report Forms (ICH GCP 8.3.24)

1) All personnel listed on From FDA 1572 must be listed on this form. In addition, you may need to list other supporting study personnel
2) Update as personnel, roles and/or study tasks change.

Study Tasks May include the following:

1) Obtain Informed Consent
2) Obtain Medical History
3) Physical Exams
4) Includstion/Exclusion
5) Drug Dispensing
6) Drug Accountability
7) AE/Con Med Assessment (ongoing)
8) CRF Completion
9) CRF Signature
10) Query Completion
11) Query Signature
12) Update/Maintain IRB/Feg Docs

Print Full Name	Signature	Initials	Role in Study	Study Tasks*	Duration From:	To:	Investigator Initials