
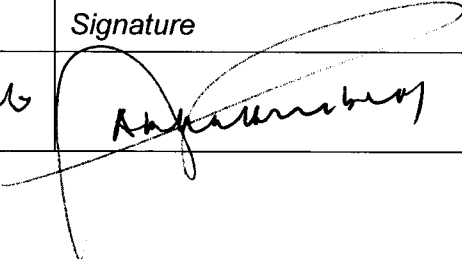


<b>Title:</b>	<b>Electronic source documents</b>	
Version:	01	
Effective date:	01-JAN-2008	
<i>Author name</i>	<i>Signature</i>	<i>Date</i>
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**Electronic source documents**

According to GCP, the investigator should provide access to source documents for the purpose of monitoring the trial. A site may use computer systems to store source documents, for example laboratory test results.

In that case the site has the following options for site visits by a monitor:

- ◆ Provide the monitor with printed copies of the required data
- ◆ Provide the monitor read-only access to the system, limited to records of patients included in the trial
- ◆ Provide the monitor with assistance from an authorized site employee who will show the required records on the screen