Title:	**Electronic source documents**	
Version:	01	
Effective date:	01-JAN-2008	
Author name	*Signature*	*Date*
P. Westveer		05-feb-2009
Approver name	*Signature*	*Date*
A. SCHATTENBERG		5-2-09

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