

Study member consent form(s)

All studies processing patient identifiable data at [National Health Service Central Register \(NHSCR\)](#) require informed consent from the people involved.

To gain adequate informed consent the National Health Service Central Register for Scotland recommends the following wording:

'I understand that information held by the NHS and records maintained by the National Health Service Central Register for Scotland may be used to keep in touch with me and follow up my health status'.

If the study subjects agree to this it provides consent for flagging or tracing for death/cancer and consent for providing Health Authority details in the future.

For studies, which do not have adequate informed consent to allow General Register Office to process identifiable patient data at NHSCR, the fact of and cause of death can be supplied.