## AuSCR Research Task Group approved projects

Title	Telephone follow-up interviewing in a national stroke registry – a process evaluation of
	respondent comments
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AuSCR role	Data provision
Approved	23 February 2014
Status	Completed
Summary	It is proposed that this article would be a reflection piece around respondent comments noted during AuSCR telephone follow-up interviews over approximately a one year period (from October 2012). Rich data are often available through the spontaneous remarks of either registrants or their proxies (e.g. carers/family members). These data could be used as part of process evaluation for this aspect of the registry operations and to inform follow-up procedures for AuSCR, and potentially other registries or research projects.
	The aim of this paper is to illustrate the strengths and limitations of telephone interviewing in a health registry for people who choose not to return mailed surveys. Research questions could include why people fail to respond to mailed-out questionnaires and what are some examples of patient/carer perceptions about the questionnaires and the registry.
Publication	Barclay-Moss K, Lannin NA, Grabsch B, Kilkenny M, Cadilhac DA <b>Stroke survivor follow-up</b> in a national registry: lessons learnt from respondents who completed telephone interviews, <i>International Journal of Stroke</i> 2019 14(2):112-114. doi:10.1177/1747493018806190