What's New in ANZDATA Collection

NON-REGISTRY TRIALS

Executive Summary

The ANZDATA Advisory Committee and Clinical Working Groups agreed to the new data collection element and this is implemented within the Online Electronic Data Collection tool (https://services.anzdata.org.au) and as a new A3 Survey and ancillary (A4) paper forms. Refer to the Registries Data Set Specification Document for more information.

Data Element

Non-Registry Trials are Ethics approved Clinical Trials that a patient is enrolled in during the survey period. For example, an Industry Drug Trial or Clinical Trial.

This data is optional.

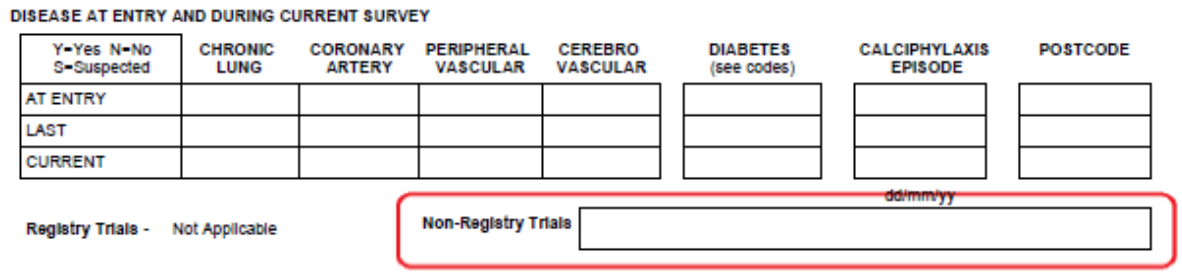
Investigating long term outcomes post clinical trial or local investigator driven trials can be facilitated by capturing this information within a Registry. Future research linking these identified trials would require appropriate Ethics and Governance approvals.

The item is stored in the ANZDATANonRegistryTrial table and is a free text field.

Collection of Data Element

This can now be reported at annual survey (via Paper) or in Real-time during a survey period online

Figure 1 – Paper Form – Non-Registry Trials



DISEASE AT ENTRY AND DURING CURRENT SURVEY

Y=Yes N=No S=Suspected	CHRONIC LUNG	CORONARY ARTERY	PERIPHERAL VASCULAR	CEREBRO VASCULAR	DIABETES (see codes)	CALCIPHYLAXIS EPISODE	POSTCODE
AT ENTRY							
LAST							
CURRENT							

dd/mm/yy

Registry Trials - Not Applicable

Non-Registry Trials

Figure 2 – Screen View – Non-Registry Trials