Participant Identification Number:	Date incident occurred (dd/mm/yyyy):

ADVERSE EVENT REPORTING FORM

Living well and enhancing active life: The IDEAL-2 study

1. Reporting

Adverse Event reported to:	By:	On: (dd/mm/yyyy)
Principal Investigator		
Chief Investigator (if serious)		
Sponsor if (serious)		
Other personnel (Please Specify)		
Adverse Event recorded electronically		

2. Adverse Event

Date researcher informed of/made aware of Adverse Event (dd/mm/yyyy)
Summary of Adverse Event

3. Any other relevant information

Please provide any additional information relevant to the Adverse Event

Report completed by:	Date

A copy of this form should be placed in the Investigators site file