A R A D

Australian Rheumatology Association Database

Reporting card for adverse events in ARAD patients receiving biologic therapy

Return this card to the ARAD data management centre in the reply-paid envelope provided.

Patient details:

Title_____ First name_____Middle name_____ Surname_____

DOB: ___ / ___ / ___ (dd / mm / yy) ☐ male ☐ female

Current bDMARD:

☐ Etanercept ☐ Infliximab ☐ Adalimumab ☐ Anakinra ☐ Rituximab ☐ Orencia

Adverse event: Date of adverse event: ____ / ____ / ____ (dd / mm / yy) (onset or diagnosis)

☐ infection ☐ malignancy ☐ other side effect → specify:_____

Severity:

☐ mild ☐ moderate ☐ severe ☐ fatal

Brief description:_____

_____

_____

_____

Outcome:

☐ recovered date of recovery ___ / ___ / ___ (dd / mm / yy) ☐ not yet recovered

Biologic therapy ☐ continued ☐ discontinued temporarily (_____weeks/months)

☐ discontinued permanently ☐ switched bDMARD to _____

Rheumatologist's details:

Title_____ First name _____ Surname_____

Signature _____ date ____/____/____ (dd / mm / yy)