



Document Number:	7.1
Document Name:	Freedom of information policy
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1.0 Purpose

The registry promotes the freedom of information. Members of the renal community have the right to access documents and information held by the registry.

2.0 Policy Details

2.1. Patients have the right to access the data they have contributed to the registry. Patients have the right to opt-out of their data being collected by the ANZDATA registry and also to withdraw their data at any point in time.

2.2. Renal units have the right to access identified data concerning their past and present patients.

2.3. With appropriate ethics approval, the renal and wider health service community have the right to access de-identified registry data relevant to a specific project.

3.0 Exhibits / Appendices / Forms

Nil

4.0 Document History

Revision	Date	Author	Description
1.0	19/05/2017	K Dansie	Creation