The Twenty Seventh Report

Australia and New Zealand Dialysis and Transplant Registry

2004



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Edited by Leonie Excell and Stephen McDonald

FUNDED BY

Commonwealth Department of Health and Ageing Kidney Health Australia New Zealand Ministry of Health

SUPPORTED BY

AMGEN Australia Pty Ltd Novartis Pharmaceuticals Australia Pty Ltd Janssen-Cilag Pty Ltd Fresenius Medical Care Australia Roche Products Pty Ltd Wyeth Australia Pty Ltd

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Editors: Leonie Excell and Stephen McDonald

Publications based upon ANZDATA Registry information reported here or supplied upon request, must include the citation as noted above and the following notice:

The data reported here have been supplied by the Australia and New Zealand Dialysis and Transplant Registry. The interpretation and reporting of these data are the responsibility of the Editors and in no way should be seen as an official policy or interpretation of the Australia and New Zealand Dialysis and Transplant Registry.

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This is the 27th Annual Report from the ANZDATA Registry. It is a comprehensive and detailed account of the delivery of dialysis and transplantation services in Australia and New Zealand in the year 2003. Once again all of the Australian and New Zealand Renal Units have contributed their patients' data to this Registry and we remain confident that there is 100% reporting.

This is the first Report for which Dr Stephen McDonald and Mrs Leonie Excell have taken sole responsibility for editing. Dr McDonald continues in his role as AMGEN Fellow in Epidemiology and has also had a major input into the composition and compilation of this Report.

Staff of the Registry, continue to provide dedicated and excellent service. Lee Excell continues in her role as Manager, Brian Livingston provides our Information Technology expertise and data analysis and Lis Steinmetz has provided administrative support. Bianca Byrne has continued in her role as Biostatistician. She has provided statistical and database analysis and has allowed us to respond rapidly to requests to the Registry from Contributors and Others. Unfortunately Bianca resigned her position during 2004 and we wish her well in her future career endeavours.

In 2004, data collection has also begun in a number of new ventures. Data has been collected for the Peritonitis Registry as well as a Living Donor Kidney Registry. Analysis of the donor data has not as yet been performed and we look forward to being able to provide information in the near future. A short report on the peritonitis is contained in this volume.

The major funding for the Registry comes from the Australian Commonwealth Department of Health and Ageing. In addition funds continue to be provided from the Kidney Health Australia and the New Zealand Ministry of Health.

We are also fortunate to receive generous "non tied" Grants from AMGEN Australia Pty Ltd, Novartis Pharmaceuticals Australia Pty Ltd, Janssen-Cilag Pty Ltd, Fresenius Medical Care Australia, Roche Products, Pty Ltd, and Wyeth Australia Pty Ltd, in 2003 and 2004.

The Internet Based Data Exchange Scheme will become completely operational by January 2005. This will enable data entry from contributing units. A change to the data collection timing for the Registry will also take place in 2005. There will be one annual data collection rather than the two as previously. In addition the Internet Based Data Exchange mechanism will be used for Real Time Data Entry of key events such as the entry of a new patient, transplantation and death.

The ANZDATA Registry Executive and ANZDATA Registry Advisory Committee are Subcommittees of the Dialysis, Nephrology and Transplant Committee of the Australia and New Zealand Society of Nephrology and Kidney Health Australia. The ANZDATA Registry Advisory Committee currently consists of:

A/Prof Rowan Walker (Chair) Prof Graeme Russ (Chair of ANZDATA Executive) Dr Stephen McDonald (AMGEN Fellow in Epidemiology) Mrs Leonie Excell (Registry Manager) A/Prof Tim Mathew (AKF Representative) A/Prof Steven Chadban (Manager/Transplantation) Professor Jeremy Chapman (Manager/Cancer) Dr Angela Webster (NOVARTIS Cancer Fellow) A/Prof Jonathan Craig (Manager/Paediatrics) Dr Mark Marshall(Manager/Haemodialysis) A/Prof David Johnson (Manager/CAPD) Dr Ian Dittmer (NZ Representative) Dr John Agar Dr Frank Ierino Dr Grant Luxton Ms Denise Tomlinson (Nursing Representative) Ms Mardi Thompson (Client Representative)

In addition small Working Groups continue to work in each of the specialty areas and have been responsible for analysis of data in their specialty as well as the production of manuscripts for publication in this report and in the international literature.

Graeme Russ Chair ANZDATA Executive

EDITORS' COMMENTS

This report sees the continuation of the evolutionary changes in the report of the last few years. The appendices continue to grow, with an increasing amount of tabular material which we hope will be of interest. There are no extra "chapters" this year; rather the new material has been incorporated into the individual chapters. This includes examination of the trends in mortality over time, use of erythropoietic agents, calcium and phosphate data, trends in BMI, and material from the peritonitis registry. Important changes have also occurred in the cancer chapter, with substantially altered methodology and reference ranges.

PRIVACY

In December 2001 changes to the Commonwealth Privacy Act were introduced which have led to changes to the collection of personal information. Essentially these extend to the private sector a number of changes based around 10 "National Privacy Principles" (NPP's). A detailed exposition of these can be found at the Privacy Commissioner's website (www.privacy.gov.au). Briefly, however, health information is treated as "sensitive" information, which must usually be collected and handled with consent of the person, unless certain conditions are met.

Each Australian State has also enacted similar provisions which cover practice and patients in public hospitals.

COLLECTION OF DATA

ANZDATA spent some time during 2002 formulating an appropriate response to these issues including seeking advice from a variety of sources. The approach taken has been that of a "opt-out" consent, whereby patients are distributed information outlining the nature and purpose of the information collected, offered an opportunity to view that data and ask questions, and the opportunity to request withdrawal of part or all of their data. This approach is explicitly suggested for Registries by the Privacy Commissioner in his "Guidelines for the Health Sector". To this end ANZDATA has circulated to all participating hospitals a patient information sheet, for each hospital to use (or a locally modified version if appropriate) to inform patients.

At the time of data collection each unit is asked to certify that they have complied with measures under the relevant privacy act.

USE OF DATA

ANZDATA does not release data identifiable by patient name. Results are published/released in tabular or graphic format. On occasion, when data identifying particular hospitals is involved, consent from the Director of the relevent renal unit is sought prior to the release of information.



ANZDATA REGISTRY

AUSTRALIA AND NEW ZEALAND DIALYSIS AND TRANSPLANT REGISTRY

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Important Privacy Information

As part of routine medical care of people receiving treatment with dialysis or kidney transplantation, your kidney specialist collects certain information about the patients they treat. All kidney specialists throughout Australia and New Zealand report this information every 6 months to the Australia and New Zealand Dialysis and Transplant Registry (ANZDATA). ANZDATA collects the information for the purpose of monitoring treatments and performing analyses to improve quality of care for people with kidney failure.

1. What is ANZDATA?

ANZDATA is an organization set up by the Australian Kidney Foundation and the Australia and New Zealand Society of Nephrology to monitor dialysis and transplant treatments. ANZDATA is funded by the Australian and New Zealand Governments and the Australian Kidney Foundation.

2. What information is collected about you ?

This information includes your name, age, gender, racial origin, hospital of treatment, some aspects of your medical condition (such as whether you have diabetes) and details about the type of kidney treatment you are receiving (dialysis or transplant).

We **DO NOT** collect details about your address, telephone number, medical insurance, or non-medical matters such as occupation, income, etc

3. Is personal data ever released ?

The identity of people in the database **IS NOT released publicly nor in any reports**. Measures have been put into place to ensure the security of all collected information.

4. What is this information used for ?

The information is used primarily for quality assurance, investigating patterns of kidney disease, and planning appropriate health services. We release reports on a variety of topics, including an Annual Report examining the rates and treatment of kidney failure in Australia and New Zealand. We also have a major role in ensuring the quality of patient care by sending to each kidney unit each year a report outlining their activity. These reports also compare the outcome of the treatment they provide with that of other units throughout the two countries. Reports are also produced at a state and national level, and from time to time analyses are also produced for renal units, government health departments and industry concentrating on particular aspects of renal failure management eg peritoneal dialysis, transplantation, haemodialysis.

5. Can you see what personal information ANZDATA collects and the reports that it produces ?

Individuals are able to view their own information on request. You can request alterations if you believe it is inaccurate. You may also opt not to have your treatment included in this database, and you should let your kidney specialist know if this is the case. You can also choose not to have some information (e.g. racial origin) recorded. However, if your information is not included in the Registry, the ability to compare results in Australia and New Zealand or to analyse the results of different treatment methods and for different patient types (eg diabetics) will be compromised.

The national reports and much other material produced by ANZDATA are available free on the Internet at www.anzdata.org.au, or they can be sent to you on request to the address above. Your kidney specialist will also have copies of many of the reports.

If you wish to discuss any of the issues raised here, please let your doctor know or telephone the ANZDATA Registry direct on 08 8222 6704. You may also write to us (ANZDATA Registry, C/- The Queen Elizabeth Hospital, 28 Woodville Road SA 5011) or send us an e-mail (anzdata@anzdata.org.au).

GUIDELINES FOR DATA RELEASE

The policy for release of data to investigators, renal units and others was revised during 2002 and is summarised on the Website. ANZDATA encourages the analysis, use and citation of its data, and receives many data requests annually which vary in size and complexity. At times these overwhelm the limited resources within the Registry, and must be prioritised. Generally, formal requests for data are preceded by a period of consultation with a member of the Registry staff. Requests are welcome from Renal Physicians, other staff members of Renal Units, Charitable Bodies, Academic Institutions, Government Departments and Industry. Requests dealing with identifiable Hospital data will only be fulfilled with the explicit consent of the relevant Hospital Units.

ATTRIBUTION OF PUBLICATIONS

The policy on attribution of publications which incorporate ANZDATA sourced data was revised during 2002, following a period of consultation with participating physicians.

Where a member of a participating unit has analysed data provided by ANZDATA and subsequently prepared a manuscript, then "ANZDATA Registry" should be acknowledged as a secondary institution in addition to the author's Hospital or University. This applies whether the primary data analysis is performed by the author or by ANZDATA staff. Where the author is an ANZDATA office holder or staff member then the primary attribution should be "ANZDATA Registry".

Where ANZDATA data is only a minor portion of the work, then it may be more appropriate to acknowledge the source explicitly in the "Acknowledgements" section.

In both cases the disclaimer on page ii of this report should be included.

In all cases the source and treatment of the data should be made clear in the "Methods" section. Preferably the abstract (and keywords if applicable) should also include "ANZDATA" which would allow for searching Registry publications. ANZDATA Registry 2004 Report

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1. Wording

Throughout this report 'treatment' refers to renal replacement therapy, including haemodialysis, peritoneal dialysis and transplantation

HD	= haemodialysis	CAPD = continuous ambulatory peritoneal dialysis
APD	= automated peritoneal dialysis	ESRD = end stage renal disease

2. Data collection

ANZDATA collects information from all renal units in Australia and New Zealand every 6 months. Currently this is by a paper-based system, with manual completion of the form and manual data entry. No formal audit mechanism is in place at this stage.

For transplants, HLA matching and panel reactive antibodies are obtained direct from the Tissue Typing laboratories in each State.

3. Inclusion criteria

Included in the Registry are all patients receiving renal replacement therapy where the intention to treat is longterm, i.e. medical opinion is that renal function will not recover. Cases of acute renal failure are excluded. People who move overseas permanently are censored at date of last treatment (or departure in the case of transplant recipients).

4. Modality attribution

The initial mode of dialysis is determined at 90 days after first treatment, to allow for early changes and maturation of access. Other transfers (between modalities, or from satellite to hospital HD etc.) are not analysed if less than 30 days, except for transfers between dialysis centres to which a 60 day rule is applied to allow for holiday movements.

5. Underlying renal disease

This is recorded by the treating hospital according to a modified EDTA coding system (details on back of survey form).

6. Deaths

Deaths are coded by the treating hospital using a modified EDTA coding system (details on back of survey form). Where a recent change in location or modality preceded death by <30 days, the death is attributed to the modality / location at time of death, although some analyses differ and this is stated for those analyses. All deaths following transplant surgery are attributed to transplantation.

7. Co-morbid conditions

These are recorded by the treating hospital. No definitions are supplied; the treating clinician is asked to record whether the patient has coronary artery disease, chronic lung disease, cerebrovascular disease, peripheral vascular disease or diabetes according to their clinical opinion on a yes / suspected / no basis.

8. Transplant Waiting List

The active transplant waiting list includes listing at any stage during the survey period.

9. Derived measures

9.1 Haemoglobin

Haemoglobin is recorded as the last available measurement before the end of the survey period.

9.2 Erythropoietic agents

Erythropoietin agent use is recorded as "yes" if these agents were used at any time during the survey period.

9.3 Iron Studies

Iron studies are requested within the last three months of the survey period.

9.4 Estimated creatinine clearance

Where creatinine clearance is estimated from serum creatinine at entry or post transplantation, the Cockroft-Gault equation is used [1].

Cl_{cr}=(140-age)*weight/(814*Cr_{serum})[*0.85 if female]

The weight term used for this is lean body mass, calculated using the equation LBW=(0.9*[height-152])+(50 if male, 45.5 if female) [2].

9.5 Urea reduction ratio / Kt/V

Results are requested in one of these formats, using the stop flow method on a mid-week dialysis. Single pool Kt/V is collected, along with the method used.

For conversion of URR to Kt/V urea the formula used [3] is

Kt/V = 0.023*PRU - 0.284 (note that PRU = percent reduction in urea and not URR).



9.6 **Body Mass Index**

Body mass index (BMI) is calculated as weight (kg)

 $(height(m))^2$

	The standard NH&MRC categories are used:	underweight	$< 20 \text{ kg/m}^2$	normal	$20-24.9 \text{kg/m}^2$
		overweight	25-29.9 kg/m ²	obese	$> 30 \text{kg/m}^2$
9.7	Peritoneal Dialysis measures	-	-		-
	These are the standard measures, often calculate	ed by compute	rised patient mana	gement progra	ıms.
9.7.1	Residual renal function				

Residual renal function

The measure used is the arithmetic mean of urea and creatinine clearance from a 24-hour urine collection and serum creatinine & urea.

9.7.2 **Peritoneal equilibration test**

The ratio of dialysate to plasma glucose is used, following a 4 hour dwell of a 2 litre 2.5% bag of dialysate, performed within 6 months after initiation of PD.

10. **Rates & Measures**

10.1 **Incidence** rates

Except where otherwise stated, quoted incidence rates are per calendar year, and are expressed per million population.

10.2 **Prevalence** rates

Except where otherwise specified, prevalence rates are point prevalence rates at 31 December 2002.

10.3 **Population denominator**

The population estimates used are the estimated resident populations (ERP) for the year 2002, released by the Australian Bureau of Statistics and Statistics New Zealand. Figures used are those for the June quarter.

For both countries, the statistics bureaux record indigenous status on a self-identification basis. For Australia, there has been considerable change in the propensity to self-identify as indigenous, such that a number of estimates are released by the ABS [4]. For this report, the low range projections have been used.

10.4 Survival rates

For transplant recipients, survival rates exclude those who were transplanted overseas or were recipients of multiple organ grafts.

Graft survival (unless otherwise qualified) includes both cessation of graft function (i.e. return to dialysis) and patient death.

Patient survival for transplant recipients - rates for fixed periods are calculated according to the life-table method and include an adjustment to the risk-set of ¹/₂ of those censored without failure over the interval to create an "average" risk set.

Patient and technique survivals for Haemodialysis and Peritoneal Dialysis are based on the dialysis modality at 90 days after first treatment for patients not grafted during that period. Patients are followed up until they are either grafted (at which point they are censored) or until they have a 'permanent' change of dialysis modality or until death or most recent follow up date. A 'permanent'change of dialysis is defined as any change in excess of 30 days.

10.5 Death and other event rates

Rates are expressed per 100 person years at risk (unless otherwise stated).

Come analyses include survival of all patients, others exclude the first 90 days of followup. This is stated in the individual analyses.

10.6 Age standardisation

All rates are crude, not age-standardised. The age distribution of the populations for Australia and New Zealand are given in Appendix I.

11. Database

Data is stored on a relational database using ORACLE version 8I.

12. Statistics

Statistical analyses were performed using SPSS release version 10.0.7 and Stata version 8.2.

13. References

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ANZ

xiv

20 - TYPE OF DIALYSIS

5 - RACIAL ORIGIN Caucasold Australian Aborigine Chinese

PRIMARY RENAL DISEASE contu-cor Peter barum reprintinguti 201 Peaterum reprintinguti 201 Peaterus rentrai values 202 Peaterusted megaureter 203 Osthursted megaureter 203 Nethorized failed bladder and ureters 203 Nethologic bladde

- Cook Islander
 Samoun
 Tongan
 Tongan
 Pacific Islander other (specify) 68 68 61
 - sian
- Vietnamese Other (specify) Patient objects to answering question °°₽∓88
- Mixed race coded by patient's asse

6 - PRIMARY RENAL DISEASE

- CARDIAC Results of ANCA (Anti Neutrophil Cytoplasmic Antibody) test in association with glomerulonephritis should be entered in box marked OTHER
 - 100 Presumed GN, type undefined histologically (no biopsy) 110 Feast actionsing Strin (incucking) syntholaisi 13 Messangiocapillary GN with subendorhelai 132 Messangiocapillary GN with intramembranous obpatis (dome orchoru) 20 Messangiocapillary GN with intramembranous obpatis (dome octorol disease)

- Myocardal ischamia (presumed)
 Myocardal ischamia and infarction
 Myoradia ischamia and infarction
 Myoradaamia
 Hemorinagio parte
 Hemorinagio parte
 Cause uncertain
 Cardiac anneel cause of cardiac failure
 Chino cause of cardiac failure
 Thin cause of cardiac failure
 - VASCULAR

130 Membranous GN 140 Extra and intra capillary GN (extensive

- Palmonary embolus
 Controvensular aconomication
 Contrometarian hiermortinage
 Haemonimage from transplant aconos site
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 Heemonimage from elsewhere (apacity)
 Bowel Hetrolion

censents: - pricinary major programs.two 151 Massnapils profilerative (g/A+ prosinos) 152 Massnapils profilerative (g/A+ prosinos) 153 Massnapils (g/A) massnapils 154 Massnapils (g/A) massnapils 155 Massnapils (g/A) massnapils 156 Massnapils (g/A) massnapils 156 Massnapils (g/A) massnapils 157 Massnapils

- INFECTION
- Please enter code for nature of infective organism, after the code for site of infection Please specify type of organism ag Staph, CMV, Candida, etc
 - - eg 321 Lung infection bacterial (staph) 322 Lung infection viral (CMV)

Complete the requested details reparding, date, identity of drug. humber of doses given, and reason for administration, acording to the following codes Record in order of administration, each separate course of such drugs; a second course of the same drug should be separately

NUMBER OF

DOSES

Daclizumab (Zenepax)

OKT3

Spontaneous fall in se.creatinine by 10% within 24 hours

52 - IMMEDIATE FUNCTION

Spontaneous fail in secreatinine by 10%, first recorded between 25-72 hours

~ _

34 - PET TEST (Required Once Only per patient)

Standard Peritoneal Dialysis Equilibration Test performed 1-6 months after initiation of PD (2.5% 2 litre exchanges)

Provide dialysis/plasma creatinine at 4 hours Range 0.1 - 1.2

TYPE OF AGENT

Record actual number of doses given

Intravenous Immunoglobulin Basilixmab (Simulect)

Other monoclonal (specify)

Poor immediate function. No spontaneous fall in se.creatinine within 72 hours: but no dialysis needed No immediate function. No spontaneous fall (> 10%) in secreatinine; dialysis required within 72 hours

.....

Rituximab Polyclonal anti T cell REASON FOR USE

56 - MONOCLONAL / POLYCLONAL

THERAPY

51 - TOTAL ISCHAEMIA (HOURS)

Post dialysis.urea: Blood is again drawn from the 'artenial' needle and this should occur within 20 seconds after cessation of the blood pump (alternatively the pump can be turned down to 50 millimit) – this is to avoid

problems with recirculation

Pre dialysis urea: Blood should be drawn from the 'arterial' needle immediately prior to dialysis, at a mid-week dialysis session

(Pre. dialysis.urea - post dialysis.urea,) × 100 = URR%. Pre dialysis urea

UREA REDUCTION RATIO %

Kt/V (for HD patients) Range 0.5 - 2.2 A Urea Reduction Ratio % (URR) B KIV (by BIOSTAT) C KIV (by UKM) D KIV (by DAUGIRDAS – single pool) E KIV (other method – specify)

33 - TYPE OF ACCESS USED IF INITIAL Treatment was haemodialysis

Leave blank if initial renal replacement treatment was not

alysis.

From time of donor renail artery interruption or aortic clamp, until time of release of renail artery in the recipient (clamp off)

00 Other (specify) 01 Donor malignancy 02 Malignancy invading graft

MISCELLANEOUS

6 Monozygotic (identical) twin 7 Dizygotic (non-identical) twin 8 Other related living donor (specify)

10 Daughter 11 Husband 12 Wife 33 Cousin 14 Unrelated Rving donor (specify)

DRUG THERAPY 90 Complexations of taugy interpay requiring reduction or withdrawal of staroid androir immunosuppressants 1 Non-compliance with therapy – causing graft taillance 35 Rejection biolowary 15 reduction due to imferioren 53 Rejection biolowary 15 reduction due to inferioren

- ANS
 Bacterial

 22 Unitary track
 2 Viral

 23 Unitary track
 3 Viral

 24 Wound
 4 Protocoa

 25 Wound
 5 Protocoa

 26 Performance
 6 Protocoa

 27 Wound
 5 Protocoa

 28 Performance
 4 Protocoa

 29 Performance
 6 Protocoa

 21 Differ
 5 Performance

 22 Differ
 5 Performance

 23 Differ
 5 Performance

 24 Protocoa
 5 Performance
- SOCIAL and hencentration
 Polleration Construction of the constructi

- Withdravel for psycho-social reasons
 Fateri related further treatment (specify reason)
 Tanner related further treatment (specify reason)
 Tanner posses for any other reason (specify reason)
 Kondental death (specify)
 Withdrawel for cereborascular common and conditions
 Withdrawel for cereborascular common and conditions
 Withdrawel incertained valuation common and conditions
 Withdrawel indext to cereborascular common and conditions
 Withdrawel indext to cally sea socies of fifticulies (WF, Fockhof, etc)
 - - - - MISCELLANEOUS

Enter the total daily dose for each drug where applicable; if an unlisted drug is used, enter the name in the space provided marked OTHER

57 - TOTAL DAILY DRUG DOSE

Prophylaxis
 Treatment for acute rejection
 Other (specify)

53 – DISEASE IN GRAFT Histologically prover

Complete this section for FUNCTIONING or FAILED GRAFTS

39 DIALYSATE CREATININE CLEARANCE (per week)

Total (includes a 24 hour urine and dialysate

Range 10 - 200 litres/week Litres/week /1.73m² Body Surface Area

DIALYSATE WEEKLY KIV

6

39 to 41 - PD CLEARANCE STUDIES

Generated from a 24 hour collection of PD effluent and urine

Y = Disease rocurrence - primary renal disease and disease in graft the same

 D = De novo glomerulonephritis
 – primary renel disease known and not the same G = Giomenulonephritis in graft - primary renal disease unknown or not biopsled

The initial drug dose (at zero months) is <u>the **first oral maintenance dose**: do **NOT** enter the intravenous loading doses administered at or shortly after transplantation</u>

(31-Mar-04)

In cases of giomenulonephritis, where histological confirmation of recurrence may be uncertain, enter as G

Only those drugs taken at the listed intervals should be entered; where necessary provide the dose recorded on the closest day preceding the requested time interval

Congenital renal hypoplasia and dysplasia Loss of single kidney (specify - e.g. trauma, surgery)

XV

- Hepatic failure (specify)
 Unamate custod by graf failure
 Bancrostifis
 Bancrostifis
 Bancrostifis
 Bancrostifis
 Bancrostifis
 Bancrostific
 Bancrostific
- (per week) = mean of urea and creatinine clearance Litres/week/1.73m² Body Surface Area RESIDUAL CREATININE CLEARANCE Range 0.1-5.0 41

- 11 Haemodiarysis plate dialysers 21 Haemodiaysis nollow face dialysers 13 Haemodiarsion 14 Haemodiarsion 21 Perintonae bagis no cycler 22 Perintonae contrause antialogy (CAPD) 22 Perintonae automated (APD) 22 Perintonae automated (APD) 23 Perintonae antimated (APD)
- 42 REASON FOR TRANSFER
- 5 • BETWEEN CAPD and APD

55 - CAUSE OF GRAFT FAILURE

Anz b

- * TRANSFER FROM CAPD / APD to HD

- 10 Hyperacule rejection (within 48 hours of transpantation) A charte investion: analysism, sustained and a superfix function 40 Chronic adoptint nephropathy (slow prograssive base mail investion, not due to rocurrent original disease or auto inpedient. REJECTION

ANZDATA Registry 2004 Report

Remain Artery stencests
 Remain a fatery transcession
 Remain a fatery transcession
 Remain vessel haemonthrage (primary)
 Remain vessel haemonthrage (secondary)
 Enclosus – transcession
 Enclosus – transcession
 Removies and severation

Reurnerk / pensistent peritonitis
 Tume/ point / pensistent peritonitis
 Tume/ point / pensistent peritonitis
 Tume/ point / pensistent /

22 - UNCORRECTED CALCIUM

At end of survey, transplantation or death.

21 - DRY WEIGHT

Composed the media media media (media) (media) and media media media media media media media media media (media) (medi

Midwoek, predialysis and closest to end of Survey Midweek, predialysis and closest to end of Survey Midweek, prediatysis and closest to end of Survey

17 - CAUSE OF DEATH

23 - PHOSPHATE Not corrected for albumin

Non-viable kidney (due to pre-transplant contrical necros
 Contical necrosis post transplant (not due to rejection)
 Unstanc and bladder problems

TECHNICAL

Mesangiocapillary GN with subendothelial deposits Mesangiocapillary GN with intramembranous depos

82 Mesang 83 Mesang

GLOMERULONEPHRITIS

memory and the second se

50 - SOURCE OF DONOR KIDNEY

method used

32 - URR or Kt/V Please enter I

24 - HAEMOGLOBIN

Cadaver
 Cadaver
 Kwin, record 6 or 7)
 Brother (if twin, record 6 or 7)
 Mother

ANZDATA Registry 2004 Report



QUEENSLAND Allamanda Private Hospital **Bundaberg Base Hospital** Cairns Base Hospital Caloundra Private Hospital Goldcoast Hospital Greenslopes Private Hospital (Baxter) Hervey Bay Hospital John Flynn Hospital Mackay Base Hospital Nambour Hospital Pine Rivers Private Hospital Princess Alexandra Hospital **Rockhampton Base Hospital Royal Brisbane Hospital** St. Andrew's Private Hospital (Gambro) Toowoomba Hospital Townsville General Hospital Wesley Private Hospital **New South Wales** Children's Hospital at Westmead **Dubbo Base Hospital** East Coast Renal Service Prince of Wales Hospital Sydney Children's Hospital St. George Hospital St. Vincent's Hospital Wollongong Hospital Gosford Hospital John Hunter Hospital Lismore Hospital Mater Misericordiae Hospital Port Macquarie Community Dialysis Centre Port Macquarie Private Hospital **Royal North Shore Hospital** South West Sydney Renal Service Bankstown Hospital Liverpool Hospital Statewide Renal Services Concord Hospital Royal Prince Alfred Hospital Sydney Adventist Hospital Tamworth Hospital Western Renal Network Westmead Hospital Orange Base Hospital Wentworth Hospital **AUSTRALIAN CAPITAL TERRITORY (ACT)** The Canberra Hospital

VICTORIA

Alfred Hospital Austin and Repatriation Medical Centre **Epworth Hospital** Forest Hill Dialysis Centre (Nephrocare) **Geelong Hospital** Kew Private Dialysis Centre (Baxter) Monash Medical Centre - Adult Monash Medical Centre - Paediatric Royal Children's Hospital North West Dialysis Service Royal Melbourne Hospital St. Vincent's Hospital TASMANIA Launceston General Hospital **Royal Hobart Hospital** SOUTH AUSTRALIA Flinders Medical Centre The Queen Elizabeth Hospital **Royal Adelaide Hospital** Women's and Children's Hospital **NORTHERN TERRITORY Royal Darwin Hospital** Alice Springs Hospital WESTERN AUSTRALIA Fremantle Hospital Hollywood Private Hospital Princess Margaret Hospital for Children **Royal Perth Hospital** Sir Charles Gairdner Hospital St. John of God Private Hospital

New Zealand

Auckland CityHospital Starship Children's Hospital Christchurch Hospital Dunedin Hospital Middlemore Hospital Palmerston North Hospital Taranaki Base Hospital Waikato Hospital Wellington Hospital Whangarei Area Hospital

QUEENSLAND

Princess Alexandra Hospital (Adult & Paediatric) Director of Transplantation - Dr David Nicol Ipswich Road Woolloongabba 4102

New South Wales

John Hunter Hospital Director of Transplantation - Professor Adrian Hibberd Lookout Road New Lambton Heights Newcastle 2304

Prince of Wales Hospital (Adult & Paediatric) Director - Professor John Charlesworth Barker Street Randwick 2031

Royal North Shore Hospital Director - Dr Lloyd Ibels Pacific Highway St Leonards 2065

Royal Prince Alfred Hospital Director - Associate Professor Josette Eris Missenden Road Camperdown 2050

St George Hospital Director of Transplantation - Professor John Kelly Montgomery Street Kogarah 2217

St. Vincent's Hospital Director - Dr Tim Furlong Victoria Street Darlinghurst 2010

The New Children's Hospital at Westmead Director - Dr Elisabeth Hodson Cnr Hawkesbury and Hainsworth Street Westmead 2145

Westmead Hospital Director - Professor Jeremy Chapman Cnr Hawkesbury and Darcy Road Westmead 2145

VICTORIA

Alfred Hospital Director - Professor Napier Thomson Commercial Road Prahran 3181

Austin & Repatriation Medical Centre Director - Dr David Power Burgundy Road Heidelberg 3084

Monash Medical Centre Paediatric Director - Dr Amanda Walker 246 Clayton Road Clayton 3165

Monash Medical Centre Adult Director - Professor Robert Atkins 246 Clayton Road Clayton 3165

VICTORIA cont

Royal Children's Hospital Director - Dr Colin Jones Flemington Road Parkville 3052

Royal Melbourne Hospital Director - Professor Gavin Becker Parkville 3052

St. Vincent's Hospital Director - Dr Brendan Murphy 41 Victoria Parade Fitzroy 3065

South Australia

The Queen Elizabeth Hospital Director - Professor Graeme Russ 28 Woodville Road Woodville 5011

Women's and Children's Hospital Director - Dr Paul Henning 72 King William Road North Adelaide 5006

WESTERN AUSTRALIA

Princess Margaret Hospital for Children Director - Dr Ian Hewitt Roberts Road Subiaco 6008

Royal Perth Hospital Director - Dr Ashley Irish Wellington Street Perth 6001

Sir Charles Gairdner Hospital Director - Dr Brian Hutchison Verdun Street Nedlands 6009

New Zealand

Auckland City Hospital Director - Dr John Collins Park Road Grafton, Auckland

Christchurch Hospital Director - Dr Richard Robson Riccarton Avenue Christchurch

Starship Children's Hospital Director - Dr William Wong Park Road Grafton, Auckland

Wellington Hospital Director - Dr Grant Pidgeon Riddiford Street Newtown, Wellington South QUEENSLAND

ANZ DATA

Atherton Satellite - Cairns Base Hospital Cairns Private Hospital Satellite - Cairns Base Hospital Home Hill Satellite - Townsville General Hospital Innisfail Hospital - Cairns Base Hospital Ipswich Satellite - Princess Alexandra Hospital Logan Satellite - Princess Alexandra Hospital Mt Isa Satellite - Townsville General Hospital Noosa Satellite - Nambour Hospital Palm Island Satellite - Townsville General Hospital Redcliffe Satellite - Royal Brisbane Hospital Robina Satellite - Goldcoast Hospital Vincent Satellite - Townsville General Hospital NEW SOUTH WALES Ballina Satellite - Lismore Hospital Bankstown Hospital - South West Sydney Renal Service Bathurst Hospital - Orange Hospital Blacktown Satellite - Westmead Hospital Brewarrina Hospital Campbelltown Satellite - South West Sydney Renal Serviced Coffs Harbour Base Hospital Coonamble Hospital Dame Eadith Walker - Statewide Renal Services Dubbo Base Hospital Eora Cottage - Prince of Wales Hospital Grafton Hospital - Lismore Hospital Lakehaven Satellite - Gosford Hospital Lanceley Cottage - Royal North Shore Hospital Lindfield Private Dialysis (Gambro) Liverpool Community Centre - South West Sydney Renal Service Maitland Hospital - John Hunter Hospital Muswellbrook - John Hunter Hospital Nita Reed House (Taree) - John Hunter Hospital Norfolk Island Hospital - Statewide Renal Services Orange Base Hospital - Westmead Hospital Port Macquarie Community Dialysis Centre Port Macquarie Private Hospital Shellharbour - Wollongong Hospital Shellharbour - Wollongong Hospital Singleton Satellite - John Hunter Hospital Sydney Adventist Hospital Sydney Dialysis Centre Wagga Wagga Base Hospital Wansey Satellite - John Hunter Hospital Wentworth Satellite - Westmead Hospital AUSTRALIAN CAPITAL TERRITORY (ACT) Canberra Community Dialysis Centre VICTORIA Angliss Hospital Ararat Hospital Austin Training Satellite - Austin & Repatriation Hospital Bacchus Marsh Hospital Bairnsdale Hospital Ballarat Health Services

Bendigo Hospital Berwick Hospital Broadmeadows Hospital Brunswick Satellite Casterton Hospital Caulfield Satellite Coburg Satellite Cohuna Hospital Colac Hospital Corryong Satellite Cranbourne Satellite Dandenong Satellite Daylesford Hospital Echuca Hospital Edenhope Hospital Epping Dialysis Unit Epworth Hospital Forest Hill Private Dialysis (Nephrocare) Frankston Satellite Gambro - Diamond Valley Hospital Geelong Hospital Goulburn Valley Hospital Hamilton Hospital Hastings Hospital Heidelberg - Austin & Repatriation Hospital Horsham Satellite Kew Private Dialysis Centre (Baxter) Kyneton Hospital

La Trobe Regional Satellite

ANZDATA Registry 2004 Report

VICTORIA CONT Lorne Hospital

Maryborough District Health Service Mildura Hospital Mitcham Hospital Moorabbin Satellite Myrtleford Hospital Nauru (overseas) - Alfred Hospital Nauru (overseas) - Monash Medical Centre Adult Newcomb Satellite North East Kidney Service - Austin & Repatriation Hospital Northern Hospital Satellite Omeo District Hospital Orbost Hospital Peter James Centre Portland Hospital Robinvale Hospital Rosebud Hospital Sale Hospital Sandringham Satellite Seymour Hospital St. Arnaud Hospital St. George's Hospital Sunshine Satellite Swan Hill Hospital Terang Satellite Timboon Hospital Wangaratta Hospital Warnnambool Hospital Werribee Mercy Hospital Western Gippsland Hospital Williamstown Satellite Wodonga Hospital Wonthaggi Hospital Yarawonga District Hospital Yarram Hospital TASMANIA North West Renal Unit, Burnie - Launceston Hospital SOUTH AUSTRALIA Berri Hospital Ceduna Satellite Hartley Private Hospital (Fresenius) Lyell McEwin Satellite Modbury Private Dialysis (Fresenius) Murray Bridge Hospital Noarlunga Satellite Centre Payneham Private Dialysis (Baxter) Port Augusta Hospital Port Lincoln Satellite Centre Wayville Satellite Centre NORTHERN TERRITORY Bathurst Island Hospital - Royal Darwin Hospital Community Health Centre - Alice Springs Hospital Katherine Dialysis Unit - Royal Darwin Hospital Nightcliff Community Centre - Royal Darwin Hospital Tennant Creek Hospital - Alice Springs Hospital WESTERN AUSTRALIA Albany Satellite Armadale Satellite Bunbury Satellite Geraldton Hospital Joondalup Satellite Unit Kalgoorlie Dialysis Unit Kimberley Dialysis Centre - Royal Perth Hospital Melville Satellite Midland Private Satellite Dialysis Centre (Baxter) Peel Health Campus - Mandurah Pilbara Dialysis Unit [Port Hedland] - Royal Perth Hospital Royal Perth Rehabilitation Hospital - Royal Perth Hospital

NEW ZEALAND

Bay of Islands Hospital - Whangarei Hospital Carrington Satellite - Auckland City Hospital Greenlane Hospital - Auckland City Hospital Manukau Satellite - Middlemore Hospital Middlemore Hospital Porirui Satellite - Wellington Hospital

During the calendar year 2003 (the period covered by this Report), the following manuscripts based on ANZDATA material appeared in peer-reviewed journals.

Polkinghorne KR, McDonald SP, Marshall MR, Atkins RC, Kerr PG. Vascular access practice patterns in the New Zealand haemodialysis population. *Am J Kid Dis. 2003; 43:696-704.*

Polkinghorne KR, McDonald SP, Atkins RC, Kerr PG. Epidemiology of vascular access in the Australian haemodialysis population. *Kidney Int. 2003; 64:1893-1902.*

Cass A, Cunningham J, Snelling P, Wang Z, Hoy W. Urban disadvantage and delayed nephrology referral in Australia. *Health Place. 2003; 9:175-182.*

Cass A, Cunningham J, Snelling P, Ayanian JZ. Late referral to a nephrologist reduces access to renal transplantation. *Am J Kidney Dis. 2003; 42:1043-1049.*

Chapman J, Russ G. Geographic variance in access to renal transplantation in Australia. *Transplantation 2003; 76:1403-6*.

Li SQ, Cass A, Cunningham J. Cause of death in patients with end-stage renal disease: assessing concordance of death certificates with registry reports. *Aust N Z J Public Health. 2003; 27:419-424.*

McDonald SP, Collins J, Johnson DW. Obesity is associated with worse peritoneal dialysis outcomes in the Australian and New Zealand patient populations. *J Am Soc Nephrol. 2003; 14:2894-2901.*

McDonald SP, Russ GR. Current incidence, treatment patterns and outcome of end-stage renal disease among indigenous groups in Australia and New Zealand. *Nephrology. 2003; 8:42-48. Kidney Int. 2003; 63:s123-s127.*

Stewart JH, Buccianti G, Agodoa L, et al.

Cancers of the kidney and urinary tract in patients on dialysis for end-stage renal disease: analysis of data from the United States, Europe, and Australia and New Zealand. *J Am Soc Nephrol 2003; 14:197-207.*

A number of further manuscripts have been published in 2004 and are listed below.

Lim WH. Is there a role for peritoneal dialysis in remote aboriginal patients with end-stage renal disease in Australia? *Nephrology (Carlton) 2004; 9:S126-8.*

McDonald S, Collins J, Rumpsfeld M, Johnson D. Obesity is a risk factor for peritonitis in the Australian and New Zealand peritoneal dialysis patient populations. *Perit Dial Int 2004; 24:340-346.*

McDonald S. Indigenous transplant outcomes in Australia: What the ANZDATA Registry tells us. *Nephrology (Carlton) 2004; 9 Suppl 4:S138-43.*

McDonald SP, Craig JC. Long term survival of children with end-stage renal disease. *N Engl J Med 2004; 350:2654-2662.*

McDonald SP, Marshall MR, Kerr PG, Russ GR. Erythropoietic agents, iron and hemoglobin - What happens beyond the trial setting: Observational data from the ANZDATA Registry. *Hemodialysis Int 2004; 8:257-264.*

Polkinghorne KR, McDonald SP, Atkins RC, Kerr PG. Vascular access and all-cause mortality: a propensity score analysis. J Am Soc Nephrol 2004; 15:477-86.

Rumpsfeld M, McDonald SP, Purdie DM, Collins J, Johnson DW. Predictors of baseline peritoneal transport status in Australian and New Zealand peritoneal dialysis patients. *Am J Kidney Dis 2004; 43:492-501.*

Stewart JH, McCredie MR, McDonald SP. Incidence of end-stage renal disease in overseas-born, compared with Australian-born, non-indigenous Australians. *Nephrology 2004; 9:247-252.*

Stewart JH, McCredie MR, Williams SM, McDonald SP. Interpreting incidence trends for treated end-stage renal disease: Implications for evaluating disease control in Australia. *Nephrology 2004; 9:238-246*.

Stewart JH, McCredie MRE, McDonald SP. The incidence of treated end-stage renal disease in New Zealand Maori and Pacific Island people and in Indigenous Australians. *Nephrol Dial Transplant 2004; 19:678-85.*



SUMMARY

KEY SUMMARY POINTS FROM THE REPORT

AUSTRALIA

- There were 13,625 patients (685 per million) receiving renal replacement therapy (RRT) at 31 December 2003. Of these, 5,951 (299 per million) had a functioning kidney transplant and 7,674 (386 per million) received dialysis treatment.
- 1,953 patients commenced RRT in Australia in 2003 (98 per million). The intake varied from 262 per million population in the Northern Territory to 75 per million in Tasmania.
- The mean age at commencement was 59.3 years.
- 27% of new patients had glomerulonephritis attributed as their cause of end stage renal failure, 26% diabetic nephropathy, and 15% hypertension.
- Of patients <65 years of age and receiving dialysis treatment, 35% were on the active kidney transplantation waiting list. This proportion varied between 14% in the Northern Territory and 50% in the Australian Capital Territory. Only 7% of Aboriginal/Torres Strait Islander patients <65 years were on the transplant waiting list.
- For the first time, access in use at first dialysis, and calcium and phosphate data are reported.
- The death rate per 100 patient years was 15.0 for dialysis dependent patients (haemodialysis 14.7, peritoneal dialysis 15.9) and 2.4 for those with a functioning kidney transplant (cadaver donor 3.0, live donor 0.8).
- Of the 1,121 deaths among dialysis dependent patients in 2003, 40% were due to cardiovascular causes, 13% to infection, 22% to withdrawal from treatment and 7% from malignancy.
- Of the 139 deaths among patients with kidney transplants, 23% were due to cardiovascular causes, 30% due to malignancy and 17% to infection.
- There has been a 6% increase in the total number of dialysis dependent patients, and the number of patients increased in all States.
- The numbers of peritoneal dialysis dependent patients increased 2% from 1,785 to 1,823 in 2003.
- There were 543 kidney transplant operations performed in 2003, a transplant rate of 27 per million population.
- Of these, 40% (218 grafts) were from live donors, compared to 38% (230 grafts) in 2002. 22% of live donor operations were performed without the recipient receiving prior dialysis therapy.
- For primary cadaver grafts performed in 2002, the 12 month patient and graft survival rates were 97% and 94% respectively.
- The five year primary cadaver recipient and graft survival for operations performed in 1998 were 88% and 78% respectively.
- There were 5,951 functioning kidney transplants in Australia at 31 December 2003, a prevalence of 299 patients per million (a 4% increase over 2002).



KEY SUMMARY POINTS FROM THE REPORT

NEW ZEALAND

- There were 2,865 patients (715 per million) receiving renal replacement therapy (RRT) at 31 December 2003. Of these, 1,166 (291 per million) had a functioning kidney transplant, and 1,699 (424 per million) received dialysis treatment.
- 449 patients (112 per million) commenced RRT in 2003.
- The mean age at commencement was 56.7 years.
- Diabetic nephropathy accounted for 40% of new patients and glomerulonephritis 26%.
- Of patients <65 years of age, 25% were on the active kidney transplantation waiting list. 23% of Maoris and 13% of Pacific Islanders <65 years of age were on the transplant waiting list.
- The death rate per 100 patient years was 15.8 for dialysis dependent patients (haemodialysis 15.0, peritoneal dialysis 16.9) and 2.3 for those with a functioning kidney transplant (cadaver donor 2.4, live donor 2.0).
- Of the 263 deaths among dialysis dependent patients in 2003, 40% were due to cardiovascular causes, 11% to infection, 26% to withdrawal from treatment and 5% from malignancy.
- Of the 26 deaths among patients with a kidney transplant, 3% were due to cardiovascular causes and 38% due to malignancy.
- The number of patients who were dialysis dependent at 31 December 2003 (1,699) was an increase of 6% over the previous year. 59% of all dialysis dependent patients were receiving home dialysis. 77% of these were on peritoneal dialysis.
- The reported haemoglobin and use of erythropoietic agents have both increased over recent surveys.
- There were 111 kidney transplant operations performed in 2003, a rate of 28 per million population.
- The percentage of live donors in 2003 was 40%.
- For primary cadaver grafts performed in 2002, the 12 month patient and graft survival rates were 97% and 92% respectively.
- The five year primary cadaver recipient and graft survival for operations performed in 1998 were 82% and 73% respectively.