Legal	<b>Entity</b>	Name:
LUBA	LIICICY	Hailic.

PO Box 11060 Adelaide S.A. 5000

South Australia Health and Medical Research Institute Limited (T/A SAHMRI), the host agency for the Australian and New Zealand Dialysis and Transplant Registry (ANZDATA) and the Australian and New Zealand Organ Donation Registry (ANZOD)

Zediana organi ponation negistry (hivzob)
Legal Entity Number:
961428
Contract Period:
1 July 2020 to 30 June 2022
Contract Number:
360263 / 01
Reporting Period:
01/01/2021 to 30/06/2021
Invoice Due Date:
20/07/2021
20/01/2021
Contact person:
Kylie Hurst
Registry General Manager
Tababaa ayaabaa
Telephone number:
+61 8 8128 4747
Email:
kylie@anzdata.org.au
Postal address:
ANZDATA Registry
c/- SAHMRI

#### ADDITIONAL INFORMATION

#### **Compliance Statement:**

The organisation remains under the business name ANZDATA (Australia and New Zealand Dialysis and Transplant Registry) and ANZOD (Australia and New Zealand Organ Donation Registry) trading under the auspices of the SAHMRI (South Australian Health and Medical Research Institute) ABN 54 141 228 346.

Project materials remain in use in accordance with the current funding agreement and service provision continues to occur as per agreed timeframes.

A table with date of receipt of annual data output and proportion of data submitted electronically by each DHB contributing unit:

Centr Code	Hospital/Centre Name	Notification	Method 2020
STAR	Starship Children's	18/01/2021	Online
DUDN	Dunedin	01/03/2021	Online
WHAN	Whangarei	01/03/2021	Online
WELN	Wellington	27/04/2021	Online
HAWK	Hawkes Bay	27/04/2021	Online
WKTO	Waikato	03/05/2021	Paper
MIDM	Middlemore	06/05/2021	File Transfer
AUCK	Auckland City	15/06/2021	Online
TARA	Taranaki	16/06/2021	Online
CHCH	Christchurch	18/06/2021	Online
WMAT	Waitemata	30/06/2021	Online
PALM	Palmeston North	Outstanding at time of reporting	Online

### Status of real time reporting to DHBs

Reports are as listed in appendix A

## Update on activity during the reporting period related to:

- Data collection and processing continues to be streamlined.
  - Progress continues with transitioning units from paper to electronic data submission.
  - All, but one, unit took up the 2020 survey data collection via online data entry, with Middlemore the only centre to develop and utilise the Registry's file import functionality.
  - The ANZDATA NZPDR Data Exchange Agreement was executed, and technical support was provided to the New Zealand Peritoneal Dialysis Registry, for the implementation of webservice technology (application programming interface) to allow real-time data capture and submission between registries.
  - Next steps, in data collection, is to progress with linkage with the Ministry's Data and Digital from New Zealand national data collections.
  - The TeachPD trial continues with ongoing data collection and support.

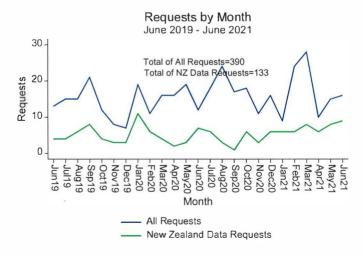
- o the ANZDATA NZ Blood Service Data Exchange Agreement (DEA) continues to be developed. This will enable streamlined and comprehensive organ wait list and transplant activity reporting and analysis. A privacy impact assessment is to be undertaken before the DEA can be signed off, to reassure compliance with recent changes to the NZ Privacy Act.
- Living kidney and deceased organ donation data collection continues.
- Development of New Zealand specific infographic representation of data collected continues.

### Collaboration with Aotearoa New Zealand Working Group continues.

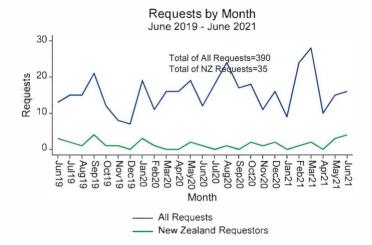
- o the group convened twice in the reporting period with a changeover in convenor to Dr Michael Collins due to the stepping down of Rachel Walker.
- o additionally, new members joined including an advanced trainee.
- the group has expanded the generation of research ideas and provided ongoing assistance on Aotearoa NZ specific data requests and promoting ANZDATA across the sector.
- expressions of interest were sought for a specific ANZDATA Quantitative Research Project (in collaboration with University of Otago). The successful applicant was Reshma Shettigar and the project has been progressing well.
- a guidance document was developed to provide ANZDATA advice for data requests involving New Zealand patients. This guidance takes into consideration patient ethnicity, disadvantaged groups such as children or minority populations and Māori participants to ensure that data requestors are aware of the context in which New Zealand patients share their information and that these data are managed and disseminated in a way that is used to benefit New Zealand patients.

### · Analytical support for New Zealand reporting

- Responding to requests for assistance with projects and enquiries for additional or new information continues. Projects including New Zealand data are circulated to the Registry's New Zealand Working Group and, in some cases, the New Zealand Renal Advisory Board when required for their input.
- o Figure 1 Requests by Month for Projects including NZ Data (Jun 2019-Jun 2021)



## o Figure 2 - Requests by Month for Projects from NZ Requestors (Jun 2019-Jun 2021)



# o Figure 3 - Requests by Quarter for Projects for only NZ Data (Jun 2019-Jun 2021)

