# Virtual Crossmatch Newsletter No. 3





The Virtual Crossmatch (VXM) Working Group is pleased to advise the next stage of the transition to VXM will commence from 18 October 2021.

# What you need to know

- All transplant units should ensure that patients who are on a transplant waitlist have now been enrolled in OrganMatch.
- ▶ The Organ Offer List (OOL) is changing and will no longer include complement-dependent cytotoxicity (CDC) results as part of the initial report. This means that OOL reports will be available sooner. An updated OOL report with CDC results will be published once they become available.

# **Enrolling waitlist patients in OrganMatch**

- ▶ All transplant waitlist patients must be enrolled in OrganMatch in order for tissue typing labs to complete HLA antibody screening and process transplant offers.
- Visit the <u>OrganMatch website</u> for further information on how to access the system and check patient enrolment status.

For transplant waitlist patients it is critical that:

- up to date serum is available for HLA antibody testing (within the last three months)
- any HLA antibodies have been identified and antigens for exclusion updated by the tissue typing lab
- details of any potential patient sensitising events such as blood transfusions are added into OrganMatch as soon as possible – this can be done from the Transplantation Portal directly.

## Changes to the Organ Offer List (OOL)

- 1 When there is a deceased donor, HLA typing will occur and the tissue typing lab will generate an initial OOL using OrganMatch this will:
  - **a include** a DSA assessment detailing the presence or absence of Donor Specific HLA antibodies
  - **b not include** CDC crossmatch results.
- 2 The initial OOL will be used by the DonateLife Agency and Transplant Units to make organ offers and commence allocation processes in most cases significantly earlier than currently provided.
- **3** Once CDC crossmatch results are available, a final OOL will be issued.

### What will OOL reports look like?

NSW Transplantation & Immunogenetics Services

**ORGAN OFFER LIST** 



| State Restricted Waiting |                          | Rank: 3        |                             | Score:    | Score: 40,000,110 |                    |  |
|--------------------------|--------------------------|----------------|-----------------------------|-----------|-------------------|--------------------|--|
| Donor                    | Name Withheld - D21-0879 | Donor Hospital | Bankstown Lidcombe Hospital | Recipient |                   | Eligible for offer |  |

#### Histocompatibility assessment

Overall match assessment

#### Histocompatibility summary

Class I DSA identified

CDC crossmatch is in progress - CDC crossmatch results to follow

### Donor HLA typing profile

| Α   | В   | С   | DRB1   | DQB1 | DQA1 | DPB1   | DPA1 | DRB3 | DRB4 | DRB5 |
|-----|-----|-----|--------|------|------|--------|------|------|------|------|
| *02 | *35 | *04 | *03:01 | *02  | *01  | *02:01 | *01  | *02  |      |      |
| *.  | *50 | *17 | *13    | *06  | *05  | *.     | *.   | *03  |      |      |

#### Recipient HLA typing profile

| A      | В      | С      | DRB1   | DQB1   | DQA1   | DPB1   | DPA1   | DRB3   | DRB4 | DRB5 |
|--------|--------|--------|--------|--------|--------|--------|--------|--------|------|------|
| *02:03 | *46:01 | *01:02 | *03:01 | *02:01 | *01:02 | *04:01 | *01:03 | *02:02 |      |      |
| *33:03 | *58:01 | *03:02 | *12:02 | *05:02 | *05:01 | *21:01 | *.     | *03:01 |      |      |

#### DSA Assessment

#### Assessment

Class I DSA identified



#### Where can I find out more information?

Further information about the project can be found below or by visiting the <a href="OrganMatch website">OrganMatch website</a>. A webinar hosted by TSANZ featuring VXM Clinical Lead A/Prof Ross Francis and VXM Laboratory Lead Rhonda Holdsworth will be held on 5 October 2021. Further information and can be found on the TSANZ website.

# Why are we transitioning to Virtual CrossMatch?

The Australian organ donation and transplantation system currently uses CDC crossmatches to help determine compatibility between organ donors and transplant recipients. Internationally, many transplant programs have moved to conducting virtual crossmatches (VXM) which can provide a more rapid assessment of compatibility without compromising transplant outcomes for low risk recipients.

This shift internationally has resulted in the equipment and reagents required for CDC being phased out, and Australian stocks will run out within 12–18 months. The aim in Australia is to transition to VXM for most transplant offers. A small number of physical crossmatches will still be required for selected high risk recipients – eventually this will be a flow crossmatch (FXM) instead of a CDC crossmatch.

#### When will the transition to VXM occur?

The transition to VXM will occur in three phases.

#### Phase 1 Phase 2 Phase 3 Phase 2a - October 2021 From July 2022 Now ▶ At the time of a deceased donor All transplant waitlist patients listed VXM processes will be introduced in OrganMatch. offer, an early OOL list w/o CDC for all transplant recipients. results will be generated using Frequency of HLA antibody ► A small number of prospective DSA assessment. screening (Luminex) increases from FXM may be performed in identified one to four times a year (and after ► CDC crossmatching will still be done sensitised recipients. any sensitising events). and an updated OOL generated Retrospective FXM will be later - this will allow for a faster Clinical transplant units work performed if the recipient is turnaround of suitability results than with tissue typing laboratories sensitised. is currently experienced. to identify antigens for exclusion for sensitised patients and list in Phase 2b - January 2022 OrganMatch. For kidney and pancreas patients – Donation offers will be made based CDC crossmatching will be limited on these exclusions in phase 2. to sensitised recipients only. All heart and lung transplant recipients will continue to have a CDC. ▶ Retrospective flow crossmatches (FXM) can be performed if required.

Regular post-implementation review will occur to ensure the effectiveness and safety of the National Histocompatibility Guidelines.