Update Post-Transplant Histocompatibility Data Collection

OPTN Histocompatibility Committee

OPTN ORGAN PROCUREMENT AND TRANSPLANTATION NETWORK

Purpose of Proposal

- Update post-transplant histocompatibility data collection instruments to be consistent with current testing methods
- Add data collection for virtual crossmatching
- Generate Discrepant HLA Typings reports for all potential HLA critical discrepancies

Background

- Donor Histocompatibility Form
 - Filled out within 60 days post-transplant by the original typing lab
 - Contains data on donor HLA typings performed
- Recipient Histocompatibility Form
 - Filled out within 60 days post-transplant by each recipient lab
 - Contains data on recipient HLA typing, HLA antibody screening, crossmatching, and donor retyping
- HLA Discrepant Typings Report
 - Filled out within 60 days after a non-equivalent HLA value is discovered between different HLA typings within the OPTN Computer System
 - Generated for discrepancies for A, B, and DR loci for kidney and pancreas donors and recipients

Proposal

- Donor Histocompatibility Form
 - Remove a net of two data elements related to separate dates and sample sources for HLA typings for Class I and Class II
 - Remove two data elements related to typing method (DNA vs. serology)
- Recipient Histocompatibility Form
 - Remove a net of four data elements related to separate dates and sample sources for HLA typings for Class I and Class II
 - Remove four data elements related to typing method (DNA vs. serology)
 - Remove outdated testing response options
 - Clarify multiple data collection fields/response options
 - Add data collection related to virtual crossmatching

Proposal

- Discrepant HLA Typings Report
 - Allow viewing/searching of report and data after submission
 - Add report information as read-only on relevant Donor and Recipient Histocompatibility forms
 - Update the list of discrepancy reasons and add definitions for clarity
 - Remove data collection for "discrepancy not resolvable"
 - Generate for HLA discrepancies for all organs and all loci (about 70 per year)

Rationale

- Updating and clarifying data collection will allow for more consistent and easier usage
- Data collection on virtual crossmatching will
 - Inform recipient treatment
 - Evaluate impacts of the practice on recipient outcomes, graft outcomes, and cold ischemic time
- Data collection for all potential critical discrepancies will
 - Increase awareness of HLA critical discrepancies
 - Allow for a system-wide perspective of critical HLA discrepancies
 - Better inform future policy updates related to critical HLA discrepancies

Member Actions

- Histocompatibility labs will need to be aware of revised data collection requirements
- Labs may have a slightly increased number of Discrepant HLA Typings reports to complete
 - There are about 70 discrepancies across the country the Discrepant Typings Report would generate for, some of which are already being generated

What do you think?

- Is the proposed list of discrepancy reasons comprehensive and clear? Are there any additional reasons you would recommend adding, or any you would recommend clarifying or taking away?
- Would the proposed changes to the Donor and Recipient
 Histocompatibility Forms be collected within discrete fields within a
 Laboratory Information System (LIS)? Please specify which data elements
 may not be collected discretely by all labs if relevant.
- Do you have usability recommendations for any of the post-transplant histocompatibility data collection instruments?

Additional Questions?

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Provide Feedback

Submit public comments on the OPTN website:

- January 23 March 19, 2024
- optn.transplant.hrsa.gov



Regional Meeting Information

Visit <u>https://optn.transplant.hrsa.gov/about/regions/regional-meetings/</u> for the latest regional meeting information and meeting materials

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Thank You For Listening!

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