POLICY STATEMENT

This policy outlines the minimum requirements for Confirmatory HLA Typing (HLA-CT) of patients and unrelated volunteer adult donors.

BUSINESS SECTION/DEPARTMENT

Repository and Laboratory Services Department

PURPOSE

This policy is meant to ensure that a potential donor is the source of the HLA assignment determined at the time of recruitment typing, and that the potential recipient is the source of the HLA assignment determined at the time of the unrelated donor search request. This policy is not meant to be used as a requirement to define final patient and donor match criteria. Transplant centers may decide to consider typing for additional HLA loci (i.e., DQB1, DPB1) or to resolve alternative HLA assignments that remain after the NMDP HLA-CT typing requirements are met.

SCOPE

This policy applies to National Marrow Donor Program (NMDP) Personnel and, where relevant, the NMDP Network. The policy content additionally may extend to other entities and persons, pursuant to contact.

RELATED DOCUMENTS

Form 22: available on the Network Web Site

Form 117: available on the Network Web Site

DEFINITIONS

Not applicable

RESPONSIBILITIES

Repository and Laboratory Services Department: develop and manage policy, publish/post policy so that it may be accessed by affected audiences.
REQUIREMENTS

Sample and Timing Requirements

HLA-CT of the potential recipient should be performed on a second independent sample prior to infusion and, preferably, prior to the unrelated donor search request. HLA-CT of the potential adult donor must be performed on a second independent sample prior to collection, preferably prior to patient conditioning.

HLA Typing Requirements

High resolution HLA-A, B, C, and DRB1 typing must be performed and reported on both the patient and all primary and back-up donors requested for work-up before HPC, Marrow or HPC, Apheresis collection can proceed. The common and well-documented (CWD) alleles, including CWD null alleles, that need to be clearly distinguished in order to meet the criteria for high resolution are described in the 2013 manuscript by Mack et al. 2013 (referenced below) or the most current published update to the CWD list. The CWD listing can be accessed at http://igdawg.org/cwd.html.

The NMDP does not require resolution of CWD alleles within G groups (defined at http://hla.alleles.org). However, laboratories are required to test for the following non-expressed alleles when they exist as a possibility within the assigned G group. A list of these null alleles within G groups is provided below.

<table>
<thead>
<tr>
<th>Null Allele</th>
<th>HLA G Group</th>
<th>Location of Polymorphism</th>
</tr>
</thead>
<tbody>
<tr>
<td>A*01:04N</td>
<td>A*01:01:01G</td>
<td>Exon 4</td>
</tr>
<tr>
<td>A*03:21N</td>
<td>A*03:01:01G</td>
<td>Exon 4</td>
</tr>
<tr>
<td>A*24:09N</td>
<td>A*24:02:01G</td>
<td>Exon 4</td>
</tr>
<tr>
<td>A*24:11N</td>
<td>A*24:02:01G</td>
<td>Exon 4</td>
</tr>
<tr>
<td>A*68:11N</td>
<td>A*68:01:02G</td>
<td>Exon 1</td>
</tr>
<tr>
<td>B*15:01:01:02N</td>
<td>B*15:01:01G</td>
<td>Intron 1</td>
</tr>
<tr>
<td>B*51:11N</td>
<td>B*51:01:01G</td>
<td>Exon 4</td>
</tr>
<tr>
<td>C*04:09N</td>
<td>C*04:01:01G</td>
<td>Exon 7</td>
</tr>
</tbody>
</table>

Serology may be used as an alternative to testing for the DNA polymorphism defining these null alleles. This approach should only be used in cases where the serology clearly distinguishes the expected specificities at the relevant locus. The final results of both the DNA and serology testing must then be reported to the NMDP.
Reporting Results to the NMDP

Final reporting of high resolution HLA typing results must meet one of the following criteria:

1) must contain only one unambiguously assigned genotype or

2) may contain multiple alternative genotypes if one includes two CWD alleles and the others do not include any alleles listed as CWD with the following exceptions:

a. Laboratories are not required to resolve CWD alleles within an HLA region that encodes identical protein sequences in the antigen recognition site (i.e., G assignment), with the exception of CWD null alleles as described above.

b. When alternative genotypes include combinations with one CWD allele plus an allele not designated as CWD, the resolution to a single genotype may not be required. If the laboratory possesses information or data relevant to the particular ambiguity being evaluated which provides evidence that the resolution to a single genotype may not be required, the laboratory may decide to claim a deviation from this policy and to report the result as completed.

Important to note: there have been numerous cases submitted to the NMDP and to the American Society for Histocompatibility and Immunogenetics demonstrating the clinical relevance of resolving these types of ambiguities. Therefore, it is highly recommended that laboratories critically evaluate each case and resolve these alternative genotypes as appropriate. This decision must be documented by the Laboratory Director and communicated to the NMDP together with the final typing results reported on the completed Form 22 and/or 117.

Along with other required forms, submission of patients' final laboratory reports to document decisions made or to supplement final HLA typing results being reported is highly encouraged. The laboratory report must clearly include all unresolved ambiguous HLA assignments. HLA laboratories must continue to adhere to all other accrediting agency standards and policies associated with high resolution DNA typing as defined by the American Society for Histocompatibility and Immunogenetics, the College of American Pathologists, or the European Federation for Immunogenetics.
REFERENCES


An electronic copy of this policy can be found at the NMDP Bioinformatics Website: http://bioinformatics.nmdp.org/Policies/Policies.aspx

REVISION HISTORY

<table>
<thead>
<tr>
<th>Revision</th>
<th>Brief Description of Revision</th>
</tr>
</thead>
<tbody>
<tr>
<td>P00079 rev. 1</td>
<td>Changed document type to policy and revised HLA typing</td>
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<tr>
<td></td>
<td>requirements. Previously controlled as A00261.</td>
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