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| <b>DATE EFFECTIVE: 12/31/11</b> | <b>List of Medical Tests</b> | <b>DOC: FRM.068</b><br><b>REV: B.01</b> |
|---------------------------------|------------------------------|---|

Testing is performed by Memorial Blood Center, which is under contract with Fairfax Cryobank and Cryogenic Laboratories, Inc.  
*Memorial Blood Center, 737 Pelham Blvd., St. Paul, MN 55114. CLIA ID # 24D0663800*

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|--|---|
| Test <b>Hepatitis B Surface Antigen</b>                    | Format EIA                                |
| Tradename(s) <a href="#">Genetic Systems HBsAg EIA 3.0</a> | Sample Serum/Plasma                       |
| Manufacturer Bio-Rad Laboratories                          | Use Donor Screen                          |
| Device Identifier <a href="#">32591, 32592, 25258</a>      | FDA Date Approved <a href="#">1/23/03</a> |

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|--|---|
| Test <b>Hepatitis B Core Antibody (IgG + IgM)</b>        | Format EIA                                |
| Tradename(s) <a href="#">Ortho HBc ELISA Test System</a> | Sample Serum/Plasma                       |
| Manufacturer Ortho-Clinical Diagnostics, Inc.            | Use Donor Screen                          |
| Device Identifier <a href="#">930740</a>                 | FDA Date Approved <a href="#">4/18/91</a> |

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| Test <b>Hepatitis C Antibody</b>                                     | Format EIA  |
| Tradename(s) <a href="#">Ortho HCV Version 3.0 ELISA Test System</a> | Sample Serum/Plasma                                 |
| Manufacturer Ortho-Clinical Diagnostics, Inc.                        | Use Donor Screen                                    |
| Device Identifier <a href="#">930740, 930750</a>                     | FDA Date Approved <a href="#">5/20/96, 2/18/09*</a> |

\* Donor screening test was approved 5/20/96 and was approved for living, cadaveric on 2/18/09.

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| Test <b>HIV1,2,0/HCV/HBV NAT</b>                      | Format PCR                                 |
| Tradename(s) <a href="#">COBAS TaqScreen MPX Test</a> | Sample Plasma                              |
| Manufacturer Roche Molecular Systems                  | Use Donor Screen                           |
| Device Identifier <a href="#">04584252 190</a>        | FDA Date Approved <a href="#">12/30/08</a> |

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|---|--|
| Test <b>HIV 1,2/ Antibody plus O</b>                                | Format EIA                               |
| Tradename(s) <a href="#">Genetic Systems HIV-1/HIV-2 Plus O EIA</a> | Sample Serum/Plasma                      |
| Manufacturer Bio-Rad Laboratories                                   | Use Donor Screen                         |
| Device Identifier <a href="#">32588,32589,25256</a>                 | FDA Date Approved <a href="#">8/5/03</a> |

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|--|---|
| Test <b>HTLV I/II Antibody</b>                                 | Format ChLIA                              |
| Tradename(s) <a href="#">Abbot Prism HTLV I/II Test System</a> | Sample Serum/Plasma                       |
| Manufacturer Abbott Laboratories                               | Use Donor Screen                          |
| Device Identifier <a href="#">6E50-68</a>                      | FDA Date Approved <a href="#">1/16/08</a> |

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|--------------------------|------------------------------|---------------------------|
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|--------------------------|------------------------------|---------------------------|

Test **Chlamydia trachomatis / Neisseria gonorrhoea\*** Format TMA  
 Tradename(s) [Gen-Probe Aptima Combo 2 Assay](#) Sample Genital swab/Urine  
 Manufacturer [Gen-Probe, Inc.](#) Use [Diagnosis](#)  
 Device Identifier [1032, 301130](#) FDA Date Approved [8/9/05](#)  
 \* Performed at Viomed Laboratories, Inc, 6101 Blue Circle Drive, Minnetonka, MN 55343. CLIA ID # 24D0400424

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Test **Cytomegalovirus (CMV) Total Antibody** Format IgG plus IgM  
 Tradename(s) [Immucor Capture CMV Total Antibody Test System](#) Sample Serum/Plasma  
 Manufacturer [Immucor, Inc.](#) Use [Donor Screen](#)  
 Device Identifier [0088](#) FDA Date Approved [12/22/95](#)

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Test **Treponema pallidum (Syphilis, IgG + IgM)** Format Microhemagglutination (MHA)  
 Tradename(s) [Beckman Coulter PK Anti-TP System](#) Sample Serum/Plasma  
 (formerly Olympus PK Anti-TP System)  
 Manufacturer Fujirebio Inc Use Donor Screen  
 Device Identifier [PH3000](#) FDA Date Approved [2/21/03](#)

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Test **Syphilis FTA-ABS** Format Treponemal  
 Tradename(s) Syphilis Antibody Detection by FTA-ABS Sample Serum/Plasma  
 Manufacturer MarDX Diagnostics Use Confirmatory  
 Device Identifier [31-8010](#) FDA Date Approved [NA](#)

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Test **Syphilis – RPR** Format Rapid Plasma Reagin (RPR)  
 Tradename(s) BD Macro-VueRPR Card Tests Sample Serum/Plasma  
 Manufacturer Becton Dickinson and Company Use Diagnosis  
 Device Identifier [274449](#) FDA Date Approved [see package insert](#)