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|  | Technical Data Sheet | Ref: TDS-008 |
| | Donor Donkey Serum | Rev: 0 |

| PRODUCT DESCRIPTION | | |
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| Biowest USA Donor Donkey Serum is a high-quality, aseptically collected serum derived from the clotted whole blood of healthy donkeys. Sourced from the United States, this serum undergoes rigorous testing and quality control to ensure purity, consistency, and traceability. | | |
| COUNTRY OF ORIGIN | | |
| The Biowest USA Donor Donkey Serum is sourced and collected in the United States, adhering to strict animal health and safety protocols to ensure the highest quality and traceability standards. | | |
| Catalog Number | Description | Origin(s) |
| S2800 | Donor Donkey Serum | United States |
| INTENDED USE | | |
| Biowest USA Donor Donkey Serum is intended for in vitro use in cell culture, research, and further manufacturing. It is not for human or animal consumption and is not an Active Pharmaceutical Ingredient (API). This product is strictly for laboratory and research applications. | | |
| COLLECTION SOURCE | | |
| Biowest USA Donor Donkey Serum is derived from clotted whole blood that is aseptically collected. Each batch is rigorously controlled from serum collection through all stages of treatment, production, and final packaging at our facilities. The serum is collected and treated in full compliance with USDA regulations to ensure quality and safety. | | |
| QUALITY CONTROL PARAMETERS | | |
| 1.1 | pH: monitored and reported for each batch, with specifications provided in the Certificate of Analysis (CoA). | |
| 1.2 | Osmolality: determined by the lowering of the freezing temperature, calibrated against standard solutions. Osmolality specifications are provided in the Certificate of Analysis. | |
| 1.3 | Endotoxin: tested using USP 85 Photometric Quantitative Techniques. Test results are reported in the Certificate of Analysis. | |
| 1.4 | Hemoglobin: measured by spectrophotometry. Results are provided in the Certificate of Analysis. | |
| 1.5 | Total Protein: determined using a colorimetric assay. Results are reported in the Certificate of Analysis. | |
| 1.6 | Sterility Testing: tested for the absence of aerobic and anaerobic bacteria, fungi, and yeast in compliance with 9CFR regulations. | |
| 1.7 | Mycoplasma: tested via culture-based methods or PCR (Polymerase Chain Reaction), depending on customer requirements. Tested for absence of Mycoplasma. | |
| 1.8 | Virus Testing: Not Applicable | |
| 1.9 | Other Testing: Additional testing may be available upon request. | |
| FILTRATION | | |
| Final Filter Size: 0.2µm | | |
| TREATMENT PROCESS | | |
| Not Applicable | | |
| STORAGE CONDITIONS | | |
| Store at ≤ -10°C, protected from light to maintain the serum’s integrity. | | |
| SHELF LIFE | | |
| 5 years from the date of manufacture when stored under the recommended conditions. | | |

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| HANDLING INSTRUCTIONS | |
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| 2.1 | Thawing: Thaw the serum in a refrigerator (2°C to 8°C) or at room temperature. Avoid rapid thawing methods to preserve protein integrity. |
| 2.2 | Aliquoting: For optimal preservation, aliquot the serum after thawing using aseptic techniques. |
| 2.3 | Storage after Thawing: If not all serum is used, thawed serum may be stored at 2°C to 8°C for up to 26 weeks without significant loss in quality, provided sterility is maintained. |
| 2.4 | Repeated Freeze/Thaw Cycles: To maintain serum quality, avoid repeated freeze/thaw cycles. Always refreeze aliquots, not the entire bottle. |
| PRECAUTIONS AND SAFETY | |
| 3.1 | For Research Use Only: Not for human or animal consumption. |
| 3.2 | Protective Equipment: Always wear appropriate PPE, such as gloves, lab coats, and face protection, when handling the serum. |
| 3.3 | Aseptic Handling: Ensure that serum is handled under aseptic conditions (e.g., laminar flow hood) to prevent contamination. |
| REGULATORY INFORMATION | |
| The serum is treated in accordance with strict animal health and safety standards to ensure the highest quality and traceability. Biowest USA adheres to all relevant regulatory guidelines and ensures that the product is safe, traceable, and compliant for research and laboratory applications. | |
| DISPOSAL INSTRUCTIONS | |
| Dispose of unused serum and packaging according to local regulations for biological materials and hazardous waste. | |
| CERTIFICATE OF ANALYSIS (CoA) | |
| A Certificate of Analysis is provided with each batch and contains detailed specifications, including test results for sterility, endotoxin levels, and other quality control parameters. | |
| DISCLAIMER | |
| This product is not intended as an Active Pharmaceutical Ingredient (API). It is intended for research, diagnostics, and medical device manufacturing only. Before use, users should refer to the Certificate of Analysis (CoA) for specific lot details. | |

For further information or inquiries, please contact Biowest USA Customer Service.