



PRODUCT DATA SHEET

Ready-to-Screen Tissue BLOTS[™] *Human Normal Skin Cell Tissue Blot*

| Catalog #: | TB52 | | | |
|--------------------|---|---|--|--|
| Lot #: | | | | |
| Components: | Protein blot of Human Normal Skin Cell Tissue samples arranged as follows: | | | |
| | Lane 1 Protein Marker* Lane 2 Epidermal Keratinocytes Adult (NHEK-Ad) Lane 3 Epidermal Keratinocytes Neo (NHEK-Neo) Lane 4 Epidermal Keratinocytes Neo Pool (NHEK-Neo Pool Lane 5 Dermal Fibroblast Adult (NHDF-Ad) Lane 6 Dermal Fibroblast Neo (NHDF-Neo) Lane 7 Microvascular Endothelial Adult (HMVECd-Ad) Lane 8 Microvascular Endothelial Neo (HMVECd-Neo) Lane 9 Lane 10 | 211,806 121,020 bl)100,216 54,395 38,708 29,806 20,040 7,331 | | Myosin β-galactosidase Bovine Serum Albumin Ovalbumin Carbonic Anhydrase Soybean Trypsin Inhibito Lysozyme Aprotinin |
| Size: | 1 Blot | 7,331 | | Aprodimi |

* Lot #: 300002325-BR

Storage Condition: 4° C

Methods Involved: The proteins were isolated from various <u>human normal skin cell tissues</u> by preparing a tissue homogenate in the presence of protease inhibitors. Protein samples (50μg) from each tissue were solubilized in SDS-lysis buffer and electrophoresed in a 10 well, 4-20% SDS-polyacrylamide gradient gel, followed by electroblotting on PVDF membrane.

Quality Control: Proteins isolated from each lot were run on 4-20% gel and stained with G-Biosciences RapidStain™ to check for its quality. Actin antibody was used to test the separation and transfer of protein from each lot.

Instructions for Use: Remove the blot (membrane) from the pouch and wash with an appropriate buffer (1X TBST or PBST) 1-2 times. Block the membrane with a protein blocking agent; e.g., G-Biosciences NAP™-BLOCKER or BLOT-QuickBlocker™, and incubate with the primary and secondary antibodies diluted in blocking solution, following the standard protocol. Develop the blot with chemiluminescent or chromogenic detection reagents for the detection of the specific protein.



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