

# 1. Technical Data Sheet

<b>Summary</b>	TruCytes™ TBMNK synthetic cells are lyophilized cell mimics that feature TBMNK biomarkers with scatter coordinates (FSC and SSC) that match lymphocyte, monocyte and granulocyte populations for lyse and wash conditions.																		
<b>Application</b>	<p>TruCytes™ TBMNK is intended to provide positive and negative signal detection for specified surface biomarkers that are targeted by specific antibodies. The following list shows the sub populations of synthetic cells that comprise the TBMNK:</p> <ul style="list-style-type: none"> <li>• CD4+ T-cells: CD45+, CD3+, CD4+</li> <li>• CD8+ T-cells: CD45+, CD3+, CD8+</li> <li>• B-cells: CD45+, CD3-, CD19+</li> <li>• NK cells: CD45+, CD3-, CD16/56</li> <li>• Classical monocyte: CD45+, CD14+</li> <li>• Non-classical monocyte: CD45+, CD16+</li> </ul> <table border="1" data-bbox="608 1088 1169 1379"> <thead> <tr> <th>TBMNK Biomarker</th> <th>Clone of Antibody</th> </tr> </thead> <tbody> <tr> <td>CD3</td> <td>SK7</td> </tr> <tr> <td>CD4</td> <td>SK3</td> </tr> <tr> <td>CD8</td> <td>SK1</td> </tr> <tr> <td>CD16</td> <td>B73.1</td> </tr> <tr> <td>CD19</td> <td>SJ25C1</td> </tr> <tr> <td>CD45</td> <td>2D1</td> </tr> <tr> <td>CD56</td> <td>NCAM16.2</td> </tr> <tr> <td>CD14</td> <td>M5E2</td> </tr> </tbody> </table> <p>*NOTE: TruCytes™ TBMNK Synthetic Cells have been designed to work with TBMNK antibody reagent kit containing the clones listed in the table below. Users are recommended to test the clones other than the listed below to determine the compatibility.</p> <p>It is an ideal process control for assays that have measured readouts using flow cytometry.</p> <p><b>For Research Use Only. Not for use in diagnostic or therapeutic procedures.</b></p>	TBMNK Biomarker	Clone of Antibody	CD3	SK7	CD4	SK3	CD8	SK1	CD16	B73.1	CD19	SJ25C1	CD45	2D1	CD56	NCAM16.2	CD14	M5E2
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<b>Materials</b>	TruCytes™ are hydrogels that are lyophilized for stability and ease of use. Each bottle contains approximately 2.5 x 10 <sup>5</sup> particles.																		
<b>Handling and Safety</b>	TruCytes™ TBMNK contains no bio-hazardous material so it is safe to use in any environment and requires no special disposal. See SDS at <a href="http://www.slingshotbio.com">www.slingshotbio.com</a> .																		

<p><b>Instructions for Use</b></p>	<ol style="list-style-type: none"> <li>1.1. Tap down the vial to ensure that all lyophilizate is collected at the bottom of the vial.</li> <li>1.2. Add 250 <math>\mu</math>L of PBS buffer. Make sure to not contact/disturb the lyophilizate with your pipette until the pellet has been soaked in the buffer. Gently pipette up and down to mix and ensure that all contents are fully dissolved before proceeding. Take the appropriate amount of reconstituted material out for each test.</li> <li>1.3. Add an appropriate amount of staining antibody (typically 5 <math>\mu</math>L standard test). Vortex mixture on high for 2 seconds to mix thoroughly.</li> <li>1.4. Incubate the mixture at RT in the dark for 15 - 30 min.</li> <li>1.5. Add 1 mL of 1X PBS, vortex, then transfer the reconstituted mixture to an appropriate tube.</li> <li>1.6. Centrifuge tube at 500 rcf for 5 min.</li> <li>1.7. Carefully pipette off the liquid without disturbing the pellet of particles.</li> <li>1.8. Repeat steps 5 - 7 to wash the particles again. Recommended: Perform 2 total washes to prevent non-specific binding.</li> <li>1.9. Add 100 <math>\mu</math>L of 1X PBS or staining buffer to the tube.</li> <li>1.10. View and acquire particles on FSC-A and SSC-A using the same instrument settings as real whole blood cells.</li> <li>1.11. For best results, set the flow rate on the cytometer to low.</li> </ol>
<p><b>Storage</b></p>	<p>Store lyophilized products at -20 °C upon receipt. Upon reconstitution, products should be used within seven (7) days when stored at 2 - 8 °C.</p>
<p><b>Expiration</b></p>	<p>One year from the date of manufacturing.</p>
<p><b>QC Data</b></p>	<p><b>Figure 1. Scattergram and Gating for TBMNK.</b> The following scattergrams show the gating strategy for proper detection of T, B, M, and NK subset populations.</p>