

**S1 Table. Quality controls (QCs) composition used in the qPCR assay.** Bovine RBC was 1:2 diluted with TE buffer pH 8.0 and different synthetic dsDNA (gBlocks, IDT) were used to spike the different QCs. RBC: red blood cells.

<b>Quality Control</b>	<b>Purpose</b>	<b>Content</b>
<b>QC1</b>	Negative control: No synthetic dsDNA added	<ul style="list-style-type: none"> <li>• Bovine RBC</li> <li>• 1x TE Buffer pH 8.0</li> </ul>
<b>QC2</b>	Positive control: Screening negative sample	<ul style="list-style-type: none"> <li>• Bovine RBC</li> <li>• 1x TE Buffer pH 8.0</li> <li>• <i>ACTB</i></li> <li>• <i>SMN1</i> (Exon-7)</li> <li>• <i>HBB</i> (CD-6)</li> <li>• sj-TREC</li> </ul>
<b>QC3</b>	Positive control: Screening positive sample	<ul style="list-style-type: none"> <li>• Bovine RBC</li> <li>• 1x TE Buffer pH 8.0</li> <li>• <i>ACTB</i></li> <li>• <i>SMN2</i> (Exon-7, c.840C&gt;T)</li> <li>• HbS allele (<i>HBB</i>: c.20A&gt;T)</li> </ul>