

Specification

**Bis-Tris for buffer solutions**

**A1025**

<b>Physical Description:</b>	Solid
<b>Product Code:</b>	A1025
<b>Product Name:</b>	Bis-Tris for buffer solutions
<b>Specifications:</b>	Assay (titr.): min. 99 % pH (1 %; H <sub>2</sub> O): 8.8 - 9.6 Heavy metals (as Pb): max. 0.0003 % Loss on drying: max. 1 % Fe: max. 0.0005 % A (1 cm/0.1 M in H <sub>2</sub> O) 340 nm: max. 0.02 280 nm: max. 0.04
<b>WGK:</b>	1
<b>Storage:</b>	RT
<b>Molecular Formula:</b>	C <sub>8</sub> H <sub>19</sub> NO <sub>5</sub>
<b>M:</b>	209.24 g/mol
<b>CAS:</b>	6976-37-0
<b>EINECS:</b>	230-237-7
<b>CS:</b>	29221900
<b>Comment</b>	<p>Bis-Tris is an important buffer for protein and nucleic acid systems. Additionally it is used as a substitute for cacodylic acid buffer systems (4). The working concentration, mentioned in the literature (ref. 1-3), is 10 - 25 mM.</p>
<b>Bibliography</b>	<p>(1)Massie, B. <i>et al.</i> (1995) <i>Bio/Technology</i> <b>13</b>, 602-608Improved vector for the production of HSV-ribonucleotide reductase R1 and R2. (2)Merabet, E. &amp; Ackers, G.K. (1995) <i>Biochemistry</i> <b>34</b>, 8554-8563Calorimetric analysis of the Icl-repressor binding to the DNA operator site. (3)Petri, V. <i>et al.</i> (1995) <i>Biochemistry</i> <b>34</b>, 9977-9984Characterization of the binding of the TATA-binding factor to adenovirus E4 promoter. (4)Scopes, R.K. ed. (1994) <i>Protein Purification (Principles and Practice)</i> <b>3rd Edition</b>, p. 352; Springer-Verlag.</p>

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