

## Specification

### Bromophenol Blue Sodium Salt for electrophoresis

**A3640**

<b>Physical Description:</b>	Solid
<b>Product Code:</b>	A3640
<b>Product Name:</b>	Bromophenol Blue Sodium Salt for electrophoresis
<b>Specifications:</b>	Assay (photometr.): min. 90 % $\lambda_{\text{max.}}$ (pH 4.6): 590 - 595 nm Loss on drying: max. 5 % Transition interval: passes test
<b>WGK:</b>	1
<b>Storage:</b>	RT
<b>Molecular Formula:</b>	$\text{C}_{19}\text{H}_9\text{Br}_4\text{NaO}_5\text{S}$
<b>M:</b>	691.94 g/mol
<b>CAS:</b>	34725-61-6
<b>EINECS:</b>	252-170-2
<b>CS:</b>	29349990
<b>Comment</b>	<p>Bromophenol blue (BPB) and xylene cyanol FF (XC) are the most widely used stains in gel electrophoresis for tracing the migration of samples on electrophoretic gels. Usually, both stains are added to the sample buffers (loading buffers; see our <i>ready-to-use loading buffers</i>) at concentrations of 0.05 % to 0.25 %. In the vertical polyacrylamide gel electrophoresis it may happen, that the gel is leaking or air bubbles may come up. If the gel is stained with 1 - 5 mg bromophenol blue, this may be detected much easier and earlier. In case that the stain will interfere with other applications (e. g. silver staining), a pre-run of the gel will remove bromophenol blue. It does not influence the migration of the samples. Add the solid stain before adding the catalyst (4). If DNA is dissolved in bidistilled water (pH 5.5 - 6.0) - not in TE buffer - both stains, as well bromophenol blue as xylene cyanol FF may change the migration rate during the electrophoresis run. The bands run faster than expected (5). In <b>denaturing</b> gels, the dye migrates with sizes like the following oligonucleotide: <b>Polyacrylamide (%)</b></p> <p><b>Bromophenol blue Xylene cyanol</b> (according to ref. 1, 4) 5 35 bases 130 bases 6 26 bases 106 bases 8 19 bases 75 bases 10 12 bases 55 bases 20 10 bases 28 bases</p>

#### AppliChem GmbH

Ottoweg 4 • D-64291 Darmstadt • Phone +49 6151 9357 0 • Fax +49 6151 9357 11 • [info.de@itwreagents.com](mailto:info.de@itwreagents.com) • [www.itwreagents.com](http://www.itwreagents.com)  
 CEO Joan Roget • Commerzbank Darmstadt • Bank 508 800 50 • Account 0186989900 IBAN DE24 5088 0050 0186 9899 00 • Swiftcode DRESDEFF508 • Finanzamt Darmstadt 07 228 16476 • Register court Darmstadt HRB Nr. 7340

### **Bibliography**

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