

# DATA SHEET

Version: 2  
Revision date: 17/02/23

## 1. Identification

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|---------------------|---|
| <b>Product name</b> | <b>10x Tris-Acetate-EDTA (TAE) Buffer, pH 8.3</b> |
| <b>Cat. No</b>      | BR0020  |

## 2. Description

TBE and TAE are the most commonly used buffers for DNA and RNA polyacrylamide gel electrophoresis. Both buffers are useful because they have a basic pH, this allowing migrations of the DNA through the gel toward the positive anode. TBE and TAE buffers are used for the analyses of DNA products resulting from PCR amplification, DNA purification, or DNA cloning experiments. TBE has high resolution for separating smaller DNA fragments but it is complicated recovery DNA from gel. TAE has a low ionic strength and buffering ability, it used to separating DNA larger than 1500 pb and easily recover the DNA from gel.

## 3. Specifications

**Chemicals:** Analytical grade.  
**RNase/DNase activity:** non-detectable  
**Format:** 10X solution  
**Volume:** 1000 ml  
**pH:** 8.3± 0.05 at 25 °C

## 4. Shipping and Storage specifications

TBE and TAE buffers are shipped at room temperature.  
Store the product in a dry place at room temperature.

## 5. Applications

- Running buffer and gels for RNA analysis native and denaturing
- Polyacrylamide and agarose gels
- Nucleic acid electrophoresis
- Transfer buffer in Northern Blotting

## 6. Directions for use

Dilute 10x solutions 10:1 to make a 1x working solution.  
1x buffer will contain 40 mM Tris, 20 mM acetic acid and 1 mM EDTA.

## 7. Further information

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| <b>Product Use Limitation</b> | This product is developed, designed and sold exclusively only for research purposes use. The product was not tested for use in diagnostics or for drug development, nor is it suitable for administration to humans or animals.  |
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