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## Product Information

### MOPS sodium salt SigmaUltra

Product Number **M5789**  
Store at Room Temperature

#### Product Description

Molecular Formula:  $C_7H_{14}NO_4SNa$

Molecular Weight: 231.2

CAS Number: 71119-22-7

$pK_a$ : 7.2 (25 °C)

Effective buffering range: pH 6.5 - 7.9

$\Delta pK_a/\Delta T$ : -0.015<sup>1</sup>

Synonyms: 3-morpholinopropanesulfonic acid sodium salt, 3-(N-morpholino)propanesulfonic acid sodium salt

Trace elemental analyses have been performed on the SigmaUltra MOPS sodium salt. The Certificate of Analysis provides lot-specific results. SigmaUltra MOPS sodium salt is for applications which require tight control of elemental content.

The zwitterionic buffer MOPS is a structural analog to the Good buffer MES. The Good buffers were developed in the 1960's for general use in biochemistry to meet the following criteria:

- midrange  $pK_a$
- maximum water solubility and minimum solubility in all other solvents
- minimal salt effects
- minimal change in  $pK_a$  with temperature,
- chemical and enzymatic stability,
- minimal absorption in visible or UV range
- reasonable ease of synthesis.<sup>2</sup>

The  $pK_a$  of MOPS (7.2) is closer to physiological pH than that of MES (6.1), and thus MOPS may be more suitable as a physiologically relevant buffer.

MOPS buffer has been utilized in the culture of cells in such systems as *E. coli*, *Cryptococcus neoformans*, cultured human keratinocytes, and thermophilic methanogenic bacteria.<sup>3,4,5,6</sup> In protein studies, MOPS has been used in an X-ray crystallographic study of the ADP-binding site of succinyl-CoA synthetase from *E. coli*, in the characterization of the Rieske-type

ferredoxin BphF, and in an electron microscopy analysis of the engineered protein betabellin-15D.<sup>7,8,9</sup>

An investigation of the interaction of various buffers, including MOPS, with plasmid sized DNA by free solution capillary electrophoresis has been reported.<sup>10</sup> A protocol describes the use of MOPS in an electrophoresis buffer for the separation of RNA in agarose gels.<sup>11</sup> A procedure for preparative-scale separation of proteins by displacement chromatography that incorporates MOPS buffer has been published.<sup>12</sup>

### **Precautions and Disclaimer**

For Laboratory Use Only. Not for drug, household or other uses.

### **Preparation Instructions**

This product is soluble in water (231 mg/ml).

### **Storage/Stability**

Solutions of MOPS are not completely stable when autoclaved in the presence of glucose.<sup>13</sup> Solutions of MOPS turn yellow when autoclaved, indicating that MOPS is unstable to autoclaving.

### **References**

1. Ellis, K. J. and Morrison, J. F., Buffers of constant ionic strength for studying pH-dependent processes. *Methods Enzymol.*, **87**, 405-426 (1982).
2. Good, N. E., et al, Hydrogen ion buffers for biological research. *Biochemistry*, **5(2)**, 467-477 (1966).
3. Tucker, D. L., et al., Gene expression profiling of the pH response in *Escherichia coli*. *J. Bacteriol.*, **184(23)**, 6551-6558 (2002).
4. Petrou, M. A., and Shanson, D. C., Susceptibility of *Cryptococcus neoformans* by the NCCLS microdilution and Etest methods using five defined media. *J. Antimicrob. Chemother.*, **46(5)**, 815-818 (2000).

5. Sando, G. N., et al., Induction of ceramide glucosyltransferase activity in cultured human keratinocytes. Correlation with culture differentiation. *J. Biol. Chem.*, **271(36)**, 22044-22051 (1996).
6. Foster, M. S., et al., Improved methods for the cultivation of strictly anaerobic, extremely thermophilic methanogens. *Biotechniques*, **15(6)**, 996-998, 1000, 1002 (1993).
7. Joyce, M. A., et al., ADP-binding site of *Escherichia coli* succinyl-CoA synthetase revealed by x-ray crystallography. *Biochemistry*, **39(1)**, 17-25 (2000).
8. Couture, M. M., et al., Characterization of BphF, a Rieske-type ferredoxin with a low reduction potential. *Biochemistry*, **40(1)**, 84-92 (2001).
9. Lim, A., et al., Engineering of betabellin-15D: a 64 residue  $\beta$ -sheet protein that forms long narrow multimeric fibrils. *Protein Sci.*, **7(7)**, 1545-1554 (1998).
10. Stellwagen, N. C., et al., DNA and buffers: are there any noninteracting, neutral pH buffers? *Anal. Biochem.*, **287(1)**, 167-175 (2000).
11. *Molecular Cloning: A Laboratory Manual*, 3rd ed., Sambrook, J. and Russell, D.W., CSHL Press (Cold Spring Harbor, NY: 2001), p. 7.32.
12. Narahari, C. R., et al., Displacement chromatography of proteins using a self-sharpening pH front formed by adsorbed buffering species as the displacer. *J. Chromatogr. A*, **825(2)**, 115-126 (1998).
13. *The Merck Index*, 12th ed., Entry# 6346.

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