

## Product Information

### 2,6-Diaminopurine hemisulfate salt

Product Number **D 3289**  
Store at Room Temperature

#### Product Description

Molecular Formula:  $C_5H_6N_6 \bullet \frac{1}{2} H_2SO_4$   
Molecular Weight: 199.2  
CAS Number: 69369-16-0  
Melting Point: 302 °C<sup>1</sup>  
 $\lambda_{max}$ : 241 nm, 282 nm (pH 1.9);<sup>1</sup> 247 nm,  
280 nm (0.1 M phosphate, pH 7.0)<sup>2</sup>  
Extinction Coefficient:  $E^{mM} = 9.55$  (241 nm),  
10 (282 nm) (pH 1.9);<sup>1</sup> 7.57 (247 nm), 9.05 (280 nm)  
(0.1 M phosphate, pH 7.0)<sup>2</sup>  
Synonyms: DAP hemisulfate; 2-aminoadenine  
hemisulfate; 1*H*-purine-2,6-diamine hemisulfate;  
2,6-diamino-9*H*-purine hemisulfate<sup>1</sup>

2,6-Diaminopurine is an adenine analogue that is an antagonist of naturally occurring purines.<sup>1</sup> DAP can base pair with thymidine in DNA, and with uracil in RNA, to give three Watson-Crick hydrogen bonds. This alteration in base pairing properties has led to the use of DAP as a structural probe of molecular recognition between ligands and DNA.<sup>2</sup>

DAP has been incorporated into anhydrohexitol nucleosides for the preparation of hexitol nucleic acids, and subsequent hybridization studies with DNA and RNA.<sup>4</sup> Ligase ribozymes that contain DAP and uracil have been prepared by *in vitro* evolution, and have been found to catalyze the template-directed joining of two RNA molecules.<sup>5</sup> A DAP moiety has been incorporated in the synthesis of various L- $\beta$ -(2*S*,4*S*)- and L- $\alpha$ -(2*S*,4*R*)-dioxolanyl nucleosides as potential anti-HIV compounds.<sup>6</sup>

The cyanophage S-2L is capable of using DAP in place of adenine in its DNA.<sup>7</sup> *Salmonella typhimurium* is also able to utilize DAP as a purine source.<sup>8</sup>

#### Precautions and Disclaimer

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

#### Preparation Instructions

This product is soluble in formic acid (50 mg/ml), with heat as needed, yielding a clear to hazy, yellow to yellow-green solution.

#### References

1. The Merck Index, 12th ed., Entry# 3028.
2. Montgomery, J. A., Holum, L. B., Synthesis of Potential Anticancer Agents. XI. N<sup>2,6</sup>-Alkyl Derivatives of 2,6-Diaminopurine. *J. Am. Chem. Soc.*, **80(2)**, 404-408 (1958).
3. Bailly, C., and Waring, M. J., The use of diaminopurine to investigate structural properties of nucleic acids and molecular recognition between ligands and DNA. *Nucleic Acids Res.*, **26(19)**, 4309-4314 (1998).
4. Boudou, V., et al., Base pairing of anhydrohexitol nucleosides with 2,6-diaminopurine, 5-methylcytosine and uracil as base moiety. *Nucleic Acids Res.*, **27(6)**, 1450-1456 (1999).
5. Reader, J. S., and Joyce, G. F., A ribozyme composed of only two different nucleotides. *Nature*, **420(6917)**, 841-844 (2002).
6. Kim, H. O., et al., L- $\beta$ -(2*S*,4*S*)- and L- $\alpha$ -(2*S*,4*R*)-dioxolanyl nucleosides as potential anti-HIV agents: asymmetric synthesis and structure-activity relationships. *J. Med. Chem.*, **36(5)**, 519-528 (1993).
7. Kirnos, M. D., et al., 2-aminoadenine is an adenine substituting for a base in S-2L cyanophage DNA. *Nature*, **270(5635)**, 369-370 (1977).
8. Garber, B. B., and Gots, J. S., Utilization of 2,6-diaminopurine by *Salmonella typhimurium*. *J. Bacteriol.*, **143(2)**, 864-871 (1980).

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