



## Product Information Sheet

### T834 Tobramycin Sulfate

Synonyms: O-3-Amino-3-deoxy- $\alpha$ -D-glucopyranosyl-(1 $\rightarrow$ 6)-  
O-[2,6-diamino-2,3,6-trideoxy- $\alpha$ -D-ribo-  
hexopyranosyl-(1 $\rightarrow$ 4)]-2-deoxy-D-Streptamine  
Sulfate

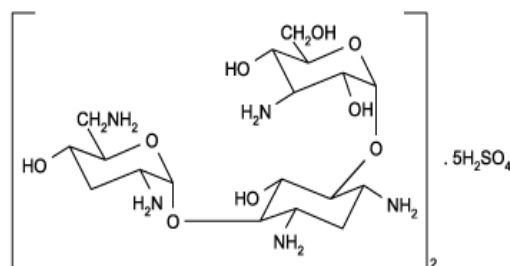
CAS: 79645-27-5

Formula:  $(C_{18}H_{37}N_5O_9)_2 \cdot 5H_2SO_4$

Mol. Weight: 1425.4

#### Properties

Form: Powder  
Appearance: White to Cream Powder  
Application: Plant Tissue Culture Antibiotic  
Solubility: Soluble in Water  
Storage Temp: 2 to 6 °C  
Other Notes: Average Activity: 634  $\mu$ g/mg



#### Application Notes

Tobramycin is an aminoglycoside antibiotic derived from *Streptomyces tenebrarius*. Its mode of action and activity are similar to those of gentamicin. It inhibits the initiation, elongation and termination of protein synthesis by binding to the 30S subunit and sometimes the 50S subunit of the bacterial ribosome.<sup>2</sup> Tobramycin is active against the *Enterobacteriaceae* and *Staphylococcus aureus*; it has been reported that tobramycin is more effective than gentamicin against *Pseudomonas* species.<sup>2,3</sup>

Minimum inhibitory concentration (MIC) of tobramycin has been reported for many different species. MIC for *Staphylococcus aureus* is less than 0.075  $\mu$ g/mL, *Escherichia coli*, *Pseudomonas*, and *Klebsiella-Enterobacter* are less than 0.65  $\mu$ g/mL, *Proteus mirabilis* and *Serratia marcescens* are 2.5  $\mu$ g/mL, and *Proteus rettgeri* is greater than 5  $\mu$ g/mL.<sup>3</sup>

Please Note: It is the sole responsibility of the purchaser to determine the appropriateness of this product for the specific plants that are being cultured and applications that are being used.

#### References

1. Merck 13, 9567
2. Martindale: The Complete Drug Reference, 35th ed., Paul S. Blake, Ed. (Royal Pharmaceutical Society, 2007), p. 316.
3. Britt, Michael R., Richard A. Garibaldi, James N. Wilfert, and Charles B. Smith. 1972. *In vitro* activity of tobramycin and gentamicin. *Antimicrobial Agents and Chemotherapy*. Vol 2(3):236-241.

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