

Dissociation Reagents

DNase I

For digestion of DNA

Catalog # 07469 25 mg
07470 100 mg
100-0683 1 g



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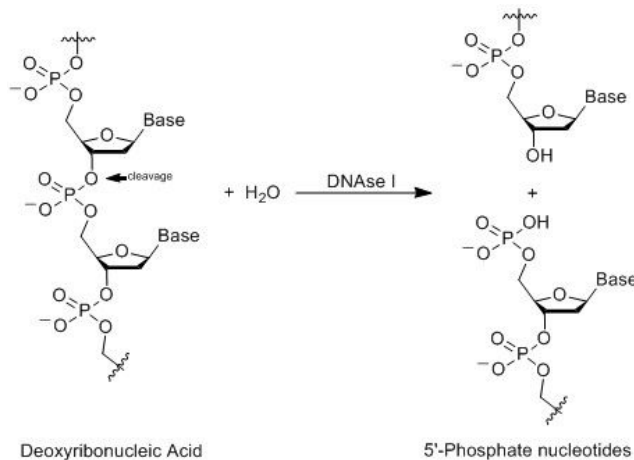
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Product Description

Deoxyribonuclease I (DNase I) is an endonuclease consisting of a single glycosylated polypeptide chain with two disulfide bonds. DNase I is often included in tissue dissociation protocols to digest DNA that has leaked into the dissociation medium as a result of cell damage. DNase I preferentially cleaves phosphodiester linkages adjacent to pyrimidine nucleotides in both single- and double-stranded DNA, yielding polynucleotides with 5'-phosphate and 3'-hydroxyl groups (Bernardi et al.). DNase I has been used for the dissociation of human tissues such as microglia (Klegeris & McGeer), cartilage (Dunham & Koch), colon (Fukushima & Fiocchi), epithelium (Fukushima & Fiocchi), liver (Vatakis et al.), lung (Fujino et al.), and neural (Fuja et al.), and for dissociation of stem cells (Kusuma et al.).

Product Information

Alternative Names: DNA endonuclease; DNA nuclease; Deoxyribonucleic phosphatase; Pancreatic DNase; Thymonuclease
Format: Lyophilized powder
Storage: Store at 2 - 8°C.
Stability: Stable as supplied for 12 months from date of receipt.
Reconstitution: Dissociation reagents can be reconstituted in a balanced salt solution or buffer of choice.
Molecular Weight: 29.1 kDa
CAS Number: 9003-98-9
Optimum pH: 7.8
Cleavage Site: DNase I preferentially splits phosphodiester linkages adjacent to a pyrimidine nucleotide. This yields 5'-phosphate terminated polynucleotides with a free hydroxyl group at the 3' position.



Cleavage site of DNase I

Please refer to the Safety Data Sheet (SDS) for hazard information.

Specifications

Source: Bovine pancreas
Activity: ≥ 2000 units/mg dry weight. See Notes for further information.

Related Products

For a complete list of dissociation reagents, as well as related products available from STEMCELL Technologies, visit www.stemcell.com or contact us at techsupport@stemcell.com.

Notes

ACTIVITY UNITS

1 unit refers to the amount of DNase I required to act on 1 mg/mL of DNA (pH 5.0, 25°C) to produce an increase in absorbance of 0.001 per minute at a wavelength of 260 nm.

References

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