

Dissociation Reagents

DNase I, ACF

Animal component-free DNase I for digestion of DNA

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|-----------|----------|---------------|
| Catalog # | 07473 | 25,000 Units |
| | 07474 | 100,000 Units |
| | 100-0273 | 500,000 Units |



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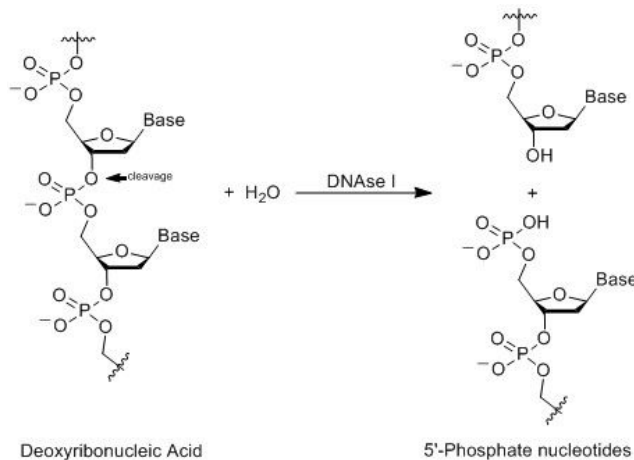
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Product Description

Deoxyribonuclease I (DNase I), Animal Component-Free (ACF) is obtained from cultures free of animal-derived materials and purified by chromatography. DNase I is an endonuclease consisting of a single glycosylated polypeptide chain with two disulfide bonds. DNase 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, ACF 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

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| 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

Specifications

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| Source: | Pichia pastoris |
| 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 causes an increase in absorbance of 0.001/minute/mL at 260 nm at 25°C, pH 5.0 when acting upon highly polymerized DNA.

References

- Bernardi A et al. (1975) The specificity of five DNAases as studied by the analysis of 5'-terminal doublets. *Eur J Biochem* 52(3): 451–7.
- Dunham BP & Koch RJ. (1998) Basic fibroblast growth factor and insulinlike growth factor I support the growth of human septal chondrocytes in a serum-free environment. *Arch Otolaryngol Head Neck Surg* 124(12): 1325–30.
- Fuja TJ et al. (2004) Asymmetric localization of LGN but not AGS3, two homologs of *Drosophila* pins, in dividing human neural progenitor cells. *J Neurosci Res* 75(6): 782–93.
- Fujino N et al. (2011) Isolation of alveolar epithelial type II progenitor cells from adult human lungs. *Lab Invest* 91(3): 363–78.
- Fukushima K & Fiocchi C. (2004) Paradoxical decrease of mitochondrial DNA deletions in epithelial cells of active ulcerative colitis patients. *Am J Physiol Gastrointest Liver Physiol* 286(5): G804–13.
- Klegeris A & Mcgeer PL. (2005) Chymotrypsin-like proteases contribute to human monocytic THP-1 cell as well as human microglial neurotoxicity. *Glia* 51(1): 56–64.
- Kusuma GD et al. (2015) Ectopic bone formation by mesenchymal stem cells derived from human term placenta and the decidua. *PLoS One* 10(10): e0141246.
- Li C-S et al. (2016) Fibromodulin reprogrammed cells: A novel cell source for bone regeneration. *Biomaterials* 83: 194–206.
- Li L & Schust DJ. (2015) Isolation, purification and in vitro differentiation of cytotrophoblast cells from human term placenta. *Reprod Biol Endocrinol* 13(1): 71.
- Moro K et al. (2015) Isolation and analysis of group 2 innate lymphoid cells in mice. *Nat Protoc* 10(5): 792–806.
- Patel J et al. (2013) Prospective surface marker-based isolation and expansion of fetal endothelial colony-forming cells from human term placenta. *Stem Cells Transl Med* 2(11): 839–47.
- Su CTE et al. (2015) An optogenetic approach for assessing formation of neuronal connections in a co-culture system. *J Vis Exp* (96): 1–9.
- Vatakis DN et al. (2012) Using the BLT humanized mouse as a stem cell based gene therapy tumor model. *J Vis Exp* (70): e4181.

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