## Close relationship between the HinfI and DpnA DNA-methyltransferase

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The amino acid sequences of some thirty different DNA methyltransferases (Mtases) have been established during the last several years. Their comparison revealed a close relationship among all the cytosine (5C) specific enzymes (1). In contrast high divergence is found between the adenine specific Mtases. Only the five Mtases dam, damT4, DpnII, EcoRV and damT2 proved to be closely related (2,3). Table I shows, that the sequence of the GANTC recognizing HinfI Mtase, which was recently published (4), does not only exhibit some identities to the isomethylomeric enzyme HhaII (5), but to a much larger extent similarities to the GATC methylating enzyme DpnA (5,6). In the alignment 42% of the positions are occupied by identical amino acids. The comparison thus defines a third group of closely related Mtases. It also supports the previous observation that recognition of different target sequences may well be achieved by Mtases whose core structure is conserved (1,2).

## Table 1

DpnA	MTKPYYNKNKMILVHSDTFKFLSKMKPESMDMIFADPPYFLSNGG ISNSGGQVVSVDKGDWDKISSFEEKH
HinfI	
DpnA	EFNRKWIRLAKEVLKPNGTVWISGSLHNIYSVGMALEQEGFKILNNITWQKTNPAPNLSCRYFTHSTETILW
HinfI	
DpnA	ARKNDKKARHYYNYDLMKELNDGKQMKDVWTGSLTKKVEK WAGK HPTQKPEYLLERIILASTKEGDY
HinfI	
DpnA	ILDPFVGSGTTGVVAKRLGRRFIGIDAEKEYLKIARKRLEAENETN
HinfI	
HinfI	RVGQTLFDKNENAICIVTQDGNVKDNEETLSIHKMSAKYLNKTNNNGWDYFYLFRNNNFITLDSLRYEYTNQ

<u>Table 1:</u> Alignment of the two DNA Mtase sequences <u>Dpn</u>A (5) and <u>Hinfl</u> (4). The alignment was performed by hand. The number of gaps/inserts was kept to a minimum. Conserved amino acids are indicated by vertical lines. The two boxes indicate regions, which are conserved in all adenine specific Mtases (7).

## REFERENCES:

- 1. Lauster, R., Trautner, T.A. and Noyer-Weidner, M. (1989). J. Mol. Biol. 206, in press.
- 2. Lauster, R., Kriebardis, A. and Guschlbauer, W. (1987). FEBS Letters 220, 167-176.
- Miner, Z. and Hattman, S. (1988). J. Bacteriol. 170, 5177-5184
- 4. Chandrasegaran, S., Lunnen, K.D., Smith, H.O. and Wilson, G.G. (1988), Gene 70, 387-392.
- 5. Lacks, S.A., Mannarelli, B.M., Springhorn, S.S. and Greenberg, B. (1986). Cell 46, 993-1000.
- de la Campa, A.G., Purushottam, K., Springhorn, S.S. and Lacks, S.A. (1987). J. Mol. Biol. 196, 457-469.
- 7. Lauster, R. (1989). J. Mol. Biol. 206, in press.