

Table 2. Steady-state kinetic parameters of the apurinic/apyrimidinic (AP) endonuclease, 3'-repair diesterase and nucleotide incision repair (NIR) activities of the Nfo proteins obtained from different preparations

Substrate	Enzymes					
	Nfo*			Nfo**		
	K_M , nMb	k_{cat} , min ⁻¹	k_{cat}/K_M , min ⁻¹ · μ M ⁻¹	K_M , nMb	k_{cat} , min ⁻¹	k_{cat}/K_M , min ⁻¹ · μ M ⁻¹
α A•T	24 ± 2.1	14 ± 0.3	580	38 ± 5.1	0.85 ± 0.1	20
THF•T	16 ± 2.0	9.3 ± 1.2	580	2.0 ± 0.5	5.4 ± 1.1	2,700
3'THF ^{NICK}	ND	ND	ND	7.7 ± 1.9	2.8 ± 0.16	360
α T•A	7.5 ± 3.0	11.0 ± 0.2	1,500	42 ± 6.2	4.8 ± 0.8	110
THF•A	3.8 ± 0.8	4.5 ± 0.2	1,200	8.0 ± 1.0	18 ± 2.2	2,200
DHU•G	41 ± 4.1	4.6 ± 0.2	110	74 ± 7	0.24 ± 0.02	3.2
DHT•A	13 ± 2.0	2.0 ± 0.1	150	38 ± 8	0.25 ± 0.01	6.6
5ohU•G	71 ± 11.0	0.9 ± 0.06	13	316 ± 44	1.6 ± 0.08	5.1
me-Fapy•C	290 ± 50	2.2 ± 0.1	7.6	250 ± 70	0.95 ± 0.1	3.8
THF•G	23 ± 1.8	21 ± 0.8	910	1.3 ± 0.2	1.7 ± 0.1	1,300
(30 mer)						

Nfo* and Nfo** were purified as described in refs. 1 and 2. Kinetic constants values in bold have been published previously (2, 3). α A, α -2'-deoxyadenosine; α T, α -thymidine; THF, tetrahydrofuran; DHU, 5,6-dihydrodeoxyuridine; DHT, 5,6-dihydrothymidine; 5ohU, 5-hydroxy-2'-deoxyuridine; me-Fapy, 2,6-diamino-4-hydroxy-5-N-methylformamidopyrimidine.

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