

Short Communication

HYDROLYSIS OF URIDINE  
DIPHOSPHATE-N-ACETYL-D-GLUCOSAMINE BY NORMAL  
AND MALIGNANT CELLS OF THE RAT

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THE OBJECTIVE of this study was to determine whether the 2-step hydrolysis of uridine diphosphate-N-acetyl-D-glucosamine (UDPGlcNAc) may serve as a marker of the malignant transformation of the rat cell. UDPGlcNAc is cleaved by nucleotide pyrophosphatase to UMP and

coma virus (RSHT) were obtained from Flow Laboratories. Rat tumour cells were cultured from fibrosarcomas induced by N-hydroxy-2-fluorenylbenzamide (RT-4 and RT-5) or from sarcomas induced by N-OH-2-FAA (D'I/I/3T

TABLE I.—*Hydrolysis of UDP[6-<sup>3</sup>H]GlcNAc by control rat embryo cells and by rat embryo*

cells exposed once or twice to N-OH-2-FAA					
		% of UDP[6- <sup>3</sup> H]GlcNAc converted			
		% of			
		Passage	UDP[6- <sup>3</sup> H]-	to [6- <sup>3</sup> H]	to [6- <sup>3</sup> H]
Cell line†	no. of cells	GlcNAc remaining*	hexose-NAc-1-P	hexose-NAc	
Exposed lines I and II	12	2 ± 1	78 ± 5	20 ± 5	
	16	4 ± 1	75 ± 5	21 ± 4	
	20	2 ± 1	79 ± 5	19 ± 4	
	24	3 ± 1	72 ± 7	24 ± 7	
	28	2 ± 1	76 ± 2	22 ± 2	
		(3 ± 1)	(75 ± 3)	(22 ± 3)	
	8	3	55	42	
	12	2 ± 1	70 ± 10	28 ± 11	
	20	14 ± 2	68 ± 9	18 ± 6	
	24	6	59	35	
Control lines III and IV	28	15 ± 10 (8 ± 6)‡	71 ± 11 (65 ± 7)‡	14 ± 1 (27 ± 10)	
	16	3 ± 1	79 ± 1	18 ± 2	
	20	3 ± 1	72 ± 6	25 ± 5	
	24	3 ± 1	74 ± 2	23 ± 1	
	28	3 ± 1	77 ± 1	20 ± 2	
Exposed lines III and IV		(3 ± 1)	(76 ± 3)	(22 ± 3)	
	16	8	70	22	
	20	17	60	23	
	24	38 ± 11	47 ± 6	15 ± 5	
	28	17 ± 4 (20 ± 13)‡	69 ± 9 (61 ± 11)‡	14 ± 6 (19 ± 5)	

\* The values in columns 3, 4 and 5 are the means + average deviations of two cell lines, except that values

hydrolysis of UDPGLcNAc by rat embryo

*GlcNAc by normal and tumour cells of rat*

cells exposed to N-OH-2-FAA raised the  
question as to whether the change in

*and hamster*

*et al.*, 1978).

peculiar to the transformed cells of the

The lack of correlation between malignant transformation of the rat embryo cell and the decrease of the cleavage of hamster. However, impairment of this

reaction is not indicative of malignant transformation of the rat cell, irrespective