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COMMON SOUTHERN CHINESE GLUCOSE-6-PHOSPHATE DEHYDROGNEASE MUTATIONS IN SINGAPORE AND TAIWAN

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Glucose-6-phosphate dehydrogenase (G6PD) deficiency is a common X-linked genetic disorder in Southeast Asia. Recently 6 nucleotide mutations have been reported to account for 80% to 90% of G6PD mutant alleles in Taiwan. A study was undertaken to find out the incidence of those common southern Chinese G6PD mutations in Singapore. Blood samples collected on fliter paper (Tokyo Roshi Kaisha Ltd, Tokyo) from 127 and 166 male G6PD deficient patients in Singapore and Taiwan respectively were tested. A non-radioactive method was used to detect the 6 common mutations. This involved the analysis of restriction fragments of DNA amplified directly from dried blood spot by polymerase chain reaction. Results of the study are summarised as follows:

Nucleotide Mutation			1388	1376	1024	493	487	95	Unknown
Singapore Hokkien Cantonese		8	20	1	0	0	1	4	
		14	18	2	0	0	7	5	
	Hakka		1	6	1	0	0	1	0
Other dialect			ct 3	6	0	0	0	0	2
Total	Chinese	(n=100)	26(26%)	50(50%)	4(4%)	0	0	9(9%)	11(11%)
	Malay	(n=25)	1	0	1	0	0	0	23
	Indian	(n=2)	0	0	0	0	0	0	2
Taiwan	Hokkien		20	69	7	10	1	12	18
	Cantonese		1	0	. 0	0	0	0	1
	Hakka		4	6	1	2	0	3	6
	Other dialec		ot 1	0	1	1	0	1	1
Total Chinese (n=166) 26(16%)			75(45%)	9(5%)	13(8%)	1	16(10%	26(16%)	

Paper to be presented by: __TAN It Koon

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