CHARACTERISATION OF THE HUMAN THYROID PEROXIDASE GENE MUTATION(S) IN PATIENTS WITH CONGENITAL DYSHORMONOGENETIC HYPOTHYROIDISM

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THESIS SUBMITTED IN FULFILMENT OF THE REQUIREMENTS FOR THE DEGREE OF DOCTOR OF PHILOSOPHY

DEPARTMENT OF MOLECULAR MEDICINE FACULTY OF MEDICINE UNIVERSITY OF MALAYA KUALA LUMPUR 2013
UNIVERSITI MALAYA

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