# **UNIVERSITI TEKNOLOGI MARA**

# PROTEOMIC PROFILING FOR POTENTIAL BIOMARKERS OF OSTEOSARCOMA PATIENTS

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Thesis submitted in fulfilment of requirements for the degree of **Master of Science** 

**Faculty of Medicine** 

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## **CONFIRMATION BY PANEL OF EXAMINERS**

I certify that a Panel of Examiners has met on 21<sup>st</sup> April 2016 to conduct the final examination of Zulaika Binti Roslan on her Master of Science thesis entitled "Proteomic Profiling for Potential Biomarkers of Osteosarcoma Patients" in accordance with Universiti Teknologi MARA Act 1976 (Akta 173). The Panel of Examiners recommends that the student be awarded the relevant degree. The Panel of Examiners was as follows:

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### ABSTRACT

Osteosarcoma (OS) is a common malignant bone tumour mainly occurs in children and young adults between the ages of 10 to 30 years. Although the advance of technology and improvement in chemotherapy treatment for OS has significantly improved the survival rate in recent years, its poor prognosis and unknown etiology continues to remain the major challenge in managing the disease progression. Therefore, the objectives of this study are to establish the serum protein profile of OS patients, to investigate the differential protein expression in pre and postchemotherapy and to identify the potential protein biomarkers relevant in OS cell migration. Serum and bone tissues biopsies were collected from healthy volunteers, pre- and post-chemotherapy OS patients. In this study, OS serum protein profile was established using a 4-plex iTRAQ analysis. Approximately 217 proteins with 104,214 spectra were acquired which includes low abundance proteins involved in biological processes and other protein classes. In addition, VCAM-1 was significantly altered (p<0.05) and shown to be highly expressed in post-chemotherapy OS serum. VCAMexpression was then verified using western and ICAM-3 blot 1 and immunohistochemistry staining to observe their roles in OS cell migrations. To date, this is the first differential protein expression study to use OS patients' serum at different stages for the protein profiling. The comprehensive data on the differentially expressed protein was successfully generated and the comparative study has revealed a significant amount of protein expressed has been significantly altered. This data could provide a new insight on the fundamental OS biological processes and further use as a potential biomarker for better OS prognosis.

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