## Redox homeostasis in the *Drosophila* testis stem cell self-renewal and maintenance

## For PhD Qualifying Examination (PQE) "Oral Component" – Open Seminar

Cancer stem cells (CSCs) are considered one of the primary causes for drug resistance and tumour recurrence. CSCs maintain low reactive oxygen species (ROS), and increased ROS by oxidizing agents show to selectively kill CSCs in leukaemia. Notably, CSCs share many properties with normal stem cells; they proliferate indefinitely, selfrenew and maintain low ROS. This suggests that elucidating the redox regulatory mechanisms in normal stem cells and characterizing the mechanisms utilized by ROS signalling to influence stem cell behaviour will yield valuable insights into how CSCs are regulated by redox homeostasis and may also lead to the development of therapeutic interventions targeting CSCs. The Drosophila testis serves as an ideal in vivo model to study stem cells behaviour. Our study showed that high ROS induced by oxidant treatment or by modulating Keap1/Nrf2 signalling activity cause a loss of germline stem cells (GSCs) by promoting a precocious differentiation of the cells. By contrast, low ROS levels induced by antioxidant treatment or modulation of Keap1/Nrf2 signalling led to an over-growth of GSCs. Our study of *Drosophila* testes demonstrates that ROS levels a vital role in the maintenance and self-renewal of stem cells.

Speaker:

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Anatomy Seminar Room, L2, MD10, Department of Anatomy, NUS.

DEPARTMENT OF ANATOMY