

UNIVERSITI TEKNOLOGI MARA

**EFFECTS OF ESTROUS CYCLE ON VASCULAR
CONTRACTILITY IN THE PRESENCE AND
ABSENCE OF ENDOTHELIUM**

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ABSTRACT

The aim of this study was to determine the possible influence of sex hormones on the contractile responses induced by phenylephrine and angiotensin II, in endothelium intact and denuded tissues. For this purpose, thoracic aorta segments from female WKY rats were used. Endothelium denuded aorta exhibited higher sensitivity to phenylephrine and angiotensin II contractility compared to endothelium intact tissues as tissue denudation remove endothelium which inhibit contractile agonist. States of estrous cycle modulated phenylephrine induced contractile responses more in endothelium intact tissues compared to angiotensin II and equally regulated responses for phenylephrine and angiotensin II in endothelium denuded tissues.

CHAPTER 1

INTRODUCTION

1.1 Background

The endothelium responds to physical and chemical stimuli via the synthesis and/or release of regulatory substances affecting vascular tone and growth, thrombosis and thrombolysis and platelet and leukocyte interactions with the endothelium (Sader & Celermajer, 2002). Vasoconstriction is a condition where there is a decrease in the diameter of the lumen of a blood vessel. Several hormones help regulate blood pressure and blood flow by altering cardiac output, changing systemic vascular resistance, or adjusting the total blood volume. Angiotensin II is a potent vasoconstrictor where it raises blood pressure by increasing systemic vascular resistance. Norepinephrine, epinephrine, antidiuretic hormone are hormones that can cause vasoconstriction. Several types of cells like white blood cells, platelets, endothelial cells and smooth muscle fibers release a wide variety of chemicals that alter blood vessel diameter. Vasoconstrictors release includes thromboxane A_2 , superoxide radicals, serotonin and endothelins (Tortora & Derrickson, 2006). α -adrenoreceptor agonist, phenylephrine is more dependent on intracellular calcium to induce contraction by myotropic action (Cadorette *et al.*, 2000).