EFFECT OF DEUTERIUM OXIDE ($D_2O$)
ON VASCULAR SMOOTH MUSCLE

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TINA MARIE McWILLIAM, B.Sc.
EFFECT OF DEUTERIUM OXIDE (D₂O) ON VASCULAR SMOOTH MUSCLE

BY

© TINA MARIE MCWILLIAM, B.Sc.

A thesis submitted to the School of Graduate Studies in partial fulfillment of the requirements for the degree of Master of Science

Faculty of Medicine
Memorial University of Newfoundland
August 1990

St. John's Newfoundland
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ISBN 0-315-65356-6
ABSTRACT

The influence of deuterium oxide (D₂O) on aortic vascular smooth muscle was investigated. The effect on receptor- and voltage-operated calcium channels was studied with the use of phenylephrine and potassium chloride respectively. D₂O depressed the phenylephrine induced contraction response in a dose dependent manner with 50% inhibition of maximum contraction observed with 60% D₂O. The effect of 60% D₂O on phenylephrine induced contraction was reversible and not dependent on an intact endothelium. Sixty percent D₂O also reduced potassium chloride induced contractions by 50% indicating an effect on voltage operated calcium channels. Studies with Bay K 8644, an L-type calcium channel activator, and calcium free buffer studies, confirm an effect on utilization of extracellular calcium sources. Inhibition of sucrose induced contraction and acetylcholine and isoprenaline induced relaxation with 60% D₂O suggest an effect on intracellular calcium mobilization or on the contractile elements themselves. D₂O also affected both the pD₂' for nifedipine and the pA₂ for prazosin. Collectively, these data indicate that D₂O acts at multiple sites.

In vivo studies indicate that chronic oral administration of 25% D₂O for 12 days reduces the blood pressure of spontaneously hypertensive (SHR) and normotensive (WKY) rats.
However, the lowering of blood pressure did not result in a corresponding change in vascular smooth muscle contractility induced with phenylephrine and potassium chloride. These results support previous evidence that a reduction in blood pressure after hypertension has developed does not reduce vascular reactivity changes. A further acute study revealed that desensitization results from chronic exposure to D₂O such that 60% D₂O produces a significant depression of contraction only in aortic rings obtained from SHR and WKY rats which had not been chronically treated with twenty-five percent D₂O.
KEY WORDS

Aorta
Calcium channels
Contraction
Deuterium oxide
Hypertension
In vitro
In vivo
Relaxation
Reversibility
Vascular smooth muscle
ACKNOWLEDGEMENTS

I would like to express deep appreciation to my supervisor, Dr. Andrew Rankin for his constant support and advice. His guidance and genuine interest in my research, along with his light hearted manner, made time spent in the laboratory very enjoyable. Thanks are extended to members of my supervisory committee, Dr. C.R. Triggle and Dr. J. Reynolds for their valuable assistance.

I am grateful for financial support provided by the Faculty of Medicine, the Graduate Students' Union, and by Dr. Rankin through a research grant from the Medical Research Council of Canada. I would like to express sincere gratitude to Atomic Energy of Canada Limited for their generous supply of deuterium oxide through Dr. A. Liepins.

Finally, I would like to thank Harold and my family for their love, support, and encouragement. Special thanks to my sister Elizabeth Ann for typing assistance.
# TABLE OF CONTENTS

## 1. INTRODUCTION

<table>
<thead>
<tr>
<th>Section</th>
<th>Title</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.1</td>
<td>Background information on deuterium oxide</td>
<td>1</td>
</tr>
<tr>
<td>1.2</td>
<td>Mechanisms of vascular smooth muscle contraction</td>
<td>7</td>
</tr>
<tr>
<td>1.3</td>
<td>Mechanisms of vascular smooth muscle relaxation</td>
<td>9</td>
</tr>
<tr>
<td>1.4</td>
<td>Competitive and noncompetitive inhibition</td>
<td>13</td>
</tr>
<tr>
<td>1.5</td>
<td>Hypertension</td>
<td>15</td>
</tr>
<tr>
<td>1.6</td>
<td>Aims of the study</td>
<td>18</td>
</tr>
</tbody>
</table>

## 2. MATERIALS AND METHODS

<table>
<thead>
<tr>
<th>Section</th>
<th>Title</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>2.1</td>
<td>Animals</td>
<td>20</td>
</tr>
<tr>
<td>2.2</td>
<td>Drugs and reagents</td>
<td>20</td>
</tr>
<tr>
<td>2.3</td>
<td>Preparation of aortic rings</td>
<td>22</td>
</tr>
<tr>
<td>2.3.1</td>
<td>Tissue isolation</td>
<td>22</td>
</tr>
<tr>
<td>2.3.2</td>
<td>Experimental apparatus</td>
<td>22</td>
</tr>
<tr>
<td>2.4</td>
<td>Experimental procedures</td>
<td>23</td>
</tr>
<tr>
<td>2.4.1</td>
<td>Phenylephrine dose response curves</td>
<td>23</td>
</tr>
<tr>
<td>2.4.2</td>
<td>Reversibility</td>
<td>23</td>
</tr>
<tr>
<td>2.4.3</td>
<td>Mechanical disruption of the endothelium</td>
<td>26</td>
</tr>
<tr>
<td>2.4.4</td>
<td>Potassium chloride dose response curves</td>
<td>26</td>
</tr>
<tr>
<td>2.4.5</td>
<td>Calcium dose response curves</td>
<td>27</td>
</tr>
<tr>
<td>2.4.6</td>
<td>Bay K 8644 dose response curves</td>
<td>27</td>
</tr>
<tr>
<td>2.4.7</td>
<td>Sucrose induced contraction</td>
<td>28</td>
</tr>
<tr>
<td>2.4.8</td>
<td>Isoprenaline induced relaxation</td>
<td>28</td>
</tr>
<tr>
<td>2.4.9</td>
<td>Acetylcholine induced relaxation</td>
<td>28</td>
</tr>
<tr>
<td>2.4.10</td>
<td>Determination of pD₂ for nifedipine</td>
<td>29</td>
</tr>
<tr>
<td>2.4.11</td>
<td>Determination of pA₂ for prazosin</td>
<td>29</td>
</tr>
<tr>
<td>2.4.12</td>
<td>Blood pressure measurements</td>
<td>30</td>
</tr>
</tbody>
</table>
3. RESULTS

3.1 Effect of D20 on aortic smooth muscle contraction
   3.1.1 Phenylephrine induced contraction
   3.1.2 Reversibility
   3.1.3 Role of the endothelium
   3.1.4 Potassium chloride induced contraction
   3.1.5 Calcium induced contraction
   3.1.6 Bay K 8644 induced contraction
   3.1.7 Sucrose induced contraction
   3.1.8 Synopsis

3.2 Effect of D20 on aortic smooth muscle relaxation
   3.2.1 Isoprenaline induced relaxation
   3.2.2 Acetylcholine induced relaxation
   3.2.3 Synopsis

3.3 Effect of D20 on the affinity of an antagonist for its receptor
   3.3.1 Noncompetitive inhibition
   3.3.2 Competitive inhibition
   3.3.3 Synopsis

3.4 Effect of D20 on aortic smooth muscle from WKY and SHR rats
   3.4.1 Blood pressure levels of WKY and SHR rats
   3.4.2 Chronic in vivo effect of twenty-five percent D20
      3.4.2.1 WKY/SHR: Potassium chloride induced contraction
      3.4.2.2 WKY/SHR: Phenylephrine induced contraction
   3.4.3 Acute in vitro effect of sixty percent D20
      3.4.3.1 WKY: Potassium chloride induced contraction
3.4.3.2 WKY: Phenylephrine induced contraction 88
3.4.3.3 SHR: Potassium chloride induced contraction 88
3.4.3.4 SHR: Phenylephrine induced contraction 91
3.4.4 Synopsis 91

4. DISCUSSION

4.1 Effect of D₂O on aortic smooth muscle contraction 97
4.2 Effect of D₂O on aortic smooth muscle relaxation 101
4.3 Effect of D₂O on the affinity of an antagonist for its receptor 102
4.4 Chronic and acute effects of D₂O on aortic smooth muscle from WKY and SHR rats 103
4.5 Summary and discussion 107

REFERENCES 111
LIST OF TABLES

Table 3-1: Effect of 60% D2O on the pD2' for nifedipine.
Table 3-2: Effect of 25% D2O consumption for 12 days on the systolic blood pressures of spontaneously hypertensive (SHR) and normotensive (WKY) rats.
LIST OF FIGURES

Figure 2-1: Experimental apparatus.

Figure 3-1: Inhibition of $1 \times 10^{-2} \text{M}$ phenylephrine induced aortic smooth muscle contraction with varying concentrations of $D_2O$.

Figure 3-2: The effect of varying concentrations of $D_2O$ on phenylephrine induced aortic smooth muscle contraction.

Figure 3-3: Reversal of the $D_2O$ inhibitory effect on phenylephrine induced aortic smooth muscle contraction.

Figure 3-4: The role of the endothelium in the effect of sixty percent $D_2O$ on phenylephrine induced aortic smooth muscle contraction.

Figure 3-5: The inhibitory effect of sixty percent $D_2O$ on potassium chloride induced aortic smooth muscle contraction.

Figure 3-6: The inhibitory effect of sixty percent $D_2O$ on the dose response curve of calcium in the presence of 60 mM potassium chloride.

Figure 3-7: The inhibitory effect of sixty percent $D_2O$ on Bay K 8644 induced aortic smooth muscle contraction.

Figure 3-8: The inhibitory effect of sixty percent $D_2O$ on 250 mM sucrose induced aortic smooth muscle contraction.

Figure 3-9: The inhibitory effect of sixty percent $D_2O$ on $1 \times 10^{-5} \text{M}$ isoprenaline induced aortic smooth muscle relaxation.

Figure 3-10: The effect of sixty percent $D_2O$ on the dose response curve constructed using $1 \times 10^{-10} \text{M}$ to $1 \times 10^{-5} \text{M}$ acetylcholine.

Figure 3-11: Four consecutive dose response curves constructed using 9.6 mM to 65.8 mM potassium chloride in the absence of $D_2O$.

Figure 3-12: Four consecutive dose response curves constructed using 9.6 mM to 65.8 mM potassium chloride in the presence of sixty percent $D_2O$. 
Figure 3-13: Noncompetitive inhibition of potassium chloride induced aortic smooth muscle contraction in the presence of nifedipine and in the absence of D$_2$O.

Figure 3-14: Noncompetitive inhibition of potassium chloride induced aortic smooth muscle contraction in the presence of nifedipine and sixty percent D$_2$O.

Figure 3-15: Four consecutive dose response curves constructed using 1x10$^{-12}$M to 1x10$^{-5}$M phenylephrine in the absence of D$_2$O.

Figure 3-16: Four consecutive dose response curves constructed using 1x10$^{-12}$M to 1x10$^{-5}$M phenylephrine in the presence of sixty percent D$_2$O.

Figure 3-17: Competitive inhibition of phenylephrine induced aortic smooth muscle contraction in the presence of prazosin and in the absence of D$_2$O.

Figure 3-18: Competitive inhibition of phenylephrine induced aortic smooth muscle contraction in the presence of prazosin and sixty percent D$_2$O.

Figure 3-19: Chronic in vivo effect of twenty-five percent D$_2$O on potassium chloride induced contraction of aortic smooth muscle from WKY and SHR rats.

Figure 3-20: Chronic in vivo effect of twenty-five percent D$_2$O on phenylephrine induced contraction of aortic smooth muscle from WKY and SHR rats.

Figure 3-21: Acute in vitro effect of sixty percent D$_2$O on potassium chloride induced contraction of aortic smooth muscle from WKY rats.

Figure 3-22: Acute in vitro effect of sixty percent D$_2$O on phenylephrine induced contraction of aortic smooth muscle from WKY rats.

Figure 3-23: Acute in vitro effect of sixty percent D$_2$O on potassium chloride induced contraction of aortic smooth muscle from SHR rats.

Figure 3-24: Acute in vitro effect of sixty percent D$_2$O on phenylephrine induced contraction of aortic smooth muscle from SHR rats.
# ABBREVIATIONS

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>ATP</td>
<td>Adenosine triphosphate</td>
</tr>
<tr>
<td>Ca&lt;sup&gt;2+&lt;/sup&gt;</td>
<td>Calcium ion</td>
</tr>
<tr>
<td>cAMP</td>
<td>Cyclic adenosine monophosphate</td>
</tr>
<tr>
<td>CaCl&lt;sub&gt;2&lt;/sub&gt;</td>
<td>Calcium chloride</td>
</tr>
<tr>
<td>cGMP</td>
<td>Cyclic guanosine monophosphate</td>
</tr>
<tr>
<td>D&lt;sub&gt;2&lt;/sub&gt;O</td>
<td>Deuterium oxide</td>
</tr>
<tr>
<td>BDRF</td>
<td>Endothelium derived relaxing factor</td>
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<tr>
<td>EDTA</td>
<td>Ethylenediaminetetraacetic acid</td>
</tr>
<tr>
<td>G-protein</td>
<td>Guanosine triphosphate binding protein</td>
</tr>
<tr>
<td>Hg</td>
<td>Mercury</td>
</tr>
<tr>
<td>IP&lt;sub&gt;3&lt;/sub&gt;</td>
<td>Inositol trisphosphate</td>
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<tr>
<td>K&lt;sup&gt;+&lt;/sup&gt;</td>
<td>Potassium ion</td>
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<td>KCl</td>
<td>Potassium chloride</td>
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<tr>
<td>KH&lt;sub&gt;2&lt;/sub&gt;PO&lt;sub&gt;4&lt;/sub&gt;</td>
<td>Potassium dihydrogen phosphate</td>
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<td>M</td>
<td>Molar concentration</td>
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<tr>
<td>mm</td>
<td>Millimeters</td>
</tr>
<tr>
<td>mM</td>
<td>Millimolar concentration</td>
</tr>
<tr>
<td>MgSO&lt;sub&gt;4&lt;/sub&gt;</td>
<td>Magnesium sulphate</td>
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<tr>
<td>Na&lt;sup&gt;+&lt;/sup&gt;</td>
<td>Sodium ion</td>
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<td>Sodium chloride</td>
</tr>
<tr>
<td>NaHCO&lt;sub&gt;3&lt;/sub&gt;</td>
<td>Sodium bicarbonate</td>
</tr>
<tr>
<td>S.E.M.</td>
<td>Standard error of the mean</td>
</tr>
<tr>
<td>SHR</td>
<td>Spontaneously hypertensive rats</td>
</tr>
<tr>
<td>WKY</td>
<td>Wistar-Kyoto rats</td>
</tr>
</tbody>
</table>
CHAPTER 1

INTRODUCTION

1.1 Background information on deuterium oxide

Deuterium, the heavy stable isotope of hydrogen, was discovered by Urey et al. in 1932 and deuterium oxide (D₂O) was first made available by Urey in 1933. Since the addition of the extra proton to hydrogen practically doubles the mass, there are great differences between the original element and the isotope. The changes include an increase in the melting and boiling point by 3.82°C and 1.42°C respectively, a 25% increase in viscosity, lowering of vapor pressure and an increase in the specific gravity to 1.1074 (Barbour, 1937). There are also solubility changes such that carbon dioxide and oxygen are both about 10 percent less soluble in D₂O than in H₂O (Katz, 1960). Deuterium oxide is found as a natural component of ordinary water at a concentration of one part in 5000 or 0.02 percent (Barbour, 1937).

It is interesting to note that the high cost of D₂O was an obstacle for prolonged in vivo studies using D₂O during the thirty years following its discovery (Thomson, 1960) and even today the high cost is a limiting factor. Since there are large differences between hydrogen and deuterium, and hydrogen is a major constituent of mammalian tissues, early studies dealt with determining the effect of D₂O on mammals. Barbour
(1937) was one of the first researchers to conduct a study of this kind and one of his fundamental conclusions was that deuterium is a depressant of biological activity and that its effects can be reversible, or not, depending upon the concentration. From this study it was also determined that mice could not survive replacement of more than one third of body water by $D_2O$; metabolic rate is increased in the early stages of $D_2O$ intoxication; and $D_2O$ has a sympathomimetic action. A similar study was conducted by Thomson (1960) who found that rats died when about one third of their body water was replaced by $D_2O$. The effects of $D_2O$ included impairment of kidney function, anaemia, disturbed carbohydrate metabolism, central nervous system disturbances, and wide effects on enzyme systems. It was concluded that the toxic action of $D_2O$ was due to the additive effect of the many changes in rates of enzymatic reactions.

Chemical and biological studies conducted by Katz (1960) reinforced the fact that $D_2O$ inhibits many enzymatic reactions. He determined that although the binding rate constants for $H^+$ and $D^+$ ions are similar, the rate constant for the unbinding of $D^+$ is much slower, demonstrating the increased stability of deuterium bonds and accounting for isotopic effects. Experimental rate constants supporting isotopic modifications due to $H^+$ and $D^+$ exchange have most recently been calculated by Michael et al. (1990).
Goodall (1958) suggested that the rate limiting process for vascular contraction may be proton transfer. He showed a three fold reduction in the development of isometric tension in frog sartorius muscle without a reduction in the amplitude of the action potential and concluded that D₂O acted on the contractile proteins with the involvement of possibly three protons. This finding sparked investigations into the effects of D₂O on the mechanical response of various muscle types.

Kaminer (1960) studied the effect of D₂O on the isolated atria of the frog and found that concentrations of heavy water between 99.8 and 25 percent decreased the force of contraction and the rate of spontaneous beating of the isolated atria. This data supports the result found by Barnes and Warren (1935) who were the first to show that 20% D₂O slows the frequency of pulsation of the excised heart of the frog. In order to elucidate the site of action of D₂O, Kaminer (1960) performed subsequent experiments on glycerol extracted psoas muscle fibres (where contractile proteins are isolated) and found that D₂O does not affect adenosine triphosphate (ATP) induced shortening of the muscle fibres. He then concluded that it is the excitation-contraction coupling which is affected by D₂O and not the contractile proteins.

Further studies on the inhibition of enzymes by Svensmark (1961) revealed that D₂O inhibited adenosine triphosphatase, an enzyme associated with muscular activity, by 50%. He then
investigated whether the inhibition was reflected in the mechanical response of skeletal muscle. Using frog semitendinous muscle he found that the amplitude of the action potential was unaffected by D₂O but there was a reduction of the frequency at which a complete tetanus could be maintained, suggesting an effect on the primary excitatory processes. There was also a reduction in twitch force and the rate of force development in D₂O which he suggested could be caused by a direct effect on the contractile elements. From this study it remained undecided whether the reduction in the force of contraction was due to a D₂O effect on the contractile elements or on excitation-contraction coupling. Also, due to inadequate controls, Svensmark argued that no definite conclusions could be drawn from the work of Goodall (1958) or Kaminer (1960).

The conflicting conclusions drawn from these studies led to increased interest in determining the site of action of D₂O. Kaminer and Kimura (1972) supported their previous conclusion by determining that the luminescent response of aequorin (a calcium sensitive luminescent protein) which accompanies contraction was abolished when H₂O was replaced by D₂O. However, it was again argued that suitable controls were lacking since the possible effects of D₂O on calcium sensitivity and kinetics of the aequorin reaction were unknown (Allen et al., 1984). Studies conducted by Allen et al. (1984) involved determining the influence of the substitution
of D₂O for H₂O on calcium transients and on contraction in both intact single skeletal muscle fibres injected with aequorin and in mechanically skinned fibres from frogs. These studies revealed that the calcium sensitivity of the myofibrils is reduced by D₂O substitution but that at saturating calcium concentrations the myofilaments generate more tension in D₂O than in H₂O. Results also indicated that calcium release from the sarcoplasmic reticulum was reduced in D₂O and it was concluded that this reduction in calcium release was a major factor in reducing twitch tension but reduced myofibrillar calcium sensitivity also played a role. Also, the rate of change of aequorin light emission brought about by D₂O substitution represented a 3 to 5 fold reduction in intracellular calcium concentration which was dependent on the concentration of D₂O and proportional to the fraction of H₂O replaced. This change in light emission, after solvent exchange, occurred too fast to be explained by exchange of intracellular water so it was postulated that the site whereby D₂O affected calcium release must be accessible to the extracellular space. It was concluded that there was time for D₂O to diffuse in or out of the T-tubules in time to produce the effects on calcium release and therefore it was proposed that the site of action of D₂O may be on excitation-contraction coupling in the T-tubules. Earlier studies (Sandow et al., 1976 and Yagi and Endo, 1976) also support D₂O effects on excitation-contraction coupling in skeletal muscle. Other evidence supporting an effect of D₂O on calcium
transients is found from the fact that D₂O reduces L-type calcium channel conductance in isolated myocytes (Prod'hom et al., 1987).

Despite wide interest in the study of the effects of D₂O on skeletal muscle, few experiments have been carried out using smooth muscle. The effect of D₂O on aortic vascular smooth muscle from spontaneously hypertensive rats (Vasdev et al., 1990a) and Dahl salt-sensitive hypertensive rats (Vasdev et al., 1990b) has been investigated. It was found that D₂O inhibited in vitro calcium uptake in aortae from these rats. Since D₂O has been demonstrated to reduce calcium uptake in vascular smooth muscle it might be expected that D₂O will also inhibit contraction of vascular smooth muscle. The purpose of the present study is to determine if this is so and to try to resolve the previously postulated mechanisms of action of D₂O on the contraction process.

The thoracic aorta was chosen to study the effect of deuterium oxide on vascular smooth muscle. The rat aorta was the vessel of choice since it is devoid of adrenergic nerves (Patil et al., 1972) and Fleisch (1974) reports that the rat aortic preparation is ideal when one wishes to investigate the effects of drugs on vascular smooth muscle devoid of nervous innervation. This condition is ideal since drugs not only act at receptors, producing an effect of their own, but may also act as nerves innervating the tissue and eliciting the release
of endogenous agonists resulting in additional tissue responses.

1.2 Mechanisms of vascular smooth muscle contraction

The most important step in the contraction or relaxation of vascular smooth muscle involves calcium mobilization (Somlyo and Somlyo, 1968). An increase in the intracellular calcium concentration at the level of the contractile proteins results in contraction of the muscle, whereas removal of calcium results in relaxation (Fleisch, 1974). Vascular smooth muscle cells may be activated either by agonists which activate receptor-operated calcium channels or by membrane depolarization which activates the opening of voltage-operated calcium channels.

There are three types of voltage dependent calcium channels. These are the T, L, and N channels which refer to transient (fast), long lasting (slow), and neuronal channels, respectively (Nowycky et al., 1985). However, only T- and L-type channels are found in vascular smooth muscle (Zelis and Moore, 1989).

In addition to receptor- and voltage-operated calcium channels there is also a leak channel which is the calcium entry pathway which allows a continuous entry of calcium into the cell that is not increased on stimulation by depolarization or agonists (Khalil et al., 1987).
Agonist activated increases in intracellular calcium concentration are mediated by a transducing guanosine triphosphate binding protein (G-Protein), which in turn activates enzymes, such as phospholipase C, to generate second messengers. Two of these second messengers are inositol triphosphate (IP$_3$) and Ca$^{2+}$ which induce the sarcoplasmic reticulum to increase the release of calcium, thereby increasing the intracellular concentration of Ca$^{2+}$ available for binding to and activation of calmodulin (Williamson, 1986). Calmodulin is a ubiquitous protein which is the major receptor of Ca$^{2+}$ and appears to mediate most of the ion's activities as a second messenger (Wang and Waisman, 1979). The calcium-calmodulin complex then activates myosin light chain kinase. Myosin light chain kinase is the enzyme which catalyses the phosphorylation of myosin; a process important for the development of contraction in vascular smooth muscle (Adelstein et al., 1981). Activation of the kinase leads to rapid cycling of the cross-bridges between actin and myosin and this interaction of the contractile proteins results in contraction of vascular smooth muscle (Khalil et al., 1987).

Depolarization induced increases in intracellular calcium concentration occur by direct entry of calcium via the T- and L-type calcium channels (Zelis and Moore, 1989).

In this study phenylephrine, an alpha-adrenergic agonist, was chosen to study the D$_2$O effect on contraction induced via
receptor mediated increases in intracellular calcium. Although phenylephrine is not completely selective for receptor-operated calcium channels (Godfraind et al., 1986) it does act mainly on this type of channel to produce contraction of aortic vascular smooth muscle (Khalil et al., 1987). Potassium chloride acts mainly on the voltage-operated calcium channel (Meisheiri et al., 1981) and was chosen to study the effect of D$_2$O on this channel type. The effects on this type of channel were also investigated using Bay K 8644 which enhances calcium influx via the voltage sensitive calcium channel under partial depolarizing conditions (Schramm et al., 1983). To further ensure no effect from receptor-operated calcium channels, they were blocked by the alpha-receptor antagonist phentolamine. Similarly, calcium free buffer studies were performed in an attempt to clarify the effects of D$_2$O on contraction induced by the influx of extracellular calcium.

Finally, the effect on sucrose induced contraction was studied to determine the effect of D$_2$O on intracellular calcium mobilization since it occurs independently of the level of membrane potential and in the absence of external calcium sources (Andersson et al., 1974).

1.3 Mechanisms of vascular smooth muscle relaxation

The two main proposed mechanisms of vascular smooth muscle relaxation involve an increase in either cyclic
adenosine monophosphate (cAMP) or cyclic guanosine monophosphate (cGMP) levels. In this study a representative was chosen from each group: isoprenaline, which is a beta-agonist and acts via an increase in cAMP levels; and acetylcholine, which results in relaxation via an increase in cGMP levels.

Various different mechanisms are thought to result in vascular smooth muscle relaxation through an increase in cAMP; including beta-adrenergic stimulation, inhibition of phosphodiesterase, and regulation of myosin light chain kinase.

Although various mechanisms have been proposed for beta-adrenergic receptor mediated relaxation of vascular smooth muscle, all theories suggest the involvement of cAMP as the second messenger (Mueller and van Breemen, 1979). Relaxation of vascular smooth muscle in response to adrenergic stimulation is mediated by $\beta_1$, $\beta_2$, or both $\beta_1$ and $\beta_2$ adrenoceptors (Lucchesi, 1989). The beta-adrenergic receptor is coupled to adenylate cyclase by a stimulatory G-protein. Activation of adenylate cyclase converts ATP to cAMP and the cAMP in turn activates a protein kinase which catalyses a protein phosphorylation and leads to a reduction in intracellular free calcium (Mueller and van Breemen, 1979). To date, it has not been determined how the intracellular calcium concentration is reduced. The proposed mechanisms by
which cAMP may decrease the calcium ion concentration available to the contractile proteins include: increased Ca\(^{2+}\) sequestration into intracellular storage sites such as the sarcoplasmic reticulum (Mueller and van Breemen, 1979); inhibition of calcium influx (Meisher and van Breemen, 1982); and increased calcium extrusion via increased activity of the Na\(^+\)–K\(^+\) pump (Scheid and Fay, 1984). At present it is thought that calcium sequestration by increased Ca\(^{2+}\) uptake into the sarcoplasmic reticulum probably plays the primary role in decreasing free intracellular calcium (Bülbbring and Tomita, 1987).

There is also evidence to support a role for increases in cAMP via inhibition of phosphodiesterase. Kaufman et al. (1987) showed that a series of dihydropyridazones had vascular smooth muscle relaxant potencies which correlated directly with their ability to inhibit an isozyme of cyclic nucleotide phosphodiesterase located in the sarcoplasmic reticulum of cardiac muscle. They suggested that the vascular relaxation produced by these agents, which produced significant increases in cAMP, was related to their ability to inhibit a vascular enzyme similar or identical to this isozyme.

It is also suggested that increased cAMP levels may cause relaxation by the regulation of myosin light chain kinase. The activity of myosin light chain kinase is regulated by cAMP-dependent protein kinase which diphosphorylates the
enzyme and weakens its ability to bind to and be activated by calmodulin. Evidence supporting diphosphorylation of myosin light chain kinase has been found by Vallet et al. (1981) in vascular smooth muscle.

In recent years there is increasing evidence in support of a role for cGMP as a mediator of vascular smooth muscle relaxation. Cyclic GMP is thought to mediate vasorelaxation by endothelium derived relaxing factor (EDRF), atrial natriuretic factor, and the nitrovasodilators (Khalil et al., 1987). The increased cGMP levels are attained by activation of guanylate cyclase and the increased levels of cGMP lead to decreased intracellular calcium levels.

In 1980, Furchgott and Zawadzki discovered that the relaxation by acetylcholine of precontracted isolated vascular preparations is dependent on the presence of intact endothelial cells. It has been demonstrated that EDRF directly activates the soluble isozyme form of guanylate cyclase (Fosterman et al., 1986) and it is thought that EDRF is a reactive oxidized material, analogous to nitric oxide, that can give rise to a free radical which activates guanylate cyclase (Murad, 1986).

Unlike EDRF, atrial natriuretic factor selectively activates the particulate form of guanylate cyclase in vascular smooth muscle (Winquist et al., 1984) and has no
effect on the soluble form of the enzyme. The endothelium
independent relaxant effect of atrial natriuretic factor is
also thought to be mediated via increased tissue levels of
cGMP (Winguist et al., 1984). However, Nakatsu and Diamond
(1989) believe that cGMP turnover in the tissue may be more
important than cGMP concentration.

Nitrovasodilators are also thought to mediate smooth
muscle relaxation by activating the particulate form of
guanylate cyclase and increasing intracellular concentrations
of cGMP (Waldman and Murad, 1987). The exact mechanism of
enzyme activation remains unknown but it has been suggested
that these agents are all capable of generating nitric oxide
and that this free radical is the activator of guanylate
cyclase.

1.4 Competitive and noncompetitive inhibition

This study involved determining the effect of D2O on the
pA2 value of the alpha-adrenoceptor antagonist prazosin which
competitively blocks the response to the alpha-adrenoceptor
agonist phenylephrine. Schild (1947) defined the pA2 value as
the negative logarithm of the molar concentration of a
competitive antagonist which reduces the effect of a double
concentration of agonist to that of a single one. The pA2
value should also be an accurate indication of the affinity of
an antagonist for its receptor (Muramatsu et al., 1990).
An agonist is a drug which when coupled with the receptor in an agonist-receptor complex will elicit a response. An antagonist on the other hand, may form an antagonist-receptor complex but it does not produce a cellular response. A competitive antagonist is one which will shift the dose response curve of an agonist to the right in a manner proportional to the antagonist concentration. However, the maximal response remains the same since the effect of the antagonist can be overcome by raising the agonist concentration. A pA₂ value may be determined by using a fixed concentration of antagonist along with graded concentrations of an agonist which acts at the same receptor.

Along with determining the D₂O effect on competitive inhibition the effect on noncompetitive inhibition was also studied. Noncompetitive inhibition is a type of antagonism which is not surmountable and therefore cannot be overcome by increasing the concentration of agonist or depolarizing agent. The noncompetitive antagonist used was nifedipine which is referred to as a calcium antagonist and blocks calcium entry via L-type calcium channels only (Zelis and Moore, 1989). Potassium chloride was used as the depolarizing agent.

Godfraind et al. (1986) define a calcium antagonist as a drug that alters the cellular function of calcium by inhibiting its entry or release or by interfering with its intracellular action. Although Church and Zsotér (1980)
suggest an intracellular action for nifedipine, Godfraind (1983) argues that the action of nifedipine is related to blockade of calcium entry.

To determine the effect of D$_2$O on the action of nifedipine, a $pD_z$' value was determined. The $pD_z$' is defined as the negative logarithm of the molar concentration of a noncompetitive antagonist which reduces the maximal effect of an agonist to 50\% of its initial value (Ariens and van Rossum, 1957).

1.5 Hypertension

In order to gain knowledge in understanding the development of hypertension in man and to find methods for its control and treatment, studies have been carried out on various animal models, the most popular species of which is the rat. Hypertension is induced in animal models by a number of methods including deoxycorticosterone acetate administration and renal artery constriction with or without removal of the contralateral kidney (Field et al., 1972) but a number of genetic models of hypertension have also been developed. The most commonly used strains include the spontaneously hypertensive rat (SHR) which was developed by brother-sister inbreeding and the Dahl salt-sensitive rat which develops hypertension due to a sensitivity to increases in dietary sodium chloride (Triggle, 1989). One problem associated with using these genetic models of hypertension is
that there is no single adequate control for the models (Clineschmidt et al., 1970) and often the results depend on the type of normotensive control which is used for the study. Clineschmidt et al. (1970) suggested the use of more than one strain of rat as a control since they found that although aortae from SHR did not show a reduced contractile response to noradrenaline in comparison to National Institute of Health Wistar rats there was a depression of contraction in comparison to Carworth Farm Wistar rats.

A further problem associated with the use of genetically hypertensive rats arises from the fact that different strains show varying results. For example, Massingham and Shevde (1971) showed that aortae from the New Zealand strain of genetically hypertensive rat exhibited reduced reactivity to norepinephrine while Clineschmidt et al. (1970) showed that the sensitivity of SHR rats was the same as controls. Furthermore, the results obtained for any particular strain vary depending on whether the work is performed in vivo or in vitro. However, generally in vitro studies show a decrease in the sensitivity of tissues from SHR rats to contractile agents in comparison to normotensive controls (Triggle, 1989).

Although many problems exist with the use of these animal models information has been gained contributing to the knowledge of the disease process of hypertension in man and to the development of drugs for treatment.
It is widely accepted that the genetically hypertensive rat demonstrates vascular responses which are different from those of normotensive controls. Most research is geared toward investigating what determines these vascular reactivity changes and if they are a cause or a consequence of the hypertension. In this study an assessment was made as to the effect of D$_2$O on the blood pressure of spontaneously hypertensive rats and Wistar Kyoto normotensive rats (WKY), and on the contractile response to both phenylephrine and potassium chloride.

Previous studies by Vasdev et al. (1990a) using SHR rats show that D$_2$O reduces the blood pressure and high in vitro calcium uptake induced by phenylephrine via receptor-operated calcium channels and by potassium chloride via voltage-operated channels in aortae. Similar studies using Dahl salt-sensitive rats showed that D$_2$O normalizes both elevated blood pressure and in vitro aortic calcium uptake (Vasdev et al., 1990b). Since D$_2$O has been shown to normalize calcium uptake in SHR rats and reduce blood pressure it may be expected that the phenylephrine and potassium chloride induced contractions of SHR aortae from rats treated with D$_2$O will not differ from those of normotensive controls. In considering this hypothesis, however, it is worth noting that vascular reactivity changes using other hypotensive agents were not reversed once hypertension had developed (Triggle, 1989). Since this study involves the use of aortae from rats which
have already developed hypertension, this latter observation should be taken into consideration.

1.6 Aims of the study

1. To determine if D$_2$O has an effect on vascular smooth muscle contraction induced by phenylephrine which stimulates calcium entry via the receptor-operated calcium channel.

2. If there is a D$_2$O effect on phenylephrine induced vascular smooth muscle contraction, to determine if the effect is reversible.

3. Since vascular endothelium releases EDRF when stimulated by a range of pharmacological agents (Furchgott et al., 1983), to investigate whether the endothelium plays a role in the D$_2$O mediated effect on vascular smooth muscle contraction.

4. To determine if D$_2$O has an effect on vascular smooth muscle contraction induced by potassium chloride which stimulates calcium entry via voltage-operated calcium channels.

5. To investigate the effect of D$_2$O on vascular smooth muscle contraction induced by Bay K 8644 which enhances calcium influx via voltage-operated calcium channels under partial depolarizing conditions.

6. To determine the effect of D$_2$O on vascular smooth muscle contraction induced by extracellular calcium influx via the voltage-operated calcium channel after depletion of intracellular calcium.
7. To assess the effect of D$_2$O on sucrose induced vascular smooth muscle contraction which occurs independently of membrane potential and in the absence of extracellular calcium.

8. To determine whether D$_2$O will have a selective effect on vascular smooth muscle relaxation induced by either isoprenaline or acetylcholine which act by increasing intracellular concentrations of cAMP and cGMP, respectively.

9. To investigate whether D$_2$O will have a selective effect on either noncompetitive inhibition of potassium chloride induced contraction by nifedipine or competitive inhibition of phenylephrine induced contraction by prazosin.

10. To examine whether D$_2$O produces a hypotensive effect in SHR and WKY rats and, if so, to determine whether there will be corresponding vascular reactivity changes.

11. To attempt to clarify the previously postulated mechanisms of action of D$_2$O.
CHAPTER 2
MATERIALS AND METHODS

2.1 Animals

For experimental procedures 2.4.1 to 2.4.11, male Sprague-Dawley rats weighing between 100-150 grams were purchased from Canadian Hybrid Farms, Nova Scotia, Canada. WKY and SHR rats (100-150 grams), for experimental procedures 2.4.12 to 2.4.14, were purchased from Charles River Inc., Montreal, Canada. At the time of sacrifice the rats weighed 300-400 grams.

The rats were maintained under standard light and temperature conditions (twelve hour light/dark period, 20-22 °C, 40-60% humidity). Housing was provided by rectangular shoe box cages, made of polycarbonate plastic, with a detachable metal rod lid (three rats per cage). The cages contained Beta-Chip wood chip bedding purchased from Charles River Inc. Diet consisted of free access to Purina Rat Chow #T-5012 (Ralston Purina Company) and tap water unless otherwise specified in the experimental procedures.

2.2 Drugs and reagents

The list of drugs and chemicals used throughout the experimental procedures is:
<table>
<thead>
<tr>
<th>Chemical Name</th>
<th>Supplier</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acetylcholine chloride</td>
<td>Sigma Chemical Company, MO, USA</td>
</tr>
<tr>
<td>Bay K 8644</td>
<td>Research Biochemicals Inc., MA, USA</td>
</tr>
<tr>
<td>Calcium chloride</td>
<td>Sigma Chemical Company, MO, USA</td>
</tr>
<tr>
<td>Deuterium oxide</td>
<td>Sigma Chemical Company, MO, USA</td>
</tr>
<tr>
<td>Ethylenediaminetetraacetic</td>
<td>Sigma Chemical Company, MO, USA</td>
</tr>
<tr>
<td>acid</td>
<td>DIFCO Laboratories, MI, USA</td>
</tr>
<tr>
<td>Isoprenaline</td>
<td>Sigma Chemical Company, MO, USA</td>
</tr>
<tr>
<td>Magnesium sulphate</td>
<td>Sigma Chemical Company, MO, USA</td>
</tr>
<tr>
<td>Nifedipine</td>
<td>BDH Chemicals, Ont., Can.</td>
</tr>
<tr>
<td>Phentolamine</td>
<td>Sigma Chemical Company, MO, USA</td>
</tr>
<tr>
<td>Phenylephrine</td>
<td>Sigma Chemical Company, MO, USA</td>
</tr>
</tbody>
</table>
Prazosin
Sodium bicarbonate
Sodium chloride
Sucrose

Sigma Chemical Company, MO, USA
Sigma Chemical Company, MO, USA
Sigma Chemical Company, MO, USA

2.3 Preparation of aortic rings

2.3.1 Tissue isolation

Rats were killed by a blow to the head and exsanguinated. The chest cavity was then surgically opened and the thoracic aorta was removed and placed in Krebs solution of the following composition (in mM): NaCl 118, KCl 4.7, CaCl₂ 2.5, KH₂PO₄ 1.2, MgSO₄ 1.2, NaHCO₃ 12.5, dextrose 11.1, and EDTA 0.01. These solutes were diluted in either deionized water or deuterium oxide, as specified in the experimental procedures, and aerated with 95% oxygen and 5% carbon dioxide. The aorta was then cleaned of all fat, blood and connective tissue and cut into sectional rings of approximately three mm in length.

2.3.2 Experimental apparatus

Aortic rings were stationed on metal hooks (made from insect pins) and attached to force transducers, by 5-0
silk thread, at one end while the other end was held stationary. Pairs of rings were lowered into 25 mL baths heated at 38°C, containing aerated Krebs solution. Tension measured by the force transducers was recorded on a Beckman R611 polygraph. A diagramatic representation of the experimental apparatus may be seen in Figure 2-1. One gram of tension was applied to the rings and they were equilibrated in the Krebs solution for a time specified in the experimental procedures.

2.4 Experimental procedures

2.4.1 Phenylephrine dose response curves

Aortic rings were allowed to equilibrate for two hours in Krebs solution (made up in 0, 20, 40, 60, or 80% D₂O by volume), the solution being changed every thirty minutes. A dose response curve was then constructed by adding consecutive doses of phenylephrine (1x10⁻¹²M to 1x10⁻⁵M) to the bath containing the aortic rings once contraction to the previous concentration had reached a plateau. Following the contraction with phenylephrine, 1x10⁻⁵M acetylcholine was added and a relaxation noted to confirm that the endothelium of the tissue was intact (also performed in protocols 2.4.2-2.4.4).

2.4.2 Reversibility

Rings were allowed to equilibrate in sixty percent D₂O Krebs for one hour; Krebs with zero percent D₂O was used
Figure 2-1: Experimental apparatus.
as a control. A phenylephrine dose response curve was then constructed using $1 \times 10^{-12} \text{M}$ to $1 \times 10^{-5} \text{M}$ phenylephrine. Rings which had been equilibrated in sixty percent D$_2$O Krebs were then re-equilibrated in regular Krebs for two hours. The second equilibration time was double that of the first because although the binding rate constants for H$^+$ and D$^+$ ions are similar, the rate constant for the unbinding of D$^+$ is slower by a factor of about 2.5 (Prod'hom et al., 1987). A second phenylephrine contraction dose response curve was then constructed.

2.4.3 Mechanical disruption of the endothelium

After the aorta had been removed, it was cut into two sections. One section had the endothelium mechanically disrupted by gently scraping the inside of the aorta with a tube of soft plastic. Three mm rings were then cut and a ring with an intact endothelium was equilibrated in the same bath as a ring with a disrupted endothelium. Tissues were set up in organ baths containing sixty percent D$_2$O Krebs or regular Krebs as the control. The tissues were allowed to equilibrate for two hours and then a phenylephrine dose response curve was performed. Acetylcholine induced relaxation was demonstrated to be absent to confirm that the endothelium scraping was effective.

2.4.4 Potassium chloride dose response curves

Rings were allowed to equilibrate in sixty percent
D2O for two hours (regular Krebs for controls). Krebs solutions were then made up with varying concentrations of potassium chloride from 9.6 mM to 65.8 mM. The concentration of sodium chloride added was also simultaneously changed so as to maintain an isotonic solution. A dose response curve was then constructed by changing the bath solutions and substituting Krebs solution containing increasing concentrations of potassium chloride.

2.4.5 Calcium dose response curves

Aortic rings were permitted to equilibrate in either sixty percent D2O Krebs or regular Krebs for two hours. A 1x10^-6M phenylephrine induced contraction was then measured to confirm the presence of the D2O inhibitory effect. The rings were then washed in calcium free Krebs solution, containing 60 mM KCl, at fifteen minute intervals until no contraction response to KCl could be elicited (at least one hour); the calcium free Krebs was also diluted with either sixty percent D2O or H2O. A calcium dose response curve was then constructed by adding increasing concentrations of CaCl2 from 0.5 mM to 5 mM.

2.4.6 Bay K 8644 dose response curves

Aortic rings were allowed to equilibrate in sixty percent D2O Krebs or regular Krebs for two hours. The Krebs solution was then substituted for Krebs solution containing 10 mM KCl; 1x10^-6M phentolamine was added to block receptor
operated calcium channels. A Bay K 8644 dose response curve was then constructed using 1x10^{-6}M to 1x10^{-5}M Bay K 8644.

2.4.7 Sucrose induced contraction

Aortic rings were permitted to equilibrate in either sixty percent D_2O Krebs or regular Krebs for two hours. A sucrose contraction response was then performed using 250 mM sucrose.

2.4.8 Isoprenaline induced relaxation

Rings were permitted to equilibrate in either sixty percent D_2O Krebs or regular Krebs for one hour. A phenylephrine induced contraction response was then measured with 1x10^{-6}M phenylephrine to confirm the presence of the D_2O inhibitory effect. After washing for one further hour the tissues equilibrated in regular Krebs were exposed to 1x10^{-6}M phentolamine to bring the contraction levels to the same level as tissues incubated in sixty percent D_2O Krebs. A second 1x10^{-6}M phenylephrine induced contraction was then performed and the precontracted tissues were relaxed with 1x10^{-4}M isoprenaline.

2.4.9 Acetylcholine induced relaxation

Following the final phenylephrine control dose response curves in the absence of D_2O and in the presence of sixty percent D_2O in experimental procedure 2.4.11 an acetylcholine dose response curve was constructed using
2.4.10 Determination of pD₂' for nifedipine

Tissues were equilibrated in either sixty percent D₂O Krebs or regular Krebs for two hours. A potassium chloride dose response curve was then constructed according to the method stated in experimental procedure 2.4.4. Subsequent response curves were constructed on the same tissues pre-incubated with three different concentrations of nifedipine: 5x10⁻⁹M, 1x10⁻⁸M, and 1x10⁻⁷M. Controls were performed simultaneously in which four dose response curves to potassium chloride were constructed without the addition of nifedipine. The pD₂' was calculated according to the following equation derived by Bickerton (1963):

\[
pD₂' = pDx' + \log [Eam / Eabm - 1]
\]

where: pDx' = the negative logarithm of the molar concentration of nifedipine

Eam = the maximal contraction height in the absence of nifedipine

Eabm = the maximal contraction height in the presence of nifedipine

2.4.11 Determination of pA₂ for prazosin

Tissues were incubated in either sixty percent D₂O Krebs or regular Krebs for two hours. A phenylephrine dose
response curve was then constructed using $1 \times 10^{-12} \text{M}$ to $1 \times 10^{-5} \text{M}$ phenylephrine. Subsequent response curves were constructed on the same tissues pre-incubated with three different concentrations of prazosin: $1 \times 10^{-9} \text{M}$, $1 \times 10^{-8} \text{M}$, and $1 \times 10^{-7} \text{M}$. Controls were performed simultaneously in which four dose response curves to phenylephrine were constructed without the addition of prazosin. The $pA_2$ was calculated according to the Schild Plot method as described by Schild (1947). The Schild Plot is a plot of $\log (X'/X)-1$ versus $-\log (A)$ where:

$$X'/X = \text{the agonist dose ratio}$$
$$A = \text{the antagonist concentration}$$

For a competitive antagonist, the line has a slope of $-1$ and the intercept on the abscissa corresponds to $pA_2$.

2.4.12 Blood pressure measurements

Systolic blood pressure was measured by tail plethysmography according to the method described by Byrom (1969); signals were amplified using a model 5A amplifier (ITTC Life Science Instruments, California, USA). Four blood pressure measurements were recorded before the animals were sacrificed.

2.4.13 Chronic in vivo studies using twenty-five percent $D_2O$

WKY and SHR rats were treated as in section 2.1
until seven weeks of age and at that time the drinking water was replaced by either distilled water or twenty-five percent D₂O in distilled water for twelve days. The animals were then sacrificed and the thoracic aorta removed. Aortic rings were equilibrated in regular Krebs for two hours following which a dose response curve was constructed using either 1x10⁻¹²M to 1x10⁻⁵M phenylephrine or 9.6 mM to 65.8 mM potassium chloride by the methods stated in experimental procedures 2.4.1 and 2.4.4 (only regular Krebs solution was used).

2.4.14 Acute in vitro studies using sixty percent D₂O

WKY and SHR rats were treated as in section 2.4.13. Experimental procedures 2.4.1 and 2.4.4 were then performed (both regular Krebs solution and sixty percent D₂O Krebs solution were used).

2.5 Analysis of data

Results are expressed as mean ± standard error of the mean. Where no S.E.M. is indicated the level of error was too small to be detected with the means of recording and analysis employed. Tissue responses are expressed as percentage maximum contraction or relaxation, where the maximum contraction or relaxation (100%) corresponds to that of the control tissue in each group.

Maximum contractions between groups were compared using an unpaired student's t-test. Significance was accepted at a
level of \( p < 0.05 \).

Plots were constructed using SigmaPlot, Scientific Graph System. The \( pA_2 \) and \( EC_{50} \) values were calculated using Pharmacologic Calculation System, version 4.1 by Tallarida and Murray (Springer-Verlag New York Inc., 1987).
CHAPTER 3

RESULTS

3.1 Effect of D_{2}O on aortic smooth muscle contraction

3.1.1 Phenylephrine induced contraction

In normal Krebs solution contraction of aortic rings using 1x10^{-9} M phenylephrine resulted in a maximum control contraction of 1.08 \pm 0.11 grams. Aortic rings incubated with increasing concentrations of D_{2}O had increasingly depressed phenylephrine induced maximal contractions (Figure 3-1). Twenty, forty, sixty, and eighty percent D_{2}O depressed the phenylephrine induced maximum contraction to 86.9\% \pm 12.6\%, 74.3\% \pm 5.0\%, 51.9\% \pm 6.5\%, and 42.4\% \pm 3.7\% of the control value respectively. The contractions were significantly depressed from forty percent to eighty percent D_{2}O.

Figure 3-2 shows the effect of varying concentrations of D_{2}O on the entire dose response curve constructed using 1x10^{-12} M to 1x10^{-5} M phenylephrine. Twenty, sixty, and eighty percent D_{2}O uniformly depress the dose response curve compared to the control curve. However, forty percent D_{2}O resulted in enhanced contraction from 1x10^{-11} M to 1x10^{-8} M phenylephrine. The dose response curve was depressed from 1x10^{-7} M to 1x10^{-5} M phenylephrine.
Figure 3-1: Inhibition of $1 \times 10^{-5}$M phenylephrine induced aortic smooth muscle contraction with varying concentrations of D$_2$O.

Columns represent mean percent maximum contraction and bars S.E.M. for eight tissues. An asterix represents a significant difference from the control contraction ($p < 0.01$).
Figure 3-2: The effect of varying concentrations of D$_2$O on phenylephrine induced aortic smooth muscle contraction.

Dose response curves were constructed using $1\times10^{-12}$M to $1\times10^{-5}$M phenylephrine. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
3.1.2 Reversibility

Reversal of the aortic smooth muscle inhibitory effect of sixty percent D$_2$O on phenylephrine induced contraction is shown in Figure 3-3. The first contraction dose response curves denoted by D$_2$Oa and H$_2$Oa are significantly different (p < 0.05). H$_2$Oa corresponds to the first control curve constructed in the absence of D$_2$O. D$_2$Oa corresponds to the first dose response curve constructed in the presence of sixty percent D$_2$O. The D$_2$O has reduced the maximum contraction to 64.9% ± 9.6% of the control value. However, the second phenylephrine dose response curve two hours following removal of D$_2$O (D$_2$Ob; Figure 3-3) is not significantly different from the second control curve (H$_2$Ob; Figure 3-3).

3.1.3 Role of the endothelium

Mechanical disruption of the endothelium had no effect on the control phenylephrine dose response curves (Figure 3-4; H$_2$O intact/scraped) or on the depressed curves of tissues incubated in sixty percent D$_2$O Krebs solution (Figure 3-4; D$_2$O intact/scraped). A level of aortic contraction similar to that previously observed was demonstrated in all tissues incubated with either regular Krebs or sixty percent D$_2$O Krebs solution.

3.1.4 Potassium chloride induced contraction

Sixty percent D$_2$O depressed the dose response curve constructed using 9.6 mM to 65.8 mM potassium chloride (Figure
Figure 3-3: Reversal of the D₂O inhibitory effect on phenylephrine induced aortic smooth muscle contraction.

\( H₂Oa/b \) represent the first and second control curves respectively. \( D₂Oa \) represents the first curve of tissues exposed to sixty percent \( D₂O \) Krebs solution; \( D₂Ob \) represents the second curve after removal of \( D₂O \). Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Figure 3-4: The role of the endothelium in the effect of sixty percent D$_2$O on phenylephrine induced aortic smooth muscle contraction.

Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
3-5). The control maximal contraction induced by 65.8 mM potassium chloride was 1.15 ± 0.10 grams and sixty percent D$_2$O depressed the potassium chloride induced maximum contraction to 51.9% ± 6.5% as compared to the control value. The difference in the potassium chloride dose response curve, constructed in D$_2$O Krebs, from the control curve, constructed in regular Krebs, was statistically significant (p < 0.02).

3.1.5 Calcium induced contraction

A calcium induced dose response curve was constructed after the contraction response to 60 mM KCl had been eliminated by washing aortic rings with calcium free Krebs solution. The level of the control maximum contraction at 5.0 mM calcium chloride was 0.93 ± 0.07 grams and sixty percent D$_2$O significantly (p < 0.05) reduced the maximum calcium contraction to 76.7% ± 5.9% as compared to control values (Figure 3-6).

3.1.6 Bay K 8644 induced contraction

The dose response curve constructed using 1x10^{-8}M to 1x10^{-5}M Bay K 8644 was significantly depressed by sixty percent D$_2$O (p < 0.05). The maximum control response in the absence of D$_2$O was 1.16 ± 0.11 grams. Sixty percent D$_2$O depressed this contraction response to 65.8% ± 11.2% as compared to the maximum control value (Figure 3-7).
Figure 3-5: The inhibitory effect of sixty percent D₂O on potassium chloride induced aortic smooth muscle contraction.

Dose response curves were constructed using 9.6 mM to 65.8 mM potassium chloride. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Figure 3-6: The inhibitory effect of sixty percent D$_2$O on the dose response curve of calcium in the presence of 60 mM potassium chloride.

Dose response curves were constructed using 1.0 mM to 5.0 mM calcium chloride. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Figure 3-7: The inhibitory effect of sixty percent D$_2$O on Bay K 8644 induced aortic smooth muscle contraction.

Dose response curves were constructed using $1 \times 10^{-9}$M to $1 \times 10^{-5}$M Bay K 8644. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
3.1.7 Sucrose induced contraction

Following incubation of aortic rings in regular Krebs a contraction response to 250 mM sucrose was performed. The amount of contraction of the control tissues to the sucrose was 0.23 ± 0.01 grams. Sixty percent D₂O significantly (p < 0.01) reduced the contraction to 55.6% ± 0.0% of the control value (Figure 3-8).

3.1.8 Synopsis

D₂O inhibited phenylephrine induced aortic smooth muscle contraction in a dose dependent manner with approximately 50% inhibition using 60% D₂O. This inhibitory effect was endothelium independent and reversible. Sixty percent D₂O also significantly inhibited potassium chloride, calcium, Bay K 8644, and sucrose induced aortic smooth muscle contraction.

3.2 Effect of D₂O on aortic smooth muscle relaxation

3.2.1 Isoprenaline induced relaxation

Rings equilibrated in sixty percent D₂O Krebs and regular Krebs were precontracted with 1x10⁻⁶M phenylephrine before relaxation with 1x10⁻⁴M isoprenaline. The level of contraction of tissues incubated in regular Krebs was 0.69 ± 0.01 grams while that of tissues incubated in sixty percent D₂O Krebs was 0.46 ± 0.01 grams. Treatment of the rings equilibrated in regular Krebs with 1x10⁻⁶M phenotolamine
Figure 3-8: The inhibitory effect of sixty percent D₂O on 250 mM sucrose induced aortic smooth muscle contraction.

Columns represent mean percent maximum contraction and bars S.E.M. for four tissues. An asterix indicates a significant difference from control values (p < 0.01).
resulted in a second contraction response to $1 \times 10^{-6}$M phenylephrine of $0.69 \pm 0.01$ grams. The second contraction response for tissues equilibrated in sixty percent D$_2$O Krebs was $0.66 \pm 0.02$ grams and this level of contraction was not significantly different from the second contraction response of tissues equilibrated in regular Krebs and phentolamine.

Sixty percent D$_2$O significantly ($p < 0.01$) reduced the relaxation response induced using $1 \times 10^{-4}$M isoprenaline to $50.3\% \pm 5.7\%$ of the maximum control response (Figure 3-9).

### 3.2.2 Acetylcholine induced relaxation

Prior to relaxation with acetylcholine, aortic rings were precontracted with $1 \times 10^{-5}$M phenylephrine. For rings equilibrated in regular Krebs solution the level of contraction was $1.39 \pm 0.08$ grams; the level of contraction for rings equilibrated in sixty percent D$_2$O Krebs solution was $1.08 \pm 0.07$ grams. Sixty percent D$_2$O significantly ($p < 0.05$) reduced the maximum relaxation to $76.7\% \pm 7.7\%$ as compared to the control value of tissues incubated in regular Krebs solution (Figure 3-10). Sixty percent D$_2$O also shifted the dose response curve of acetylcholine to the right. For tissues incubated in regular Krebs solution the EC$_{50}$ was $9.62 \times 10^{-8}$M while for tissues incubated in sixty percent D$_2$O Krebs the EC$_{50}$ was $5.64 \times 10^{-7}$M.
The inhibitory effect of sixty percent D$_2$O on $1 \times 10^{-4}$M isoprenaline induced aortic smooth muscle relaxation.

Columns represent mean percent maximum relaxation and bars S.E.M. for eight tissues. An asterix indicates a significant difference from the control value ($p < 0.01$).
Figure 3-10: The effect of sixty percent D$_2$O on the dose response curve constructed using $1\times10^{-10}$M to $1\times10^{-4}$M acetylcholine.

Points represent mean percent maximum relaxation and bars S.E.M. for six tissues.
3.2.3 Synopsis

Sixty percent D₂O reduced isoprenaline induced relaxation of aortic smooth muscle by approximately 50% and acetylcholine induced relaxation by 25%. D₂O also significantly changed the EC₅₀ of the acetylcholine relaxation response.

3.3 Effect of D₂O on the affinity of an antagonist for its receptor

3.3.1 Noncompetitive inhibition

In determination of the pD₂ for nifedipine the four control curves in the absence of D₂O were not significantly different (Figure 3-11). Likewise, the four control curves in the presence of sixty percent D₂O were not significantly different (Figure 3-12). Nifedipine at concentrations of 5x10⁻⁹M, 1x10⁻⁹M, and 1x10⁻⁸M depressed the maximum contraction response to 59.5% ± 7.1%, 38.9% ± 6.3%, and 8.3% ± 2.1% respectively of the control value in the absence of D₂O (Figure 3-13), and to 46.4% ± 5.2%, 21.4% ± 1.6%, and 3.6% ± 1.7% respectively in the presence of sixty percent D₂O (Figure 3-14).

The pD₂ for nifedipine in the absence of D₂O was 8.125 ± 0.001 whereas, the pD₂ for nifedipine in the presence of sixty percent D₂O was 8.453 ± 0.002 (Table 3-1). The difference between these values was statistically significant.
Figure 3-11: Four consecutive dose response curves constructed using 9.6 mM to 65.8 mM potassium chloride in the absence of D₂O.

Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
CONCENTRATION OF KCL (mM)

PERCENT MAXIMUM CONTRACTION

CURVE 1
CURVE 2
CURVE 3
CURVE 4
Figure 3-12: Four consecutive dose response curves constructed using 9.6 mM to 65.8 mM potassium chloride in the presence of sixty percent $\text{D}_2\text{O}$.

Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Percent maximum contraction vs. concentration of KCl (mM). The graph shows four curves labeled CURVE 1, CURVE 2, CURVE 3, and CURVE 4. Each curve represents a different concentration range of KCl, with concentrations ranging from 0 to 70 mM.
Figure 3-13: Noncompetitive inhibition of potassium chloride induced aortic smooth muscle contraction in the presence of nifedipine and in the absence of D$_2$O.

Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Figure 3-14: Noncompetitive inhibition of potassium chloride induced aortic smooth muscle contraction in the presence of nifedipine and sixty percent D₂O.

Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Table 3-1: Effect of 60% D₂O on the pD₂₅ for nifedipine.
<table>
<thead>
<tr>
<th>[nifedipine]</th>
<th>H₂O Krebs pD₂'</th>
<th>D₂O Krebs pD₂'</th>
</tr>
</thead>
<tbody>
<tr>
<td>5 x 10⁻⁹M</td>
<td>8.134</td>
<td>8.363</td>
</tr>
<tr>
<td>1 x 10⁻⁸M</td>
<td>8.197</td>
<td>8.564</td>
</tr>
<tr>
<td>1 x 10⁻⁷M</td>
<td>8.045</td>
<td>8.432</td>
</tr>
</tbody>
</table>

mean ± S.E.M. 8.125 ± 0.001  *8.453 ± 0.002
*p < 0.01
(p < 0.01).

3.3.2 Competitive inhibition

In the determination of the pA₂ for prazosin the four control curves in the absence of D₂O (Figure 3-15) were not significantly different and likewise, the four control curves constructed in the presence of sixty percent D₂O (Figure 3-16) were not significantly different. In the absence of D₂O and prazosin the EC₃₀ was 1.08x10⁻⁷M; 1x10⁻⁹M, 1x10⁻⁸M and 1x10⁻⁷M prazosin changed the EC₃₀ to 7.43x10⁻⁷M, 1.86x10⁻⁶M, and 1.01x10⁻⁵M respectively (Figure 3-17). Consequently, the pA₂ value for prazosin in the absence of D₂O was 9.81 ± 0.38 and the slope of the Schild plot was -1.10 ± 0.21. In the presence of sixty percent D₂O and in the absence of prazosin the EC₃₀ was 3.49x10⁻⁸M; 1x10⁻⁹M, 1x10⁻⁸M, and 1x10⁻⁷M prazosin changed the EC₃₀ to 1.94x10⁻⁷M, 1.23x10⁻⁵M, and 3.16x10⁻⁴M respectively (Figure 3-18). In determining the pA₂ for prazosin in the presence of sixty percent D₂O the slope of the Schild plot was -1.65 ± 0.14 and therefore no valid pA₂ value could be determined.

3.3.3 Synopsis

Sixty percent D₂O significantly changed both the pD₂ for nifedipine and the slope of the Schild plot for prazosin.
Figure 3-15: Four consecutive dose response curves constructed using $1 \times 10^{-12}$M to $1 \times 10^{-5}$M phenylephrine in the absence of D$_2$O.

Points represent mean percent maximum contraction and bars S.E.M. for six tissues.
PERCENT MAXIMUM CONTRACTION

LOG CONCENTRATION PHENYLEPHRINE (M)
Figure 3-16: Four consecutive dose response curves constructed using $1 \times 10^{-12}$M to $1 \times 10^{-5}$M phenylephrine in the presence of sixty percent D$_2$O.

Points represent mean percent maximum contraction and bars S.E.M. for six tissues.
Figure 3-17: Competitive inhibition of phenylephrine induced aortic smooth muscle contraction in the presence of prazosin and in the absence of D2O.

Points represent mean percent maximum contraction and bars S.E.M. for six tissues.
Figure 3-18:  Competitive inhibition of phenylephrine induced aortic smooth muscle contraction in the presence of prazosin and sixty percent D$_2$O.

Points represent mean percent maximum contraction and bars S.E.M. for six tissues.
3.4 Effect of D₂O on aortic smooth muscle from WKY and SHR rats

3.4.1 Blood pressure levels of WKY and SHR rats

The blood pressure levels of WKY and SHR rats may be seen in Table 3-2. The blood pressures of WKY untreated (control) rats were significantly (p < 0.01) lower than those of age matched SHR control rats. Oral consumption of twenty-five percent D₂O for twelve days significantly (p < 0.01) reduced the systolic blood pressure of SHR and WKY rats in comparison to their respective controls. SHR rats consuming twenty-five percent D₂O had reduced blood pressures which were not significantly different from that of WKY controls, i.e., 167 ± 5.3 and 161.2 ± 5.9 respectively.

3.4.2 Chronic in vivo effect of twenty-five percent D₂O

3.4.2.1 WKY/SHR: Potassium chloride induced contraction

The maximum level of KCl induced contractions for aortae of SHR control rats was significantly (p < 0.02) reduced to 64.4% ± 10.1% of the WKY control value of 0.65 ± 0.05 grams contraction (Figure 3-19). Chronic oral consumption of twenty-five percent D₂O, for twelve days by WKY rats had no significant effect on the in vitro potassium chloride induced contraction response curve when measured in normal Krebs solution. However, oral consumption of twenty-five percent D₂O by SHR rats produced a small but significant
Table 3-2: Effect of 25% D₂O consumption for 12 days on the systolic blood pressures of spontaneously hypertensive (SHR) and normotensive (WKY) rats.
<table>
<thead>
<tr>
<th>Type of rat</th>
<th>Type of water consumed</th>
<th>Systolic blood pressure (mmHg) mean ± S.E.M.*</th>
</tr>
</thead>
<tbody>
<tr>
<td>SHR</td>
<td>H₂O</td>
<td>210.1 ± 4.3</td>
</tr>
<tr>
<td>SHR</td>
<td>25% D₂O</td>
<td>167.0 ± 5.3</td>
</tr>
<tr>
<td>WKY</td>
<td>H₂O</td>
<td>161.2 ± 5.9</td>
</tr>
<tr>
<td>WKY</td>
<td>25% D₂O</td>
<td>109.0 ± 5.6</td>
</tr>
</tbody>
</table>

*Each value represents an average of four measurements; each group consisted of six rats.
Figure 3-19: Chronic in vivo effect of twenty-five percent D$_2$O on potassium chloride induced contraction of aortic smooth muscle from WKY and SHR rats.

CONTROL corresponds to muscle from rats consuming distilled water and D$_2$O to rats consuming twenty-five percent D$_2$O. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Concentration of Potassium Chloride (mM) vs. Percent Maximum Contraction.

- **WKY CONTROL**
- **WKY D2O**
- **SHR CONTROL**
- **SHR D2O**
(p < 0.05) depression of contraction, i.e., 41.0 ± 6.5% of the WKY control value.

3.4.2.2 WKY/SHR: Phenylephrine induced contraction

The maximum level of contraction induced using $1 \times 10^{-5}$ M phenylephrine was significantly (p < 0.01) different for aortic smooth muscle obtained from WKY and SHR rats (Figure 3-20). The level of contraction for SHR control rats was 63.4% ± 7.11% as compared to the WKY control value of 0.67 ± 0.06 grams.

Chronic oral administration of twenty-five percent D$_2$O had no significant effect on the phenylephrine induced contractions using aortic smooth muscle obtained from either WKY or SHR rats.

3.4.3 Acute in vitro effect of sixty percent D$_2$O

3.4.3.1 WKY: Potassium chloride induced contraction

The maximum level of contraction induced by 65.8 mM potassium chloride was not significantly different for aortic smooth muscle incubated in regular Krebs regardless of whether the WKY rats had previously orally consumed twenty-five percent D$_2$O (Figure 3-21; WKY CONT H$_2$O KREBS/WKY TREAT H$_2$O KREBS). In aortic smooth muscle obtained from WKY rats which had orally consumed twenty-five percent D$_2$O, sixty percent D$_2$O in vitro did not significantly alter the maximum contraction
Figure 3-20: Chronic in vivo effect of twenty-five percent D$_2$O on phenylephrine induced contraction of aortic smooth muscle from WKY and SHR rats.

CONTROL corresponds to muscle obtained from rats consuming distilled water and D$_2$O to rats consuming twenty-five percent D$_2$O. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Figure 3-21: Acute in vitro effect of sixty percent D$_2$O on potassium chloride induced contraction of aortic smooth muscle from WKY rats.

CONT corresponds to rats consuming distilled water and TREAT to rats consuming twenty-five percent D$_2$O in distilled water. H$_2$O corresponds to aortic rings equilibrated in regular Krebs solution and D$_2$O to aortic rings equilibrated in sixty percent D$_2$O Krebs. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
(Figure 3-21; WKY TREAT D2O KREBS). In aortic smooth muscle obtained from WKY rats which had not been in vivo deuterated by twenty-five percent D2O (Figure 3-21; WKY CONT D2O KREBS), sixty percent D2O Krebs significantly (p < 0.05) reduced the maximum contraction to 45.9% ± 8.5% as compared to the control value of 0.74 ± 0.01 grams.

3.4.3.2 WKY: Phenylephrine induced contraction

The maximum level of contraction induced by 1x10^-5M phenylephrine was not significantly different for aortic smooth muscle equilibrated in regular Krebs solution whether or not the WKY rats had been previously exposed to twenty-five percent D2O (Figure 3-22; WKY CONT H2O KREBS/WKY TREAT H2O KREBS). In aortic smooth muscle obtained from rats exposed to twenty-five percent D2O, sixty percent D2O did not significantly reduce the maximum contraction (Figure 3-22; WKY TREAT D2O KREBS). However, in aortic smooth muscle obtained from WKY rats which had not been previously exposed to twenty-five percent D2O (Figure 3-22; WKY CONT D2O KREBS), sixty percent D2O significantly (p < 0.05) reduced the maximum contraction to 62.5% ± 6.0% as compared to the control value of 0.61 ± 0.07 grams.

3.4.3.3 SHR: Potassium chloride induced contraction

The maximum level of contraction induced using 65.8 mM potassium chloride (0.21 ± 0.02 grams) was not significantly different for aortic smooth muscle equilibrated
Figure 3-22: Acute *in vitro* effect of sixty percent D$_2$O on phenylephrine induced contraction of aortic smooth muscle from WKY rats.

CONT corresponds to rats consuming distilled water and TREAT to rats consuming twenty-five percent D$_2$O in distilled water. H$_2$O corresponds to aortic rings equilibrated in regular Krebs solution and D$_2$O to aortic rings equilibrated in sixty percent D$_2$O Krebs. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
in regular Krebs solution whether or not the SHR rats had been previously exposed to twenty-five percent D₂O (Figure 3-23; SHR CONT H₂O KREBS/SHR TREAT H₂O KREBS). Sixty percent D₂O did not significantly change the maximum contraction of aortic smooth muscle obtained from either group of SHR rats.

3.4.3.4 SHR: Phenylephrine induced contraction

The maximum level of contraction induced using 1x10⁻⁵M phenylephrine (0.28 ± 0.07 grams) was not significantly different for aortic smooth muscle equilibrated in regular Krebs solution whether or not the SHR rats had been previously exposed to twenty-five percent D₂O (Figure 3-24; SHR CONT H₂O KREBS/SHR TREAT H₂O KREBS). In aortic smooth muscle obtained from rats previously exposed to twenty-five percent D₂O, sixty percent D₂O did not reduce the maximum contraction (Figure 3-24; SHR TREAT D₂O KREBS). However, in aortic smooth muscle obtained from SHR rats which had not been previously exposed to twenty-five percent D₂O (Figure 3-24; SHR CONT D₂O KREBS), sixty percent D₂O significantly (p < 0.05) reduced the maximum contraction to 54.6% ± 7.4% as compared to the control value.

3.4.4 Synopsis

Oral consumption of 25% D₂O significantly reduced the systolic blood pressures of SHR and WKY rats, with a reduction of SHR blood pressure to a level not significantly different from that of WKY control rats. Oral consumption of 25% D₂O had no significant effect on phenylephrine induced
Figure 3-23: Acute in vitro effect of sixty percent D$_2$O on potassium chloride induced contraction of aortic smooth muscle from SHR rats.

CONT corresponds to rats consuming distilled water and TREAT to rats consuming twenty-five percent D$_2$O in distilled water. H$_2$O corresponds to aortic rings equilibrated in regular Krebs solution and D$_2$O to aortic rings equilibrated in sixty percent D$_2$O Krebs. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
Figure 3-24: Acute *in vitro* effect of sixty percent D$_2$O on phenylephrine induced contraction of aortic smooth muscle from SHR rats.

CONT corresponds to rats consuming distilled water and TREAT to rats consuming twenty-five percent D$_2$O in distilled water. H$_2$O corresponds to aortic rings equilibrated in regular Krebs solution and D$_2$O to aortic rings equilibrated in sixty percent D$_2$O Krebs. Points represent mean percent maximum contraction and bars S.E.M. for eight tissues.
contraction of aortae from SHR and WKY rats; there was a small
depression of potassium chloride induced contraction in SHR
rats only. Subsequent acute exposure of aortae from SHR and
WKY rats to sixty percent D$_2$O demonstrated that significant
inhibition of contraction occurred only in aortae obtained
from rats which had not previously consumed 25% D$_2$O.
CHAPTER 4

DISCUSSION

4.1. Effect of D₂O on aortic smooth muscle contraction

Phenylephrine acts mainly on receptor operated calcium channels to produce contraction of aortic vascular smooth muscle (Khahil et al., 1987). Although phenylephrine is not completely selective for receptor operated calcium channels (Godfraind et al., 1986), it was used as a first approximation of the effects of deuterium oxide on agonist induced vascular smooth muscle contraction. Increasing concentrations of the D₂O depressed the maximal contraction in a dose-dependent manner (Figure 3-1) with an approximate IC₅₀ at 60% D₂O. Similar effects have been shown where concentrations of D₂O between 25 and 99.8 per cent decreased the force of contraction and the rate of spontaneously beating, isolated atria of the frog (Kaiminer, 1960). The entire dose response curve using 1x10⁻¹¹M to 1x10⁻⁵M phenylephrine (Figure 3-2) shows that varying concentrations of D₂O did not uniformly depress the lower end of the dose response curve; tissues exposed to forty per cent D₂O actually showed significantly increased contraction levels from 1x10⁻¹¹M to 1x10⁻⁶M phenylephrine. However, although these tissues demonstrated increased sensitivity at these lower drug doses, the level of contraction at the maximum concentration of phenylephrine was
significantly depressed in comparison to control tissues. These data, showing inhibition of phenylephrine contraction, may suggest an effect of D$_2$O on calcium entry via the receptor-operated calcium channel.

The series of experiments demonstrating reversibility of the D$_2$O inhibitory effects on vascular smooth muscle contraction (Figure 3-3) indicate the absence of tissue injury. Similar studies using skeletal muscle also show reversibility of the D$_2$O effect on the force of contraction (Svensmark, 1961).

Previous studies have shown that the endothelium does have an effect on the enhanced second phenylephrine induced contraction in the rat isolated aortic strip when carrying out multiple dose response curves in one tissue (Demirel and Türker, 1989). These workers found no diminished effect of removal of the endothelium in their first phenylephrine contraction curve. Accordingly, by utilizing the first curve only, it was confirmed that the amount of contraction produced by phenylephrine in the 0% D$_2$O Krebs controls was the same whether or not the endothelium had been disrupted (Figure 3-4). Furthermore, aortic rings equilibrated in 60% D$_2$O Krebs displayed approximately the same percentage of decreased contraction, in relation to those of control tissues, whether or not the endothelium had been disrupted. Thus the endothelium does not appear to play a major role in the D$_2$O
mediated inhibitory effect of vascular contraction. Acetylcholine was used at the end of each experiment to ensure that the endothelium scraping was effective since it produces a relaxation effect in previously contracted intact rings but has no effect on endothelium-denuded aorta (Rapoport and Murad, 1983).

Potassium chloride dose response curves were constructed to show the effect of D$_2$O on voltage-operated calcium channels since KCl acts mainly on this type of calcium channel (Meisher et al., 1981). Potassium chloride maximum contraction was reduced by 60% D$_2$O to a level similar to the inhibition of phenylephrine contraction (Figure 3-5). This suggests that D$_2$O also affects calcium uptake via voltage-operated calcium channels and supports the observations by Vasdev et al. (1990) of D$_2$O effects on $^{45}$Ca uptake by rat aorta.

Studies conducted in calcium free Krebs help to differentiate effects on extracellular calcium influx from effects on intracellular calcium mobilization. In these experiments tissues were washed in calcium free Krebs until there was no contractile response to 60mM KCl. This was considered an indication that intracellular calcium stores had been exhausted since it has been shown that the calcium release induced by high doses of pharmacological contractile agents in the absence of extracellular calcium exhausts the
intracellular calcium stores (van Breemen, 1969). Subsequent calcium induced contraction was depressed in the presence of 60% D$_2$O (Figure 3-6) confirming an action of D$_2$O on extracellular calcium influx. However, it may have been more appropriate to ensure depletion of intracellular calcium sources by washing with calcium free Krebs solution containing phenylephrine since phenylephrine mediates vascular smooth muscle contraction by mobilizing intracellular calcium sources whereas potassium depolarization is not believed to mobilize intracellular calcium (Deth and van Breemen, 1974).

Sixty percent D$_2$O similarly reduced the maximum contraction of aortic rings with Bay K 8644 (Figure 3-7) which is dependent upon extracellular calcium and facilitates activation of the voltage dependent calcium channel (Su et al., 1984). Since this protocol was carried out in the presence of phentolamine, D$_2$O effects on receptor-operated calcium channels were excluded.

These results suggest that a major action of D$_2$O is on extracellular calcium influx through voltage-operated calcium channels.

These studies do not conclusively indicate whether D$_2$O is producing selective effects only on the voltage- and receptor-operated calcium channels. There appears to be an effect on both types of channel but D$_2$O could be acting at multiple
Evidence for actions of D₂O on intracellular calcium mobilization was obtained from studies of aortic contraction with sucrose. Andersson et al. (1974) have shown that a sucrose induced contraction of vascular smooth muscle occurs independently of membrane potential level and in the absence of extracellular calcium. D₂O depression of the sucrose contraction in this study (Figure 3-8) suggests further sites of action independent of extracellular calcium sources. However, conclusions from this study are limited since the precise mechanism of sucrose induced contraction is unknown and the absolute tension achieved was very low compared with other protocols in this study.

4.2. Effect of D₂O on aortic smooth muscle relaxation

Studies of the effect of D₂O on aortic smooth muscle relaxation were performed to further elucidate the mechanism of action of D₂O. For this purpose aortic smooth muscle relaxation with isoprenaline and acetylcholine was investigated. Since isoprenaline induced relaxation was measured on tissues precontracted with phenylephrine the contraction level of tissues equilibrated in 0% D₂O was reduced by use of the alpha-adrenoceptor antagonist phentolamine. The relaxation response with the β-adrenoceptor agonist, isoprenaline, in tissues equilibrated in 60% D₂O Krebs was significantly reduced compared to control tissues
(Figure 3-9). Sixty percent D$_2$O also depressed aortic smooth muscle relaxation with acetylcholine (Figure 3-10) which requires the presence of an intact endothelium and produces relaxation by the release of EDRF. Along with depressing the acetylcholine dose response curve, 60% D$_2$O has also shifted the curve to the right. This may indicate a combination of noncompetitive and competitive inhibition. These results indicate that the D$_2$O effect is not contraction specific but can also affect vascular smooth muscle relaxation, perhaps through an action on sarcoplasmic reticulum calcium mobilization or on the contractile proteins. The fact that D$_2$O affects relaxation using both isoprenaline and acetylcholine which act by distinct mechanisms (an increase in cAMP and cGMP, respectively) again suggests that D$_2$O acts at multiple sites.

4.3. Effect of D$_2$O on the affinity of an antagonist for its receptor

In studies to this point D$_2$O acted as an inhibitory agent in each protocol. As a further attempt to make a definite conclusion as to the site of action of D$_2$O, its effects on noncompetitive and competitive inhibition were studied using nifedipine and prazosin respectively. D$_2$O significantly changed the pD$_2$' for nifedipine (Table 3-1) which selectively inhibits calcium entry through voltage dependent calcium channels (Cauvin et al., 1983) producing a decreased vascular smooth muscle contraction. There is evidence that Bay K 8644
and nifedipine bind to the same binding site (Schramm et al., 1985) but whereas nifedipine blocks calcium entry, Bay K 8644 enhances it. Since the results showed that D₂O affected both Bay K 8644 contraction and the pD₂ for nifedipine this suggests a possible interaction at the shared binding site.

The pA₂ value for prazosin in the absence of D₂O was calculated as 9.81 ± 0.38 which was very similar to that determined by Muramatsu et al. (1990) for rat aorta. Sixty percent D₂O significantly changed the slope of the Schild plot and consequently no pA₂ value could be determined in the presence of D₂O. The change in the slope of the Schild plot may indicate an alteration in the kinetics of the interaction between prazosin and its binding site.

These data again suggest multiple sites of action for D₂O.

4.4. Chronic and acute effects of D₂O on aortic smooth muscle from WKY and SHR rats

The previous studies dealt only with acute in vitro effects of D₂O. Therefore the chronic in vivo effect of 25% D₂O on normotensive (WKY) and spontaneously hypertensive (SHR) rats was also studied.

Toal and Leenen (1985) have reported that significant differences in blood pressure can be detected at four weeks of age and that SHR blood pressure progressively increases with
age. In this study, systolic blood pressure measurements confirmed that SHR control rats had significantly higher blood pressure than WKY control rats (Table 3-2). Seven weeks is clearly sufficient for the development of hypertension in the SHR rats and after that time D$_2$O treatment was started. Preliminary blood pressure measurements should have been made before D$_2$O treatment was started to ensure that WKY rats were normotensive and SHR rats were hypertensive. Since this procedure was not followed the D$_2$O treated rats were simply compared with control rats which did not receive D$_2$O. The blood pressure levels for the SHR and WKY rats in this experiment were measured by tail plethysmography but the differences in levels for SHR and WKY rats are similar to those measured by cannulae insertion into the aortic arch (Toal and Leenen, 1985). Oral consumption of 25% D$_2$O in the drinking water for 12 days reduced the blood pressure of SHR rats to a level not significantly different from that of WKY control rats. D$_2$O also reduced the blood pressure level of WKY rats. However, the blood pressure measurements of the WKY rats were high compared with those of other researchers (Vasdev et al., 1990a). The high blood pressure was probably due to the fact that the rats were not acclimatized before the blood pressure measurements were made.

The potassium chloride and phenylephrine induced contraction levels of control SHR rats were significantly lower than those of WKY rats (Figures 3-19 and 3-20). Although
in vivo studies usually indicate an increase in the contractility of vascular smooth muscle obtained from SHR rats, many in vitro studies have reported depressed contraction levels (Triggle, 1989). Spector et al. (1969) reported that aortic strips from spontaneously hypertensive rats were less responsive to contraction induced with norepinephrine, serotonin, and potassium chloride. Similarly, Field et al. (1972) reported decreased isometric tension produced in response to norepinephrine and potassium chloride in aortic rings of SHR rats compared to normotensive controls.

Oral consumption of 25% D$_2$O did result in a small depression in the in vitro potassium chloride dose response curve of aortae from SHR rats (Figure 3-19). However, after oral consumption of 25% D$_2$O there was no significant effect on subsequent in vitro potassium chloride induced aortic contractions from WKY rats (Figure 3-19) or phenylephrine induced contractions from WKY or SHR rats (Figure 3-20). Previous in vitro studies showed that 20% D$_2$O had no significant effect on phenylephrine induced aortic contraction (Figure 3-1), so it is not surprising that the in vivo effect of 25% D$_2$O was small. A higher concentration of D$_2$O was not used since Thomson (1960) found that rats died when about one third of their body water was replaced by D$_2$O.

Even though 25% D$_2$O reduced the blood pressure of SHR rats to the level of WKY control rats, the potassium chloride
and phenylephrine induced maximal contraction levels for the SHR rats were still significantly different from that of WKY control rats. It has been noted in several studies, that a reduction in blood pressure after hypertension has developed in SHR rats does not reverse vascular reactivity changes (Triggle, 1989). This may explain the apparent contradiction between blood pressure and aortic contraction following chronic D₂O treatment.

The acute studies revealed that maximum contraction with either phenylephrine or potassium chloride was not significantly different for aortic rings equilibrated in regular Krebs solution, regardless of whether the SHR or WKY rats had previously consumed 25% D₂O (Figures 3-21 to 3-24; H₂O Krebs). Even small chronic effects which were evident in the first part of the experiment (Figure 3-19; SHR D₂O) were no longer detectable after equilibration in regular Krebs solution for two hours. This may indicate that the D₂O effect is reversible as demonstrated in previous studies (Figure 3-3).

In all cases, aortic rings obtained from either SHR or WKY rats which had consumed 25% D₂O, did not show significantly decreased contraction levels when rings were equilibrated in sixty percent D₂O rather than normal Krebs (Figures 3-21 to 3-24). When potassium chloride was used as the contractile agent there was a trend towards a decrease in
contraction levels but it was not statistically significant (Figures 3-21 and 3-23). There is a very large standard error in the potassium chloride dose response curves of SHR rats (Figure 3-23) which may partly be due to the fact that the level of contraction was very small since SHR rats show decreased contraction levels, coupled with the inhibitory effects of 25% and 60% D₂O. However, from figures 3-21, 3-22, and 3-24, it is clear that rings obtained from SHR and WKY rats which had not previously consumed 25% D₂O showed significantly decreased potassium chloride and phenylephrine induced contraction levels in 60% D₂O Krebs solution. The level of inhibition in SHR and WKY rats was similar to that previously found in Sprague-Dawley rats (Figures 3-2 and 3-5). These data from the acute studies suggest that homologous desensitization or tolerance to D₂O may have occurred since 60% D₂O has a reduced effect in aortic rings from rats with prolonged exposure to 25% D₂O. This would explain why D₂O produces a significant depression of contraction levels only in aortic rings obtained from rats not previously exposed to D₂O.

4.5. Summary and conclusion

From these studies it cannot be determined at which specific stage of the contraction process D₂O is producing its effect. In studies performed by Svensmark (1961) on frog skeletal muscle, it remained undecided as to whether force
reduction was due to the effect of D$_2$O on the contractile elements or on the excitation-contraction coupling. Similarly, no definite conclusions can be drawn from the work of Goodall (1958) on frog sartorius muscle or Kaiminer (1960) on glycerol extracted rabbit's psoas, as to the exact site of D$_2$O action. The level of inhibition was approximately 25% in the Bay K 8644 and calcium contraction study compared to 50% inhibition with the sucrose contraction and isoprenaline relaxation protocols. This difference suggests a greater effect of D$_2$O on intracellular calcium mobilization or other intracellular events, rather than on the influx of extracellular calcium. However, the observations that D$_2$O does inhibit contraction with phenylephrine, potassium chloride, Bay K 8644, and calcium all suggest that a major influence of D$_2$O is on extracellular calcium influx. Further evidence suggesting a nonspecific action for D$_2$O was obtained from the fact that D$_2$O inhibited relaxation with isoprenaline and acetylcholine, and affects both the noncompetitive inhibition of potassium chloride contraction with nifedipine and the competitive inhibition of phenylephrine contraction with prazosin. Taken together, the data indicate multiple sites of action of D$_2$O on vascular smooth muscle.

Data from the acute studies suggest that homologous desensitization to D$_2$O may have occurred since D$_2$O produces a significant depression of potassium chloride and phenylephrine induced contraction levels only in aortic rings obtained from
SHR and WKY rats which had not previously consumed D$_2$O. It is not possible to isolate the mechanism of desensitization, such as down regulation of receptors, since the site of action of D$_2$O is unknown and appears to be nonspecific. Although chronic oral consumption of 25% D$_2$O reduced the systolic blood pressure in SHR and WKY rats, subsequent in vitro aortic contraction was not always similarly reduced. It may be concluded that the hypotensive effect of chronic D$_2$O administration does not entirely appear to be a consequence of vascular reactivity changes.

Tolerance to the effects of D$_2$O suggest that there have indeed been changes in membrane function and/or regulation or mobilization of calcium sources. There may be membrane receptor changes associated with D$^+$ and H$^+$ exchange since proton exchange is important in most biological reactions and isotopic effects due to D$^+$ and H$^+$ exchange have been shown to affect L-type calcium channel conductance (Prod'hom et al., 1987). Future studies should be directed toward performing receptor binding studies to determine if tolerance does occur due to down regulation of receptors. The effect of chronic in vivo exposure to D$_2$O on the pA$_2$ and pD$_2$' values of competitive and noncompetitive antagonists, respectively, may give some insight into whether receptor modifications occur due to D$^+$ and H$^+$ exchange. Finally, an analysis of calcium homeostasis in D$_2$O tolerant rats may demonstrate whether D$_2$O affects calcium regulation.
In summary, these studies indicate that deuterium oxide reduces vascular smooth muscle contraction induced through agonist and depolarization activation of calcium channels. This depression of contraction is reversible and not dependent on an intact endothelium. The action of D₂O appears not to be solely dependent on intracellular calcium mobilization or extracellular calcium utilization as determined by calcium free buffer studies and sucrose and Bay K 8644 induced contraction. Since D₂O has an effect on relaxation of vascular smooth muscle, sucrose induced contraction, and the action of both competitive and noncompetitive antagonists, this suggests multiple sites of action and perhaps an action on the contractile elements themselves. Chronic studies show that oral consumption of 25% D₂O reduces systolic blood pressure in hypertensive and normotensive rats but the hypotensive effect does not appear to be a consequence of vascular reactivity changes. Finally, chronic in vivo exposure to D₂O desensitizes aortic rings to subsequent inhibition by acute in vitro D₂O exposure.
REFERENCES


