# CARBON DIOXIDE ADMINISTRATION TO CALVES

Thesis for the Degree of M. S. MICHIGAN STATE UNIVERSITY Seref Nezihi Ersoz 1962





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#### ABSTRACT

## CARBON DIOXIDE ADMINISTRATION TO CALVES

## by Seref Nezihi Ersoz

An investigational study has been made to determine the possible surgical anesthetic effect of carbon dioxide  $(CO_2)$ .

A review of literature did not reveal satisfactory references in veterinary medicine with the exception of administration of  $CO_2$  to slaughter hogs and poultry. In human medicine,  $CO_2$  has been used during child delivery, in the attempted treatment of alcoholism, muscular rigidity and psychoneurotic cases.

The experimental equipment consisted of two  $CO_2$  tanks, one oxygen  $(O_2)$  tank, a  $CO_2$  gas analyzer (Ranarex), an electrocardiograph (Burdick EK III), and a kymograph (Watson Bird).

Initially 100%  $CO_2$  was administered until the calves fell into lateral recumbency, following this 65%  $CO_2$  and 35%  $O_2$  was used to maintain anesthesia and the following data were collected before and during  $CO_2$  anesthesia: Total leukocyte and erythrocyte counts, differential blood counts, packed cell volumes (hematocrit), hydrogen ion concentrations, arterial blood pressures, electrocardiograms, pneumograms, and body temperatures. During anesthesia response to corneal and cutaneous pain stimuli were periodically observed.

The principal findings were: 1) on the basis of experimental data no consistent significant changes were noted in the hematocrit values, total erythrocyte, leukocyte, and differential counts, 2) a consistent decrease of blood pH varying between 0.3 and 1.1, 3) an increase in blood pressure ranging from 13 to 112 mm. Hg., 4) an increase of heart rate between 30 and 120 beats per minute above normal, 5) significant changes in the conduction system of the heart during CO<sub>2</sub> administration were tachycardia, possible heart injury, and on one calf the Wenchebach phenomena were observed.

The pertinent observations were: 1) blood stains in the nasal discharge on three calves, 2) deep respiratory movements, 3) defecation at the beginning or during  $CO_2$ administration, 4) intermittent bellowing, 5) involuntary movements of the limbs of two calves, 6) a partial to complete absence of response to corneal and cutaneous pain stimuli.

The results of the findings and observations are discussed with a summary followed by references and a bibliography.

## CARBON DIOXIDE ADMINISTRATION TO CALVES

By

Seref Nezihi Ersoz

## A THESIS

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ii

Dedicated to My Instructors

Prof. Dr. S. Nejat Yalki Prof. Dr. Yasar Altan

# TABLE OF CONTENTS

## CHAPTER

I.	INTRODUCTION
II.	REVIEW OF THE LITERATURE
III.	MATERIALS AND METHODS
	Experimental Animals
	Experimental Equipment
	Methods
IV.	RESULTS AND DISCUSSION
	Physiological Data
	Hematological Findings
	Electrocardiographic Findings
	Observations on Anesthetic Effect of CO <sub>2</sub> 33
v.	SUMJARY
REFEREN	CES
BIBLIOG.	RAPHY

# LIST OF FIGURES

**.**..

.

i

1.	The CO <sub>2</sub> Apparatus
2.	The CO <sub>2</sub> Gas Analyzer (Ranarex)
3.	The Beginning of CO2 Administration
4.	Calf Under CO <sub>2</sub> Anesthesia
5.	Electrocardiograph (Burdick EK III) 24
6.	Kymograph (Watson-Bird Variable Speed 70-142) 26
7.	Pneumogram, Calf No. 1
8.	Pneumogram, Calf No. 2
9.	Pneumogram, Calf No. 3
10.	Pneumogram, Calf No. 4
11.	Pneumogram, Calf No. 5
12.	Electrocardiogram, Calf No. 1
13.	Electrocardiogram, Calf No. 2
14.	Electrocardiogram, Calf No. 3
15.	Electrocardiogram, Calf No. 4 60
16.	Electrocardiogram, Calf No. 5 62

# LIST OF TABLES

## TABLE

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1.	Hematological Findings, Calf No. 1
2.	Physiological Data, Calf No. 1
. 3.	Hematological Findings, Calf No. 2
4.	Physiological Data, Calf No. 2
5.	Hematological Findings, Calf No. 3
6.	Physiological Data, Calf No. 3
7.	Hematological Findings, Calf No. 4
8.	Physiological Data, Calf No. 4
9.	Hematological Findings, Calf No. 5
10.	Physiological Data, Calf No. 5
11.	Electrocardiographic Findings, Calf No. 1 40
12.	Electrocardiographic Findings, Calf No. 2 40
13.	Electrocardiographic Findings, Calf No. 3 41
14.	Electrocardiographic Findings, Calf No. 4 41
15.	Electrocardiographic Findings, Calf No. 5 42

#### CHAPTER I

#### INTRODUCTION

Carbon dioxide (CO<sub>2</sub>)--its production in the body, effects upon body systems, and elimination from the body-has always been of concern to anesthesiologists, as well as physiologists.

In human medicine,  $CO_2$  has been used for a number of different conditions: child delivery (MacRae, 1951), treatment of neuroses and alcoholism (LaVerne, 1953), treatment of stutterers (Smith, 1953), and muscular rigidity (Fay, 1953). The most extensive studies with CO<sub>2</sub> have been made by Meduna (1953 and 1958) who has written a text book entitled "Carbon Dioxide Therapy--A Neurophysiological Treatment of Nervous Disorders." Within recent years, an increasing interest has been taken in the anesthetic properties of CO<sub>2</sub> in veterinary medicine (Swem, 1952; Murphy, 1953; Blomquist, 1957; Wernberg, 1957; Kitchell, et al., 1960), as well as its usage in meat packing plants to produce surgical anesthetic effects before exsanguination. The "Humane Slaughter Law" passed by the 85th U.S. Congress as a public law in 1958 required slaughter houses to dispatch with domestic animals in a humane manner. Carbon dioxide has been approved for a pre-surgical anesthetic for swine, sheep,

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and calves (Title 9, Code of Federal Regulations, 1959).\*

This study was conducted as a result of interest aroused by the work of previous investigators and the apparent lack of scientific data regarding the effect of CO<sub>2</sub> on calves.

<sup>\*</sup>Regulations governing the meat inspection of the United States Department of Agriculture, United States Government Printing Office, Washington, 1960.

#### CHAPTER II

## REVIEW OF THE LITERATURE

In 1827, Hickman, cited by McQuacker (1929), first demonstrated that animals exposed to an atmosphere containing a high concentration of  $CO_2$  became anesthetized. He performed experimental surgery on a dog, puppy, mouse and rabbit using  $CO_2$  as the anesthesia inducing agent.

In 1878 Bert, cited by Loewy (1923), recommended the gas for anesthesia and reported that  $CO_2$  concentrations of 25 to 30% in the inspired air would effect paralysis of the central nervous system with cessation of corneal and other reflexes. Zuntz and Loewi (1897) and Kropeit (1898) observed that in rabbits there occurred a short excitation stage, but no such stage has been observed in cold blooded animals (Winterstein, 1911). Albitzki (1912) stated in experiments with rabbits that if the concentration of  $CO_2$  in inspired air was lower than 25 to 30% complete anesthesia was not attained. But following recovery from anesthesia they showed disturbances of balance, as well as a failure in the ability to keep upright and to change their positions.

Prausnitz (1928) showed that 25 to 30% CO<sub>2</sub> was required in the inspired air to abolish corneal and light

reflexes in experiments on dogs, cats, rabbits, guinea pigs, rats, and mice. The excitation stage was slight, and awakening occurred quickly even though the anesthesia lasted for one hour. The animals tolerated a concentration of 50% CO<sub>2</sub> resulting in deep anesthesia for 60 to 90 minutes. Unfortunately, after this period of anesthesia the animals showed convulsions and died of pulmonary edema.

Leake and Waters (1928), in experiments on dogs, found that 30 to 40%  $CO_2$  with oxygen  $(O_2)$  will anesthetize the dogs in about one minute usually without struggling. The pulse rate was usually reduced as was the respiratory rate, but the blood pressure rose rapidly. The depth of respiration was greatly increased, but breathing did not seem to be labored. Upon opening the abdominal cavity, the intestines seemed relaxed and the muscles were usually flaccid. Dogs have been maintained under fairly satisfactory anesthesia for two hours. Respiratory movements became more regular and the depth of anesthesia deeper the longer the animal received the same mixture. According to Leake and Waters (1928), the lower percentages of CO<sub>2</sub> did not seem to anesthetize as well. Concentrations higher than 40% tended to depress respiration. Cardiac function and anoxemia became a factor with concentrations above 80%. With 100% CO2, the blood pressure became lower and the heart and respiration stopped at about the same time. In another experiment one year later, these investigators demonstrated

anesthesia with 30-40%  $CO_2$  in a variety of animals obtaining comparable results. They also pointed out that it was theoretically significant that high concentrations of a metabolic waste product  $(CO_2)$  would depress the central nervous system enough to produce anesthesia. These investigators concluded that the anesthetic effect, in the case of a mixture of 30 to 40%  $CO_2$  with  $O_2$ , was not due to asphyxia. The theory being that the increased  $CO_2$  tension in the blood prevented the escape of  $CO_2$  from the cells, and that the gradual accumulation of this metabolic product may progressively depress cellular function.

In 1926, Loevenhart, Lorenz and Waters first introduced CO<sub>2</sub> inhalation therapy into psychiatry. They theorized that a disordered function of the brain may be corrected by physiological and biochemical means. The report of this study was published in 1929 but their technique has since been abandoned.

Hettwer (1938) examined the relation of excitability of frog sciatic nerve to  $CO_2$  tension of the external medium by systemic, graphic action potential threshold determinations and found a reduced irritability of isolated nerves after exposure to  $CO_2$ .

Barbour and Seevers (1941a) conducted a series of experiments to determine the influence of cold on the narcotic action of CO<sub>2</sub> on rats, rabbits and dogs in a chamber at 5 C. They reported the minimal narcotic concentrations

at this temperature as follows: rat, 11%; rabbit, 17%; and dog, 14%. The induction time varied from 3 to 24 hours. These concentrations of  $CO_2$  did not induce narcosis at 25 The same authors (1941b) showed in albino rat experi-C. ments that the maximum tolerated concentration of CO2 for the first time was between 15 to 20%. At 20% an occasional animal survived. Atmospheres containing between 25 to 30% CO2 were uniformly lethal. Survival time varied from 0.5 to 36 hours, depending on the concentration of  $CO_2$ . At 50% CO<sub>2</sub>, no animal survived longer than five hours. Death seemed to be due primarily to pulmonary injury rather than narcosis. The edema and hemorrhage noted in the lungs were also seen on all visible mucous membranes. Continuing the experiments, Barbour and Seevers (1943a) also pointed out that during the exposure of albino rats to 11% CO<sub>2</sub> for 30 days, the plasma pH fell to a level as low as 7.09 in the first half hour and remained below 7.15 during the remainder of the period. A marked reticulocytosis occurred in the rat during prolonged exposure to 11% CO2, but no corresponding increase in the number of leukocytes or erythrocytes was observed. In another experiment they (1943b) concluded that the time required to produce the state of narcosis varied with the concentration of the  $CO_2$ , the species, and previous condition of the animal.

Gellhorn and Yesinick (1941) produced convulsions in anesthetized cats by the injection of metrazol,

picrotoxin, corymyrtin, camphor, strychnine and other drugs. In the non-denervated cats, CO<sub>2</sub> exerted a greater anticonvulsant effect whereas in those that were denervated, anoxia was produced. In experiments on dogs, Draper and Whitehead (1944), and Parry, et al. (1949) succeeded in arresting respiration for 45 minutes by administering a barbiturate and CO<sub>2</sub> anesthesia and permitting the dog to rebreathe expired air. By the end of the experiments, up to 50% CO<sub>2</sub> was found in the alveolar air. In other experiments the acidity of the venous blood was determined after inspiration of different concentrations of CO2. Inspiration of 40% CO<sub>2</sub> was found to effect an average fall of blood pH from 7.42 to 6.79. The only alteration in the electrocardiogram was an increase in the amplitude of the T wave, which returned to normal following restoration of the normal blood pH.

Paulsen (1949) described an apparatus with an arrangement for continuous production of anesthetic gas mixtures consisting of  $CO_2$  and ether. In a series of experiments on rabbits Paulsen (1952) used 30 to 35%  $CO_2$  and observed that  $CO_2$  produced a marked rise of the  $CO_2$  concentration in the blood of these animals, and at the same time a fall in the blood pH from about 7.5 to 6.5. The corneal reflex also ceased at this concentration.

In human medicine, Meduna (1950, 1953) conducted a series of experiments using 30% CO<sub>2</sub> and 70% O<sub>2</sub> on

psychoneurotic individuals. This line of therapy was given to more than 40,000 patients. No complications were reported other than tongue biting (1 case) and 3 or 4 spontaneous urination (3 or 4 cases). MacRae (1951) found that inhalation of suitable mixtures of  $CO_2$  and  $O_2$  produced coma which was both safe and beneficial in delivery at child birth. He stated that as an anesthetic for delivery "CO<sub>2</sub> coma" is more satisfactory both for the mother and the baby than other anesthetics. However, some other anesthetic is needed for surgical repair because the "CO2 coma" is brief. LaVerne (1953), using higher concentrations of CO2 such as 70%  $CO_2$  and 30%  $O_2$ , demonstrated the safety of this mixture on 4,000 patients. It should be taken into consideration that the CO<sub>2</sub> treatment has some contraindications, such as: previous history of coronary attack, organic heart disease, advanced arterio-sclerosis, hypertension and emphysema.

Within recent years the veterinary profession has shown an increased interest in the anesthetic properties of  $CO_2$ . Swem (1952) and Murphy (1953) at Geo. A. Hormel & Co. employed a mixture of  $CO_2$  to immobilize the slaughter hogs for ease in shackling and sticking. By using this method, both the carcass yield and by-products were improved.

Johnson (1960) believes that the ability of CO<sub>2</sub> to produce anesthesia is by loss of function of the consciousness center of the brain because of the associated

anesthetic properties of the gas and the markedly reduced supply of  $0_2$  to the brain. This is accomplished before vital heart action ceases. It is known that an isolated nerve in high CO<sub>2</sub> tension milieu demands greater irritation to react, that is, to send out impulses. In addition, it is known that the speed of the impulse along the nerve is reduced. CO2 also has the effect of making the transfer of impulses from one nerve cell to another more difficult because the connections between the nerve cells (synapses) are partly blocked by CO<sub>2</sub>. Meduna (1953) is in agreement with this concept. On the other hand, Murphy (1953), working with hogs, contended that the CO<sub>2</sub> unites with the hemoglobin of the blood and thereby causes unconsciousness. The compound rapidly separates from the hemoglobin and is discharged by exhalation as the hog continues to breathe fresh air before and after sticking.

As a result of the swine studies, Kotula, Drewmak and Davis (1956) used 33 to 36% CO<sub>2</sub> in stunning chickens, and Blomquist (1957) and Wernberg (1957, 1962) further investigated reactions of hogs to CO<sub>2</sub> stunning during an experiment carried out by the Danish Meat Research Institute in which 70% CO<sub>2</sub> was used. Pietraszek (1958) and Bartels (1962) also outlined the effect of CO<sub>2</sub> in stunning hogs.

In the United States the Humane Slaughter Law was passed by the 85th Congress as a Public Law 85-765 and became effective August 27th, 1958. The purpose of this law

was to require the use of humane methods in the slaughter of livestock. The animals must be exposed to the CO<sub>2</sub> gas in such a way as to produce surgical anesthesia before they are shackled, hoisted, thrown, cast or cut. Carbon dioxide is a relatively cheap and non-flammable gas, commercially available in large quantities and when used with properly designed units, offers no hazard to employees. These factors have stimulated research to determine its value as a surgical anesthetic for use on animals.

## CHAPTER III

## MATERIALS AND METHODS

## A--Experimental Animals

All experiments in this study were conducted on five calves as described below:

Tag No.	Breed	Age	Sex	Weight
l	Holstein	3 months	М	260 lbs.
2	Holstein	4 months	М	335 lbs.
3	Holstein	4 months	М	350 lbs.
4	Holstein	3 months	F	272 lbs.
5	Jersey	3 months	М	360 lbs.

The above animals were housed in the Veterinary Clinic of the College of Veterinary Medicine, Michigan State University, and were fed good quality alfalfa hay.

## B--Experimental Equipment

The CO<sub>2</sub> apparatus used for induction of anesthesia consisted of:

1. Two  $CO_2$  tanks each equipped with two gauges (cylinder contents and outflow regulator) and a flow-meter (Figure 1). The tanks were connected with a copper tubing in such a manner that the gas from one or both of them passed through an Automatic  $CO_2$  Heater (Figure 1) before going

Figure 1. The CO<sub>2</sub> Apparatus

- A. CO<sub>2</sub> tanks
- B. Automatic CO<sub>2</sub> Heater
- C. Cylinder Contents Gauge
- D. Outflow regulator
- E. Flow-meter

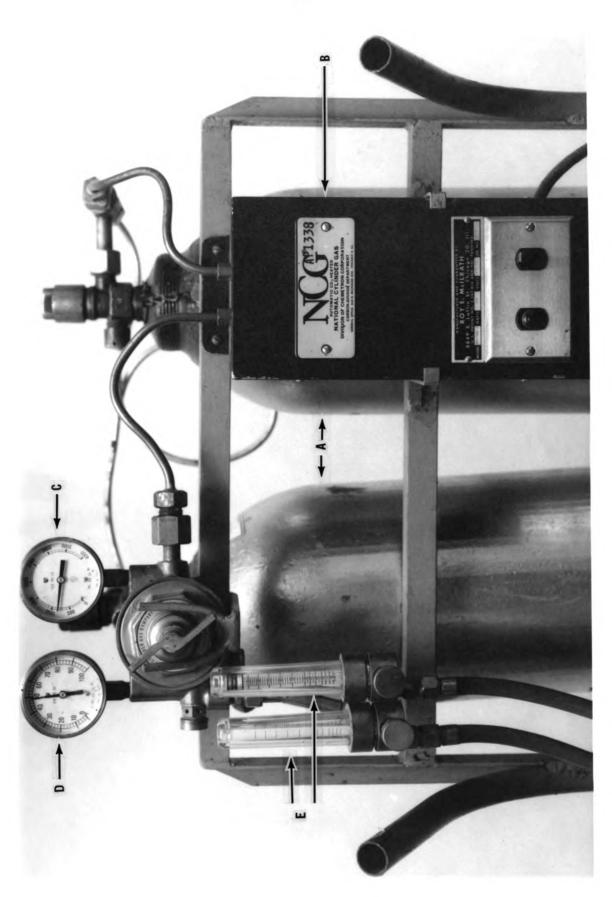


Figure 2. The CO<sub>2</sub> Gas Analyzer, Ranarex

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Figure 3. The Beginning of CO<sub>2</sub> Administration

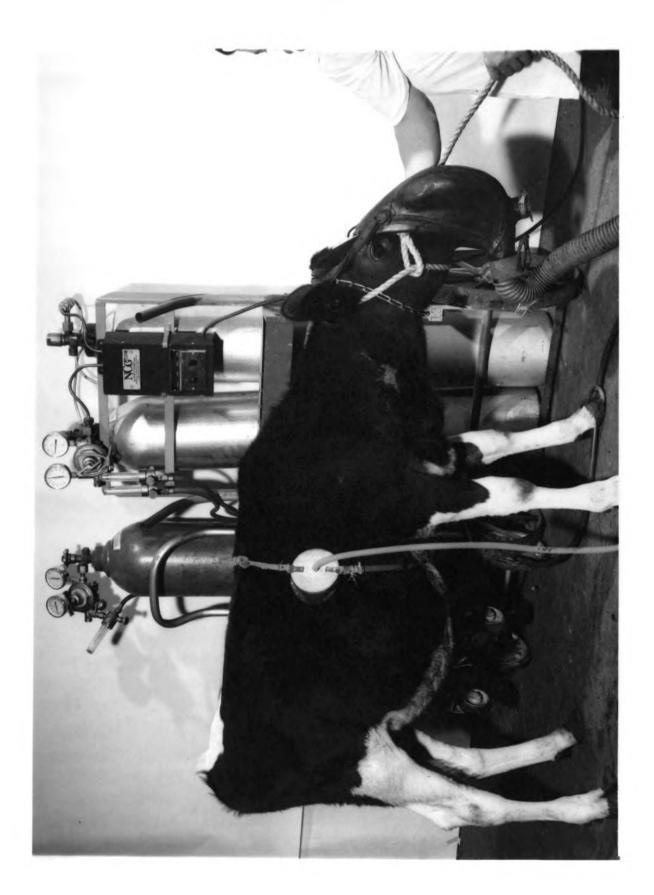


Figure 4. Calf under CO<sub>2</sub> Anesthesia



into the inhalation mask. \* This unit was mounted on a twowheeled truck.

2. One O<sub>2</sub> tank mounted on a truck and likewise equipped with two gauges (as above) and a flow-meter (Figure 1).

3. An equine protective field mask, M4, which will be called an inhalation mask.

4. A  $CO_2$  gas analyzer, Ranarex, <sup>\*\*</sup> which gave a continuous analysis of the concentration of  $CO_2$  being inhaled by the calf. The gas sampling tube was connected to the tube containing the  $CO_2-O_2$  mixture 5 ft. from the mask. At this point, the gases had flowed 25 ft. from their source and should have been thoroughly mixed.

## C--Methods

The initial stages of administering the anesthesia consisted of giving the calves  $100\% \text{ CO}_2$  flowing at a rate of 150 lts. per hour. Within 25 to 37 seconds after the calves went down, a sufficient flow of  $O_2$  was supplied to result in a  $CO_2$  concentration of 65%. Induction of anesthesia was also attempted with 65 and  $30\% \text{ CO}_2$  in combination with either  $O_2$  or air.

<sup>&</sup>lt;sup>\*</sup>Chemical Supply Officer, New Cumberland General Depot, New Cumberland, Pa.

<sup>\*\*</sup> Manual R-466, Ranarex Instrument Division, The Permutit Company, 50 West 44th Street, New York 36, N.Y., 1957.

Electrocardiographic tracings were made on a Burdick EK III<sup>\*</sup> using a paper speed of 25 mm./second. Only limb leads were used and the recordings were taken before and during CO<sub>2</sub> administration.

Blood pressures were taken directly from the carotid artery which had previously been permanently implanted beneath the skin by surgical procedures. For pressure determinations a 16 gauge  $l\frac{1}{2}$  inch hypodermic needle was inserted into the carotid artery with the point directed towards the heart. This needle was connected to a mercury manometer and the blood pressure was recorded in millimeters of mercury. A series of blood pressure determinations were taken prior to any experimentation in order to establish normal ranges. Blood pressures were also taken during CO<sub>2</sub> administration.

Frequency and depth of respiration were recorded immediately prior to and during CO<sub>2</sub> administration using a Watson-Bird Variable Speed Kymograph 70-142<sup>\*\*</sup> (Figure 6). Kymograph paper speed used for this work was 25 mm/sec.

The hydrogen ion concentration of arterial blood was determined on a Beckman pH meter, Model H-2<sup>\*\*\*</sup> using

<sup>\*</sup>Operating Manual EK III-858 for the Burdick EK III Dual-Speed Electrocardiograph, Aug., 1958.

<sup>\*\*</sup> Tools for Biophysics, Catalog No. 5-60, Phipps and Birds, Inc., Silver Spring, Maryland.

<sup>\*\*\*</sup> Beckman--Bulletin 190-B, Feb., 1948, National Technical Labs, South Pasadena, California.

a special blood electrode assembly (X 800-5) with a flow chamber.

Total erythrocyte and leukocyte counts were made on the improved Neubauer Sharpline Hemocytometer.<sup>\*</sup> The blood smears for differential counts were stained with Wright stain. The packed cell volumes were read from microcapillary tubes.

Rectal temperatures were taken before and during CO<sub>2</sub> administration. The corneal reflexes were tested by touching the cornea with a moistened finger and cutaneous pain reflexes by pricking the skin between the toes with a needle or by pinching with a forceps.

Each calf was subjected to CO<sub>2</sub> administration three times with intervening intervals varying from 2 to 8 days.

During CO<sub>2</sub> administration, a claw amputation was attempted on two calves and on one calf a 3 inch skin incision was made on the lateral surface of the foreleg (over the radius) and sutured.

<sup>\*</sup>Sharpline Hemocytometer, Improved Neubauer Ruling, Chicago Apparatus Co., Chicago, Ill.

Figure 5. Electrocardiograph Burdick EK III

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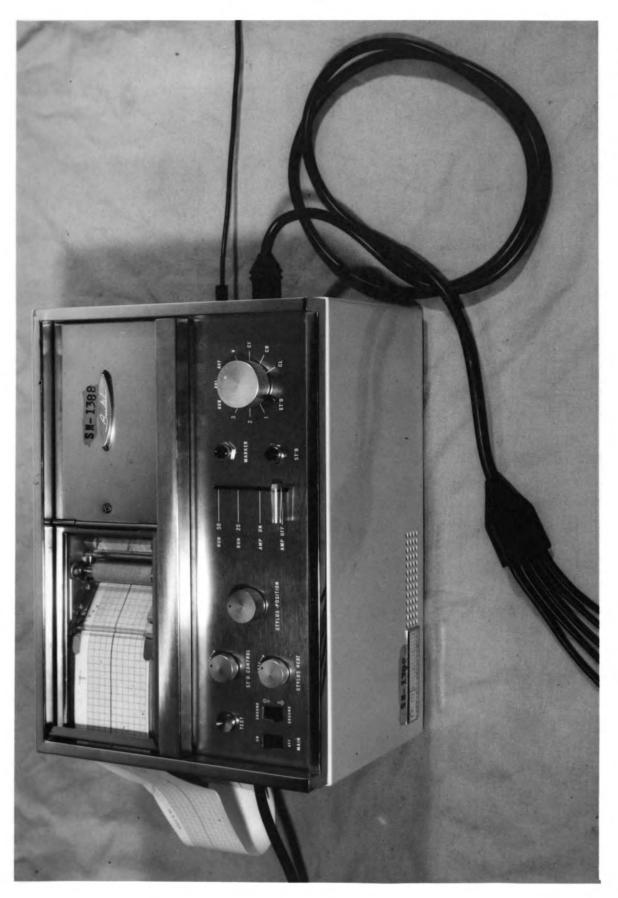
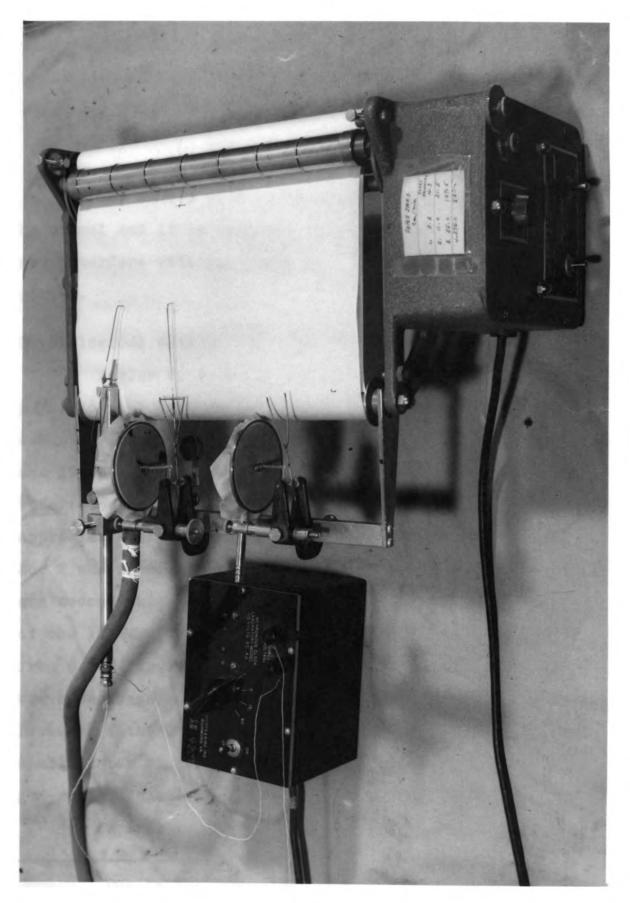


Figure 6. Watson-Bird Variable Speed Kymograph 70-142



### CHAPTER IV

### RESULTS AND DISCUSSION

The calves used in this study were considered to be normal and it is believed that their health status did not interfere with the investigational results herein reported.

### Physiological data

Tables 2, 4, 6, 8, and 10 show the experimental physiological data on each of the five calves. Before  $CO_2$  administration, the rectal temperatures varied between 101.0 and 103.0 F. The temperature of each calf taken 5 minutes before the end of  $CO_2$  administration did not reveal any appreciable elevation. However, the temperature of calf No. 2 was increased during the first (102.6 F. to 104.8 F.) and second experiments (102.4 F. to 103.8 F.). During 7 of the 15, there was an increase of temperature ranging from 0.1 F. to 0.7 F. In the remaining experiments, there was a decrease of temperature varying from 0.2 F. to 1.2 F. Because of the variations of temperatures observed and the small number of animals used, it is difficult to make a general statement regarding the effect of  $CO_2$ .

Superficial observation of the pneumograms (Figs.

<sup>\*</sup>Due to the similarity of the pneumograms only one is presented for each calf (recorded during first experiment).

7, 8, 9, 10, 11) might lead one to believe that the respiratory rates during  $CO_2$  anesthesia were increased. But actually, the rates were often the same as before  $CO_2$  administration. However, there was a marked increase in the depth of the respiratory movement. Best and Taylor (1961) state that an excess of  $CO_2$  in circulating blood depresses the respiratory center resulting in a decrease of respiratory rate. Before the  $CO_2$  administration, the respiratory frequency varied between 20 and 60 per minute; during  $CO_2$  administration it varied between 36 and 60. The most significant increase in respiratory rate was seen on calf No. 4 (Table 8, Fig. 10) where there was an increase from 20 to 44 during second experiment and from 28 to 40 during the third experiment. Also Calf No. 5 showed an increase from 24 to 48 during the second experiment.

The most significant findings of the physiological data were those pertaining to the heart rate, blood pressure, and blood pH during  $CO_2$  administration. The heart rate on each animal increased during  $CO_2$  administration. This increase varied from 40 to 120 beats per minute. The most marked increase was on Calf No. 2 (Table 4 and Fig. 12) during the first experiment (from 80 to 200). The increased heart rate is mainly due to the effect of  $CO_2$  upon cardiac and vasomotor centers, and directly on the cardio-vascular musculature (Best and Taylor, 1961).

The pH of venous blood decreased on each animal

during  $CO_2$  administration. This was apparently the result of increased amounts of carbonic acid in the blood. This fall varied between 0.3 and 1.1. Fertaining to this decrease, it is noteworthy to state that the junctional tissues are particularly sensitive to high tensions of  $CO_2$  and auriculoventricular conduction becomes markedly depressed when the  $CO_2$  excess is sufficient to cause a fall in pH of the fluids bathing the cardiac muscle fibers. At a pH of around 7.0 complete heart block occurs in human beings (Best and Taylor, 1961). The excessive fall in blood pH is probably a disadvantage to the use of  $CO_2$  in calves as a surgical anesthetic. However, there did not appear to be any serious toxic effect during the experiments.

The blood pressures also increased during experiments on each animal. According to Best and Taylor (1961) this increase is a result of high CO<sub>2</sub> tension in venous blood coming into the heart, thus enhancing the extensibility of the cardiac muscle fiber during diastole, and in consequence, exerts a more favorable effect upon the filling of the heart. The cardiac output is therefore increased and the hypertonicity of the vasoconstrictor center results in a high blood pressure. The highest increase was from 170 to 282 mm. Hg. on Calf No. 2 (Table 4) during the third experiment, and the smallest increase was on Calf No. 5 (Table 10) during the third experiment (from 157 to 170 mm. Hg.). In the

remaining experiments, the blood pressure increase ranged from 21 to 84 mm. Hg. above the recordings taken prior to  $CO_2$  administration.

The corneal reflex was present on each animal with the exception of partial abolishment on Calf No. 1 during the first and second experiments. The cutaneous pain reflexes were abolished on Calf No. 1 during first and second experiments. During the balance of the experiments, cutaneous pain reflexes were either partially abolished (forelegs or hindlegs), or were present.

### Hematological findings

The results of the hematological findings are presented in Tables 1, 3, 5, 7, and 9. The hematocrit (Hct.) values varied between 27 and 36.5% in all calves before  $CO_2$ administration. During  $CO_2$  administration in the first trials on each animal there was an increase in REC counts varying from 290,000 to 3,840,000/mm<sup>3</sup>. As an example, in Calf No. 5 (Table 9), the Hct. value remained constant during the experiment but the HEC count increased from 8,190,000 to 10,380,000. This picture may indicate that a new supply of HEC's entered the blood stream but that they were apparently smaller in size. A similar observation was made on Calf No. 3 (Table 5) which also shows an increase of REC's from 7,020,000 to 10,860,000 with only a slight increase in Hct. (27.5 to 29\%). In the remainder of the experiments there was no consistent increase or decrease in REC counts.

The WBC counts before  $CO_2$  administration varied between 5,700 and 13,200/mm<sup>3</sup>. During  $CO_2$  administration, the counts ranged from 5.760 to 13.000/mm<sup>3</sup>. The foregoing WBC counts are generally considered to be the normal values. Inconsistent variations in WBC counts may be noted before and during  $CO_2$  administration.

## Electrocardiographic findings

The electrocardiograms<sup>\*</sup> (Figs. 11, 12, 13, 14, and 15) were taken with the calves in the standing position before starting the experiments and in a recumbent position during  $CO_2$  administration. All recordings were from Lead I.

In this study with calves in the standing position P waves were positive on Calf Nos. 1, 2, 3, and 4 (Figs. 11, 12, 13, and 14) and diphasic on Calf No. 5 (Fig. 15). As is known, the P wave is variable in dairy calves, at times diphasic in Lead I and sometimes inverted (Ardsel, Kreger, and Bogart, 1959), or positive (Platner, Kibler, and Brody, 1948). The PR interval varied between 0.132 and 0.182 seconds in all calves. These data are similar to the results obtained by Platner, Kibler, and Brody (1948) who reported intervals of 0.133 to 0.167 seconds.

During CO<sub>2</sub> administration with the calves in left lateral recumbency there was a significant change in the P

Due to the similarity of electrocardiographic recordings only one is presented for each calf (recorded during first experiment).

wave and the PR interval in all calves as follows:

<u>Calf Nos. 1 and 2</u> (Figs. 11 and 12, Tables 13 and 14). In Calf No. 1 the T wave was superimposed on the P wave whereas the P wave almost disappeared on Calf No. 2. The PR interval showed a variation between 0.14 and 0.16 seconds and widened on Calf No. 1. This picture together with the results on Calf No. 2 may indicate a possible current injury of heart muscle. Both recordings present evidence of tachycardia.

<u>Calf No. 3</u> (Fig. 13 and Table 15). The P wave for this calf is superimposed by the T wave, and there is an increase (up to 0.20 sec.) in the PR interval. This increase in the PR interval is sufficient to cause one beat to drop out. This picture reveals a special type of partial AV block, sometimes referred to in man as the Wenchebach phenomena. It occurs when the PR interval increases in length with each successive cardiac cycle until the impulse, originating in the SA node, is completely blocked at the AV node and fails to initiate a QRS complex (Burch and Winsor, 1960).

<u>Calf Nos. 4 and 5</u> (Figs. 14 and 15, Tables 16 and 17). The P wave on Calf No. 4 was absent. On Calf No. 5 a pre-anesthetic diphasic P wave changed to a monophasic wave with a positive deflection during  $CO_2$  administration. Also, the PR interval increased on Calf No. 5 (0.20 to 0.22 sec.).

Before CO<sub>2</sub> administration the QRS complex was diphasic on Calf Nos. 1, 2, and 4 (Figs. 11, 12, and 14; Tables 13, 14, and 16), and positive on Calf Nos. 3 and 5 (Figs. 13 and 15; Tables 15 and 17). Arsel III, Krueger and Bogart (1959) reported similar findings. The QRS interval showed a variation between 0.04 and 0.072 second which is similar to the results reported by Platner, Kibler and Brody (1948).

During CO<sub>2</sub> administration, the QRS complex showed a negative deflection and amplitudes were increased on all calves. During the same time the QRS interval increased during each experiment ranging from 0.01 to 0.028 seconds with one exception--Calf No. 5--during the second experiment in which there was no change in the QRS interval.

Prior to CO<sub>2</sub> anesthesia the T wave was positive on Calf No. 1 (Fig. 11), and negative on Calf Nos. 2, 4, and 5 (Figs. 12, 14, and 15), and there was no T wave on Calf No. 3 (Fig. 13). During the same time the QT interval showed a variation of 0.28 to 0.36 seconds which resembled the results obtained by Platner and his colleagues (1948).

Observations on anesthetic effect of CO2

As was pointed out in Chapter II, 100%  $CO_2$  has been used to put calves down in lateral recumbency. Near the beginning of each  $CO_2$  administration most calves coughed once or twice and became recumbent within 25 to 37 seconds. There was an absence of struggling in Calves No. 1, 2, 3,

and 4 with the following exceptions: Calves No. 2 and 4 during the second experiment struggled intermittently. Calf No. 5 struggled intermittently during each of the three experiments.

During this series of 15 CO<sub>2</sub> administrations the calves occasionally emitted a slight bellowing, passed feces, and showed some involuntary movements of limbs. At the end of some of the experiments there were blood stains mixed with nasal discharge in the inhalation mask.

In an evaluation of the anesthetic effect of 65% $CO_2$  and 35%  $O_2$  for surgical procedures, a 12 cm. skin incision was made on the forelimb over the radius (lateral surface) of Calf No. 4. The incision was made and suturing was accomplished without the calf showing any evidence of pain.

Two attempts were made to perform a claw amputation operation on two calves while under the influence of 65% $CO_2$  and 35%  $O_2$ . There was not sufficient desensitization of the interdigital skin to permit this surgery.

		<u></u>			I	fferent	ial Cou	int <sup>*</sup>
E	xperiment	Hct. %	RBC 106	WBC 103	Neut. %	Lymph. %	Mono. %	Eosin. %
7 ~ +	Before	27.0	6.52	6.55	25	67	3	5
lst	During**	28.0	6.81	7.00	4	89	4	3
0	Before	29.0	6.47	7.15	8	88	4	0
2nd	During	34.5	8.56	8.30	6	91	l	2
<b>Z</b> m d	Before	35.0	6.65	5.70	18	79	2	ı.
3rd	During	36.5	6.65	5.76	21	74	2	3

Table 1. Hematological findings, Calf No. 1

Table 2. Physiological data, Calf No. 1

I	Experiment	Duration** /min.	*Temp. F.	Resp. rate /min.	Heart rate /min.	pres	• pH	<u>Reflex</u> Cornea	
7 ~ 4	Before		102.4	52	100	166	7.1		
lst	During**	17	102.5	38	160	200	6.6	PA	A
2nd	Before		102.4	36	100	166	6.9	)	
2110	During	20	102.6	40	150	210	6.6	PA	A
3rð	Before		101.1	52	80	166	7.4		
	During	45	101.8	36	120	250	6.4	Р	PA

\*Basophils were not observed on the blood smears. \*\*Taken 5 minutes before the end of CO<sub>2</sub> administration. \*\*\*Duration of CO<sub>2</sub> administration.

## Legend:

A--Abolished; PA--Partially abolished; P--Present.

				Differential Count*				
Experiment	Hct. %	RBC 106	WBC 103	Neut. %	Lymph. %	llono. %	Eosin.	
Before	29.0	5.78	8.35	25	72	2	l	
lst During**	30.0	7.11	12.55	36	60	4	0	
Before	27.0	6.65	6.60	20	<b>7</b> 8	l	l	
2nd During	27.0	6.35	10.45	18	82	0	0	
Before	29.0	6.45	6.30	26	70	4	0	
3rd During	31.0	7.10	6.73	28	68	4	0	

Table 3. Hematological findings, Calf No. 2

Table 4. Physiological data, Calf No. 2

	Experiment	Duration** /min.	*Temp. F.	Resp. rate /min.	rate	Blood pres. mm. H	рH	Reflexe Cornea	
7	Before		102.6	42	80	170	7.5		
Ŧ	st During	21	104.8	44	200	212	6.6	P	PA
2	Before		102.4	60	80	170	6.9	1	
2	nd During	30	103.8	48	120	194	6.5	P	Р
7	Before rd		101.6	52	80	170	7.3		
2	During	27	101.2	56	110	282	5.6	P	PA

\*Basophils were not observed on the blood smears. \*\*Taken 5 minutes before the end of CO<sub>2</sub> administration. \*\*\*Duration of CO<sub>2</sub> administration.

# Legend:

P--Present; PA--Partially abolished.

					Differen	tial Co	unt*
Experiment	Hct. %	RBC 10 <sup>6</sup>	WBC 103	Neut. %	Lymph. %	Mono.	Eosin. %
Before	27.5	7.02	13.20	42	53	1	4
lst During**	29.0	10.86	10.10	49	41	1	9
Before	28.0	6.16	10.60	19	77	3	l
2nd During	27.0	5.26	13.00	12	84	2	2
Before	32.0	7.06	11.25	44	55	1	0
3rd During	32.0	6.40	11.45	40	55	3	2

Table 5. Hematological findings, Calf No. 3

Table 6. Physiological data, Calf No. 3

I	Experiment	Duration <sup>**</sup>	"Temp. F.	Resp. rate /min.	Heart rate /min.	pres	• pH	<u>Reflexe</u> Cornea	
	Before		102.4	44	60	178	7.2	2	
ls	During	30	102.2	44	90	223	6.6	P	PA
2~	Before		102.6	32	60	178	7.1		
2r	During	22	103.0	52	100	212	6.5	5 P	PR
2-	Before		102.0	40	60	178	7.3	5	
31	During	27	102.6	<b>3</b> 8	120	250	6.2	2 P	PR

\*Basophils were not observed on the blood smears. \*\*Taken 5 minutes before the end of CO<sub>2</sub> administration. \*\*\*Duration of CO<sub>2</sub> administration.

Legend:

P--Present; PA--Partially abolished; PR--Partially abolished rear legs.

					Differen	tial Co	unt*
Experiment	Hct. %	RBC 106	WBC 103	Neut. %	Lymph. %	iiono.	Eosin. %
Before	35.0	8.19	10.80	8	91	0	l
lst During**	36.5	10.38	7.90	11	87	1	l
Before	28.0	7.70	9.10	24	76	0	0
2nd During	35.0	13.35	8.14	24	76	0	0
Before	32.0	8.08	11.45	16	82	2	0
3rd During	33.0	8.09	11.25	25	72	2	0

Table 7. Hematological findings, Calf No. 4

Table 8. Physiological data, Calf No. 4

E	xperiment	Duration** /min.	*Temp. F.	Resp. rate /min.	Heart rate /min.	<u>Blood</u> pres. mm. H	pH	Reflexe Cornea	
٦ -	Before		102.4	36	90	174	7.7		
ls	During	42	101.2	52	140	224	7.0	P	PA
2n	Before		102.6	24	96	174	7.5		
211	During	25	102.4	48	148	210	6.7	P	P
3 m	Before		101.8	28	90	174	7.3		
3r	During	18	101.2	32	140	220	7.0	Р	PF

\*Basophils were not observed on the blood smears. \*\*Taken 5 minutes before the end of CO<sub>2</sub> administration. \*\*\*Duration of CO<sub>2</sub> administration.

# Legend:

P--Present; PA--Partially abolished; PF--Partially abolished forelegs.

						Differen	tial Co	unt*
Ex	periment	Hct. %	RBC 106	WBC 103	Neut. %	Lymph.	Mono.	Eosin. %
1 ~ +	Before	36.5	8.19	10.80	8	91	0	1
lst	, During**	36.5	10.38	7.90	10	89	l	0
0	Before	30.5	9.49	10.20	13	84	2	1
2nd	l During	31.0	9.77	9.40	10	86	3	l
7	Before	30.0	8.25	10.50	18	78	2	2
3rd	During	32.0	10.30	8.45	23	<b>7</b> 5	2	0

Table 9. Hematological findings, Calf No. 5

Table 10. Physiological data, Calf No. 5

E	xperiment	Duration** /min.	*Temp. F.	Resp. rate /min.	rate	pres.	pH	<u>Reflexe</u> Cornea	
] -	Before		102.0	52	70	15 <b>7</b>	7.6		
ls	During	40	101.0	60	140	210	7.1	P	Р
2n	Before		101.8	20	100	157	7.6		
211	During	15	101.6	44	144	178	6.9	P	Р
3r	Before		101.6	28	70	157	7.4		
1	During	23	101.4	40	130	170	6.7	P	Р

\*Basophils were not observed on the blood smears. \*\*Taken 5 minutes before the end of CO<sub>2</sub> administration. \*\*\*Duration of CO<sub>2</sub> administration.

Legend:

P--Present.

Ē	Experiment	PR interval sec.	QRS interval sec.	QT interval sec.
	Before	0.14	0.072	0.28
lst	During	0.16	0.080	0.22
0	Before	0.15	0.048	0.28
2nd	During	0.16	0.080	0.23
7	Before	0.16	0.072	0.32
3rd	During	0.16	0.080	0.24

Table 11. Electrocardiographic findings, Calf No. 1

Table 12. Electrocardiographic findings, Calf No. 2

E	xperiment	PR interval sec.	QRS interval sec.	QT interval sec.
7 - +	Before	0.16	0.040	0.35
lst	During	No P wave	0.052	0.20
2nd	Before	0.14	0.044	0.32
Znu	During	No P wave	0.072	0.24
3rd	Before	0.16	0.040	0.30
	During	No P wave	0.064	0.20

Experiment		PR interval sec.	QRS interval sec.	QT interval sec.
7	Before	0.180	0.040	No T wave
lst	During	0.13-0.20	0.080	0.28
2nd	Before	0.182	0.040	No T wave
	During	0.13-0.20	0.070	0.28
3rd	Before	0.180	0.040	No T wave
	During	No P wave	0.078	0.30

Table 13. Electrocardiographic findings, Calf No. 3

Table 14. Electrocardiographic findings, Calf No. 4

Experiment		PR interval sec.	QRS interval sec.	QT interval sec.
lst	Before	0.160	0.060	0.36
	During	No P wave	0.070	0.28
2nd	Before	0.160	0.048	0.32
	During	Not recorded*		
3rd	Before	0.168	0.048	0.35
	During	No P wave	0.080	0.28

\*Because of the movements of calf.

Experiment		PR interval sec.	QRS interval sec.	QT interval sec.
lst	Before	0.140	0.080	0.328
	During	0.220	0.128	0.360
2nd	Before	0.132	0.080	0.320
	During	0.200	0.080	0.360
3rd	Before	0.128	0.080	0.340
	During	Not recorded*		

Table 15. Electrocardiographic findings, Calf No. 5

\*Because of the movement of calf.

Figure 7. Pneumogram, Calf No. 1

- A. Before  $CO_2$  administration
- B. During  $CO_2$  administration
- C. Time interval 1 sec.

Figure 8. Pneumogram, Calf No. 2

- A. Eefore CO<sub>2</sub> administration
- B. During  $CO_2$  administration
- C. Time interval 1 sec.

Figure 9. Pneumogram, Calf No. 3

- A. Before  $CO_2$  administration
- B. During  $CO_2$  administration
- C. Time interval 1 sec.

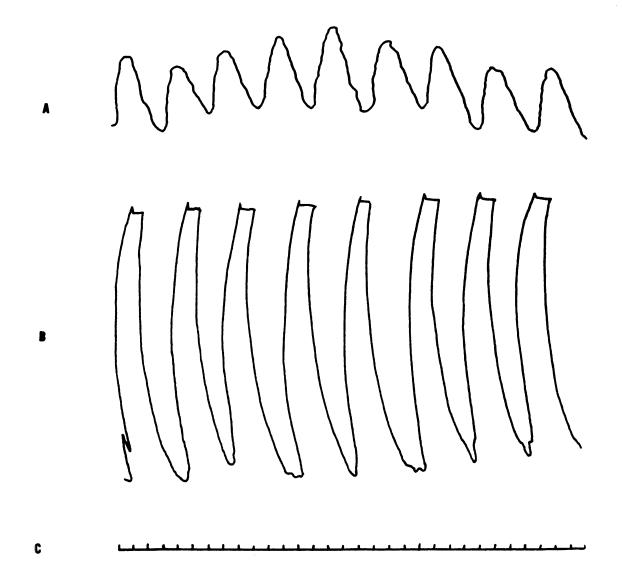


Figure 10. Pneumogram, Calf No. 4

- A. Before  $CO_2$  administration
- B. During CO<sub>2</sub> administration
- C. Time interval 1 sec.

Figure 11. Pneumogram, Calf No. 5

- A. Before CO<sub>2</sub> administration
- B. During  $CO_2$  administration
- C. Time interval 1 sec.

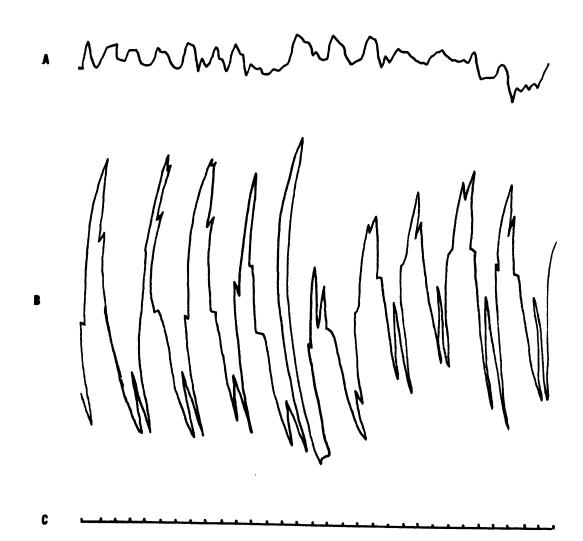


Figure 12. Electrocardiogram, Calf No. 1

A. Before CO<sub>2</sub> administration
B. During CO<sub>2</sub> administration
(Five large squares represent a time interval of 1 sec.)

,

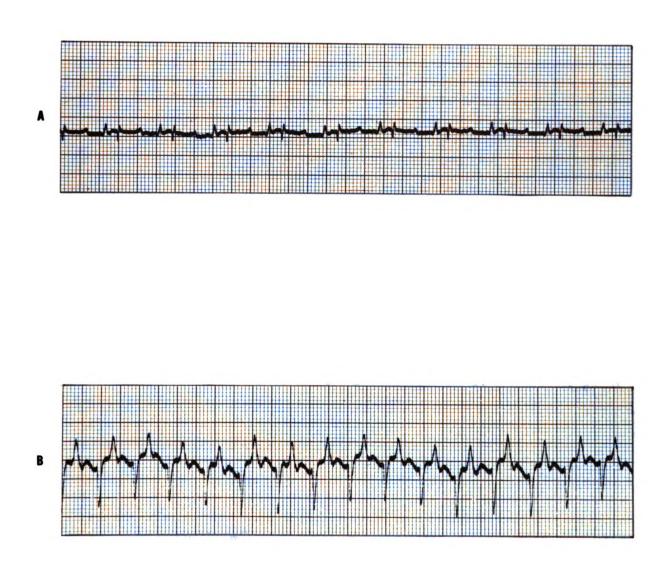


Figure 13. Electrocardiogram, Calf No. 2

A. Before  $CO_2$  administration

B. During  $CO_2$  administration

(Five large squares represent a time interval of 1 sec.)

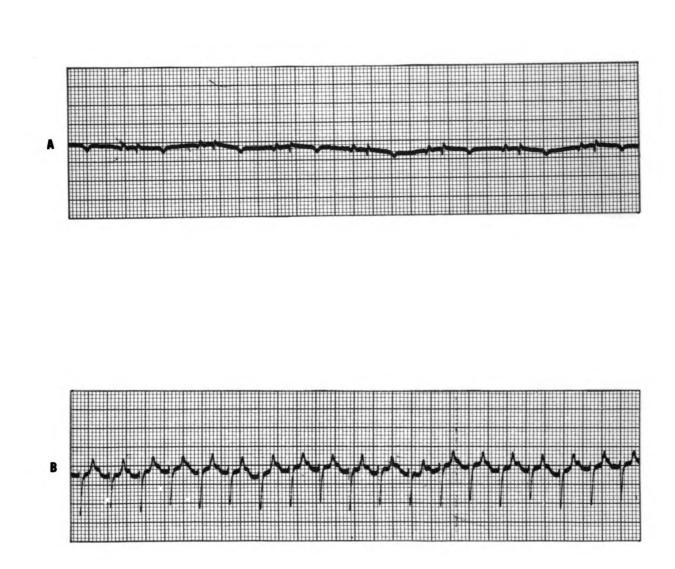


Figure 14. Electrocardiogram, Calf No. 3

A. Before CO<sub>2</sub> administration
B. During CO<sub>2</sub> administration
(Five large squares represent a time interval of 1 sec.)

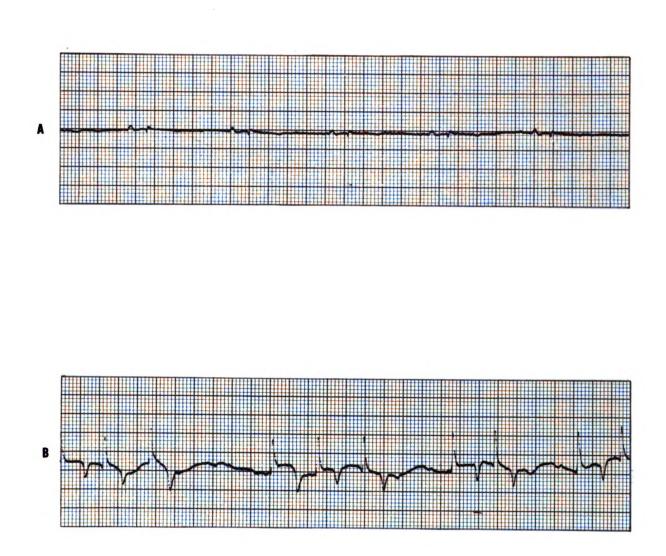


Figure 15. Electrocardiogram, Calf No. 4

A. Before CO<sub>2</sub> administration
B. During CO<sub>2</sub> administration
(Five large squares represent a time interval of 1 sec.)

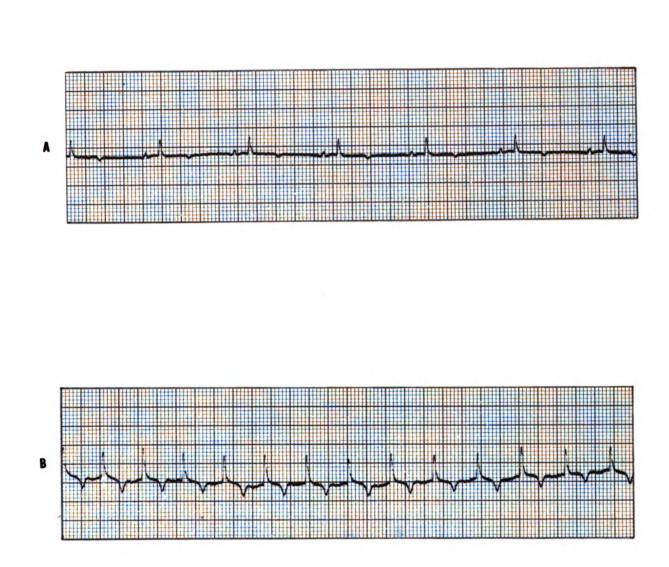
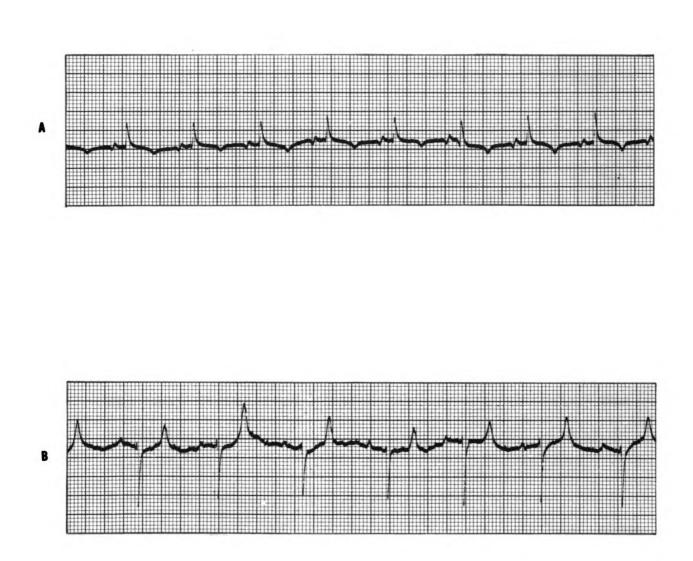


Figure 16. Electrocardiogram, Calf No. 5
A. Before CO<sub>2</sub> administration
B. During CO<sub>2</sub> administration
(Five large squares represent a time interval
of l sec.)



#### CHAPTER V

### SUMMARY

- 1. During the administration of 65% CO<sub>2</sub> and 35% O<sub>2</sub> to calves, there was a consistent decrease of blood pH varying between 0.3 and 1.1.
- During all CO<sub>2</sub> administrations, there was a definite increase in blood pressure ranging from 13 to 112 mm. Hg.
- 3. The heart rate showed an increase between 30 and 120 beats per minute.
- 4. The hematological studies conducted during CO<sub>2</sub> administration revealed only minor fluctuations from the values determined before the administration of CO<sub>2</sub>. The most significant fluctuation was noted during the first CO<sub>2</sub> experiments on each calf where there was an increase in the RBC count varying from 290,000 to 3,800,000 cells per cubic millimeter.
- 5. As evidenced by the electrocardiographic studies, there appeared to be a significant change in the conduction system of the heart during CO<sub>2</sub> administration. Tachy-cardia, possible heart injury, and on one calf (No. 3; Fig. 13), the Wenchebach phenomena (partial AV block), were observed on the electrocardiograms.

- 6. During CO<sub>2</sub> anesthesia, the depth of respiratory movements increased but the rate did not appreciably change.
- 7. The corneal reflex was present during CO<sub>2</sub> anesthesia. Cutaneous pain reflexes were sometimes absent, at other times present, especially in the interdigital areas.
- 8. Rectal temperatures did not reveal any consistent decrease or increase during CO<sub>2</sub> administration.
- 9. The nasal discharges of 3 calves contained small amounts of blood noticed upon completion of CO<sub>2</sub> administration.
- 10. Calves would not fall into lateral recumbency when  $CO_2$  was administered in concentrations of 65 or 80%, in combination with either  $O_2$  or air.

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