CHLORAL HYDRATE FOR GENERAL ANAESTHESIA IN GOATS

BY

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THESIS

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DECLARATION

"CHLORAL HYDRATE FOR GENERAL ANAESTRESIA IN GOATS" is a benefide record of research work done by me during the course of research and that the thesis has not previously formed the basis for the award to me of any degree, diploma, associateship, fellowship, or other similar title, of any other University or Society.

Hannathy, 31-7-1981. RUYSwandhan R. VISWANATIAN

CERTIFICATE

"CHLORAL HYDRATE FOR GENERAL ANAESTHESIA IN GOATS" is a record of research work done independently by Sri. R. Visuamathan under my guidance and supervision and that it has not previously formed the basis for the award of any degree, fellowship, or associateship to him.

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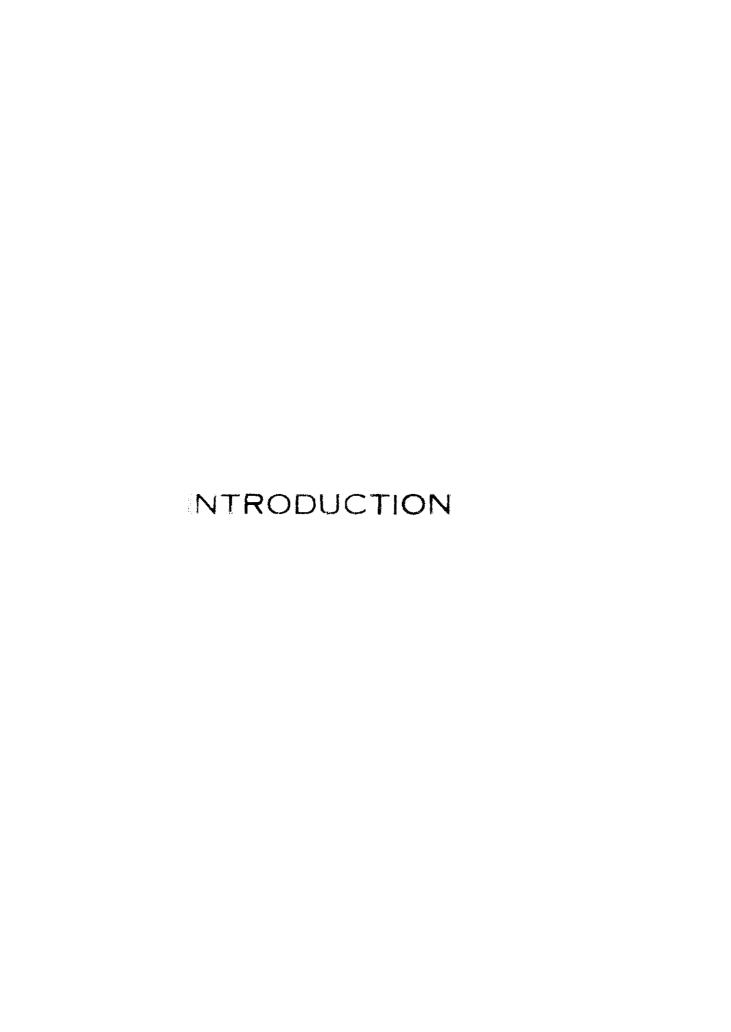
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INTRODUCTION

Veterinarians have to face the problem of assessmentialing enimals of different species and temperature. Variations assenget individuals are also semarkable. General assessments has been practiced in animals from assistant days. The advent of modern assessmentic techniques dates back to the last century. Humbert (1875) was the first to use chloral hydrate intravenessly in horses as a general assessment and sarcotic (cited by Hall, 1971 and Jones et al. 1978).

The pharmoological properties of chloral hydrate closely resemble that of trichlorosthanel and trichloroscetic scid. Trichlorosthanel is exercted partly through kidney and partly after it is conjugated in liver mainly with glucuromic scid to form urochloralic scid.

Chloral hydrate is a good hypnotic, but
a poor amesthetic. In hypnotic doses, medulary
centres are not effected but in amesthetic doses
it depresses the respiratory centre and the
vacomotor centre resulting in fall of blood pressure

(Lumb and Jones, 1973).

Chieral hydrate alone or in combination with magnesium sulphate and Newbutel had been extensively used as a general amountate in large animal practice. Chieral hydrate is administered often in goats orally, intravenously and as a retention enema. Movever, a detailed systematic study on the use of intravenous administration of chieral hydrate at different dose levels, used alone or in combination with pre-amaesthetics like Triflupromagine hydrochloride, appears to be necessary to recommend its use in clinical practice, and hence the present study.



REVIEW OF LITERATURE

Chlorel hydrate was used as a narcotic and anaesthetic in horses by Humbert as early as 1875 and was first given by intravenously at a dose of 30 to 30 grammes. Degive (1908) made use of a 20 per cent solution of chloral hydrate for intravenous injection in horses (cited by Hall, 1971 and Jones 185 al. 1978).

while (1962) had reported different doses of chloral hydrate for intravenous injection in calves, vis. a mean dose of 183.3 mg per kg bodyweight in 19 calves weighing 45.5 to 90.5 kg and a mean dose of 201.7 mg per kg bodyweight in six calves weighing 90.6 to 108.0 kg.

chlorn, hydrate in a dose of 58 to 125 mg per 1b bodyweight produced anaesthesis lasting for 16 to 50 minutes. for anaesthetising easel, Margava and Vyas

(1967) recommended a combination of chloral hydrate
and magnesium sulphate each 12 per cent at a dose
of 12 per 220 lb bedyweight.

cent solution of chloral hydrate at the rate of 0.08 to 0.1 g per kg bodyweight when administered intravenously after atropine sulphate or neuroleptics, produced narcosis within a few minutes and lasted for about half an hour. The animal recovered in one to two hours.

chloral hydrate and magnesium sulphate in a proportion of 2:1, as general anaesthetic in 156 goats of different age groups. They observed that the duration of anaesthesia varied from 30 to 90 minutes with a recovery period of five to fifteen minutes. They also observed that 13 animals which were prognant during the experimental study, delivered at full term.

Chandna et al. (1970) observed that in calves, chloral hydrate at the rate of 106 mg per kg bodyweight, showed a mean recovery time of 70 minutes. It was also

reduced to 75 mg per kg bodyweight, when used in combination with a preamesthetic. They also observed that there was significant acceleration in respiration and pulse rate but no significant variation in blood pressure. The recovery was prolonged by 10 to 14 minutes. The average duration of light amesthesis was 30 minutes.

Hail (1971) reported that chloral hydrate
at 20 per cent solution administered intravenously
was satisfactory for anaesthetising horses. The
dose of intravenous administration vary extensively,
probably due to the variation with the rate of
administration and other connected factors.

Hamsa (1971) recommended a mixture of 30 grams of chloral hydrate and seven grammes of Magnesium sulphate in 420 ml of distilled water or physiological saline for general anaesthesis in horses.

Kumar et al. (1971) made a comparable evaluation of chloromag with and without pentobarbital sodium for inducing general anaesthesia in dogs and observed

that chloromag pentobarbital sodium combination was more effective than chloromag alone.

Kumar et al. (1971) made use of a solution containing 21 grams of chloral hydrate and 10.5 grammes of magnesium sulphate in 500 ml distilled water for general amaesthesia in dogs and found that the average quantity of the solution required to produce surgical amaesthesia was 7.7, 5.7, 3.6 and 5.5 ml per kg bodyweight respectively in controls, and when methodone, fluphenssine and triflupromasine hydrochloride were used as pre-medicaments. They did not observe any untoward reaction and the duration of surgical amaesthesia lasted for 22 to 45 minutes.

Singh et al. (1971) reported that, in buffaloes, the doses of chloromag solution containing 4.2 per cent chloral hydrate and 2 per cent magnesium sulphate, can be reduced from 2.1 ml per kg to 1.6 ml per kg bedyweight when premedicated with 0.1 mg per kg chloropromatine hydrochloride or 1 mg per kg triflupromatine hydrochloride. The dose can also be reduced to 1.8 ml per kg bodyweight when promethazine hydrochloride at the rate of 0.1 mg per kg was used as premedicant, but it produced sudden fall in blood pressure.

Johani et al. (1972) used chloral and thiopenton sodium mixture intravenously for ansesthesia in buffalo calves and observed that administration of atropine sulphate 3 to 13 mg subcutaneously 30 minutes before inducing amesthesia prevented excessive salivation and regurgitation.

Singh at al. (1972) administered 12 per cent solution of chloral hydrate as general anaesthetic at the rate of 20 ml per 50 kg bodyweight in cattle, which were starved overnight. These animals recovered from aspecthesis in 15 to 20 minutes. There was increase in glucose and serum level in blood samples taken 10 minutes after the injection of a mesthetic.

Sharm at al. (1973) had reported an increase in erythrocyte count, has moglobin content and a decrease in mean corpuscular has moglobin content and colour index, during chloromag or other amesthesis in experimental dogs. These changes returned to normally after recovery. They further observed that chloromag anaesthesis produced leucopenis.

Rebesio and Mehamood (1974) used a mixture of 10 per cent solution of chloral hydrate and magnesium sulphate at the rate of 2.5 ml per kg bodyweight in twenty rams. They observed that, the mixture produced satisfactory surgical assessment with only a slight transient phase of excitement. The solution did not produce any serious depresent effect in heart rate and respiration.

Singh gt al. (1974) had reported a decrease in total crythrocyte and hasmoglobin content and an increase in leucocyte count during chloromag sedation in bovines. Increase in serum calcium and blood glucose concentration was reported by them.

Major et al. (1978) reported a significant fell in hadmoglobin content and packed cell volume along with an increase in leucocyte count following chloral hydrete anaesthesia in buffalo calves.

changes at al. (1978) made use of a mixture of chiefs hydrate and magnesium sulphate in the ratio of 2:1 in goats and the average dose was found to be 75 ml with a range of 45 - 95 ml and the

duration of anaesthesia was reported to be varying from 30 to 90 minutes and the recovery period varied from 5 to 15 minutes.

Lakshmipathy and Vijyakumar (1980) atudied the effect of Siquil at the rate of 5 mg per 100 lb body-weight reconstituted in 5 ml distilled water as slow intravenous injection in 16 buffalo calves. The rectal temperature showed a gradual fall and in four cases, temperature became sub-normal. There was an initial rise in the rate of respiration and subsequent downward trend from 18.19 a 1.69 to 13.38 \(\pm\) 1.37 at 30 minutes interval. A gradual fall in pulse rate was also recorded.



MATERIALS AND METHODS

The experimental study was conducted on 36 apparently healthy, Alpine - Malabari cross-bred bucks, aged from 17 to 30 months and weighing from 22 to 39.5 kg.

The animals were kept under observation for two weeks and treated for ecto and endo parasites, before the experiment.

These animals were divided into two groups,
vis., Group I and II, each group consisting of 18
animals. Group I and II were further subdivided
into three subgroups vis., A, B and C, each subgroups
consisting of six animals. These animals were numbered serially from 1 to 6 vis:

Group I A-1, A-2, A-3, A-4, A-5 and A-6; I B-1, B-2, B-3, B-4, B-5 and B-6; I C-1, C-2, C-3, C-4, C-5 and C-6;

Group II A-1, A-2, A-3, A-4, A-5 and A-6; II B-1, B-2, B-3, B-4, B-5 and B-6; II C-1, C-2, C-3, C-4, C-5 and C-6.

Preparation of animals.

The animals were fasted over night and were weighed before the administration of the amesthetic. The site

was prepared by elipping the hairs and painting Tr. of Todine.

In Group I, freshly prepared chloral hydrate six per cent solution was administered intravenously at the rate of

- (i) 1 ml per kg bedyweight in I A,
- (ii) 1.5 ml per kg bedyweight in I B and
- (dii) 2 ml per kg body weight in I C.

In Group II, triflupromasine hydrochloride (Siquil*) followed by freshly prepared chloral hydrate six per cent solution were administered intraveneusly at the rate of

- (i) 0.2 mg per kg bedyweight of triflupromasine hydrochloride followed by 1 ml per kg bedyweight of chloral hydrate six per cent solution in IN A.
- (11) 0.2 mg per kg bodyweight of triflupromasine hydrochloride followed by 1.5 ml per kg bodyweight of chloral hydrate six per cent solution in
 II B and
- (111) 0.2 mg per kg bodyweight of triflupromanine hydrochloride followed by 2 ml per kg bodyweight

Siguil (Vety.) a product of Sarabhai Chemicals
Private Ltd., containing 20 mg of Triflupromazine
hydrochloride in 1 ml of the solution.

of chieral hydrate six per cent solution in II C.
Technique.

The animal was controlled in the standing position. The pulse, temperature and respiration were recorded before the administration of the amaesthetic. The jugular vein was raised and a hypodermic needle, 18 B.V.G. was introduced into the jugular vein and five millilitres of blood was collected into a vial containing EDTA (Ethylene diamine tetra acetate) for haematological studies. Through the same needle, in Group I, chloral hydrate six per cent solution and in Group II, triflupromazine hydrochloride solution followed by chloral hydrate six per cent solution were administered as slow intraveneus injection.

The bleed samples were collected again after fifteen minutes of administration of the amesthetic.

The following haematological values were studied:

- 1. total erythrocytes,
- 11. total leucocytes.
- iii. Haemoglobin gm percentage and
 - iv. Packed Cell Volume percentage.

A haem cytometer was used for the estimation of the total erythrosyte and leucocyte counts.

For erythrocyte count, Hayem's fluid and for leucocyte count, Thomas fluid were used as the diluent, Haemoglobin estimation was done using Sahl's acid Haematin method making use of a haemoglobinometer. The packed cell volume estimation was done by a Vintrobe haematocrit.

During the experiment, the following observation were made

- i) Temperature, pulse and respiration before administration of amesthetic and at five minutes interval after administration of amesthetic till the 20th minute.
 - ii) Volume of annesthetic administered.
 - iii) Time taken for intravenous injection.
 - iv) Time taken for incoordination of limbs.
 - v) Time taken for disappearance of palpebral reflex.
 - vi) Time taken for assuming recumbency.
 - vii) Time taken for reappearance of palpebral reflex.
 - viii) Duration of recumbency.
 - in) Time taken for complete recovery from annesthesia.

- m) Post amesthetic observations, if any.
- xi) Hasmatological estimations immediately before and fifteen minutes after administration of anaesthetic.



RESULTS

The observations in general with respect to each group of animals are presented in Tables I to 9.

Group I A

The data are presented in Tables 1, 4 and 7.

The average bodyweight of the animals in

this group was 30.92 ± 2.70 kg. Chloral hydrate
at the rate of 1 ml per kg bodyweight as a six

per cent solution was administered intravenously.

The time taken for the intravenous injection was

50.83 ± 7.16 seconds.

The palpebral reflex disappeared by 4.13 ± 1.01 minutes. Incoordination of movements was observed by 1.50 ± 0.29 minutes. Only one animal (No.4) assumed sternal recumbersy by the 11th minute which persisted for four minutes. Palpebral reflex disappeared in four animals while it was present in two animals (No.4 and 6). Incoordination of movements was not observed in two animals (Nos. 2 and 6). Palpebral reflex reappeared by 8.13 ± 2.24 minutes. The animal became apparently normal in gait by 12.06 ± 1.58 minutes.

The temperature, pulse and respiration recorded before the administration of the anaesthetic were $102.87 \pm 0.30^{\circ}$ F, 83.33 ± 2.77 per minute and 18.33 ± 0.80 per minute respectively. The temperature during the 5th, 10th, 15th and 20th minutes after the administration of the anaesthetic was $102.83 \pm 0.41^{\circ}$ F, $102.57 \pm 0.34^{\circ}$ F, $102.67 \pm 0.31^{\circ}$ F and $102.60 \pm 0.31^{\circ}$ F respectively.

The pulse rate during the 5th, 10th, 15th and 20th minutes after the administration of the assestantian was 92.67 ± 2.77 , 91.33 ± 3.49 , 88.00 ± 2.25 and 86.00 ± 3.35 per minute respectively.

The respiration during the 5th, 70th, 15th and 20th minute after administration of the anaesthetic was 21.33 ± 0.80 , 21.33 ± 1.43 , 20.67 ± 1.09 and 18.33 ± 0.62 per minute respectively.

The hassogram recorded before the administration of the ammesthetic was as follows:

REC - 8.98 ± 1.07 million/e.m.m.,

WPC - 7650.00 ± 340.59/e.m.m.,

Hb - 9.83 ± 0.35 6 % and

PCV - 33.33 ± 2.03 %.

The hasmogram recorded fifteen minutes after the administration of chloral hydrate was as follows:

REC - 8.27 + 1.228 million/e.m.m.,

WEC - 8475.00 ± 310.85/c.m.m.,

Hb - 9.37 ± 0.53 g % and

PCV - 30.50 ± 0.563 \$

All the animals started feeding immediately after recovery but two animals were dull for few hours.

Group I B

The data are tabulated and presented in Tables 2. 5 and 8.

The average bedyweight of the animals in this group was 29.58 ± 2.72 kg. Chloral hydrate at the rate of 1.5 ml per kg bodyweight in a six per cent solution was administered intravenously. The time taken for the administration was 55.00 ± 5.92 seconds.

The palpebral reflex disappeared by 1.30 \pm 0.30 minutes. Incoordination of movements was observed by 0.88 \pm 0.13 minutes. Animals except No.1 and 6 became recumbent by 1.13 \pm 0.32 minutes. Of these one

animal (No.4) assumed lateral recumbency, two animals (Nos. 3 and 5) assumed sternal recumbency and then lateral recumbency, while animal No.2 remained in sternal recumbency alone. Salivation was observed in two animals (Nos. 3 and 4), and lacrimation was observed in three animals (Nos. 3, 4 and 5). Dilatation of pupil was noticed only in ome animal (No.2).

Palpebral reflex reappeared by 14.90 \pm 1.38 minutes. The animals were in recumbent position for 14.88 \pm 4.32 minutes. The animals became apparently normal in gait by 20.88 \pm 1.09 minutes.

The temperature, pulse and respiration recorded before the administration of the anaesthetic were $102.47 \pm 0.25^{\circ}$ F, 78.67 ± 2.46 per minute and 16.67 ± 0.67 per minute respectively.

The temperature during the 5th, 10th, 15th and 20th minute after the administration of anaesthetic was $102.77 \pm 0.33^{\circ}$ F, $102.47 \pm 0.27^{\circ}$ F, $102.30 \pm 0.25^{\circ}$ F and $102.37 \pm 0.25^{\circ}$ F respectively.

The pulse rate during the 5th, 10th, 15th and 20th minute after the administration of the amaesthetic

was 98.00 ± 2.25 , 96.33 ± 3.95 , 98.67 ± 3.25 and 99.00 ± 3.38 per minute respectively.

The respiration during the 5th, 10th, 15th and 20th minutes after the administration of the ansesthetic was 21.50 ± 1.34 , 20.00 ± 0.90 , $20.6\% \pm 1.52$ and 19.67 ± 1.49 per minute respectively.

The haemogram recorded before the administration of the anaesthetic was as follows:

RBC - 6.00 ± 0.35 million/c.m.m.,

WEC - 6808.33 + 1260.19/e.m.m.,

Hb - 9.42 + 0.27 g % and

PCV - 30.50 ± 1.67 %

The hasmogram recorded fifteen minutes after the administration of chloral hydrate was as follows:

RBC - 6.03 ± 0.39 million/c.m.m.,

WBC - 8133.33 ± 754.39/c.m.m.,

Hb - 8.20 + 0.38 g % and

PCV - 30.17 + 1.49 \$

All the animals started feeding immediately after recovery.

Group I C

The data are presented in tables 3, 6 and

The average body-veight of the animals in this group was 30.17 ± 2.28 kg. Chloral hydrate at the rate of 2 ml per kg bodyweight as a six per cent solution was administered intravenously. The time taken for the intravenous injection was 78.33 ± 8.75 seconds.

The palpebral reflex disappeared by 3.63 \pm 0.75 minutes and respected by 9.67 \pm 3.51 minutes. The incoordination of movements was observed by 1.33 \pm 0.49 minutes.

All the animals, except Nes. 2 and 6, became recumbent by 2.00 ± 1.08 minutes. Out of these, three animals (Nos. 3, 4 and 5) assumed lateral recumbency, one animal (No.I) assumed sternal recumbency. Dilatation of pupil was noticed in two animals (Nos. 3 and 4).

The period of recumbercy in this group was 12.75 ± 6.63 minutes. The animal became apparently normal in gait by 47.83 ± 12.41 minutes.

The temperature, pulse and respiration recorded before the administration of the anaesthetic were

102.00 ± 0.16°r, 80.50 ± 3.36 per minute and 20.33 ± 1.96 per minute respectively.

The temperature during the 5th, 10th, 15th and 20th minutes after the administration of the anaesthetic was $102.10 \pm 0.10^{\circ}$ F, $101.87 \pm 0.20^{\circ}$ F, $101.97 \pm 0.20^{\circ}$ F and $102.13 \pm 0.28^{\circ}$ F respectively.

The pulse rate during the 5th, 10th, 15th and 20th minutes after the administration of ansesthetic was 104.00 ± 6.12 per minute, 94.67 ± 3.17 , 93.67 ± 2.85 and 90.00 ± 3.46 per minute respectively.

The respiration during the 5th, 10th, 15th and 20th minute after administration of the amesthetic was 22.50 \pm 1.93, 20.66 \pm 0.84, 17.67 \pm 0.61 and 18.67 \pm 0.99 per minute respectively.

The hasmogram recorded before the administration of the anaesthetic was as follows:

RBC - 6.42 + 0.16 million/c,m.m.

WHC - 8250 + 381.88/e.m.m.,

Hb - 9.52 ± 0.20 g % and

PCV - 29.50 ± 1.45 %

The haemogram recorded fifteen minutes after the



administration of chloral hydrate was as follows:

REC - 5.67 + 0.18 million/c.mm.

WEC - 8416.67 + 624.72/0.mm.

Mb - 8.42 + 0.20 g % and

PCV - 28.17 ± 1.28 %.

All the animals started feeding immediately after mecovery.

Group II A

The data are presented in tables 1, 4 and 7.

The average bodyweight of the animals in this
group was 27.33 ± 2.68 kg. Triflupromasine hydrochloride solution at the rate of 0.2 mg per kg
bodyweight followed by chloral hydrate at the rate
of 1 ml per kg bodyweight as a six per cent solution
was administered intravenously. The time taken for
completing injection was 56.00 ± 12.08 seconds.

The pelpabral reflex disappeared by 2.92 ± 0.52 minutes. Incoordination of movements was observed by 4.50 ± 1.60 minutes. Out of the six animals, eally one animal (No.3), assumed the position of sternal recumbency by 10th minute which persisted for one minute. Drowsiness and droping of head were noticed in two animals (No.1 and 4).

Palpebral reflex reappeared by 10.83 ± 1.60 minutes. The animal became apparently normal in gait by 14.17 ± 2.85 minutes.

The temperature, pulse and respiration recorded before the administration of the amesthetic were 102.07 ± 0.32^{0} Y, 84.67 ± 4.46 per minute and 22.50 ± 1.66 per minute respectively.

The temperature during the 5th minute, 10th, 15th and 20th minute after the administration of the amosthetic was 102.33 ± 0.33^{6} Y, 102.13 ± 0.40 , 101.83 ± 0.36 and 101.70 ± 0.33 respectively.

The pulse during the 5th, 10th, 15th and 20th minutes after the administration of the anaesthetic was 98.00 ± 3.86 , 104.00 ± 2.00 , 93.61 ± 5.15 and 91.00 ± 3.0 per minute respectively.

The respiration during the 5th, 10th, 15th and 20th minutes after the administration of the amaesthetic was 25.67 ± 1.75 , 23.60 ± 2.46 , 23.33 + 2.30 and 20.33 ± 1.66 per minute respectively.

The hasmogram recorded before the administration of the amesthetic was as follows:

REC - 10.08 ± 1.05 million/c.mm.,

VHC - 8333.33 ± 353.33/e.mm.,

Hb - 10.13 + 0.18 g 5 and

PGV - 31.67 ± 1.12 %.

The hasmogram recorded fifteen minutes after the administration of chloral hydrate was as follows:

RBC - 9.76 + 1.54 million/c.mm.,

WEC - 8383.33 ± 174.01/e.m..

Hb - 9.72 + 0.29 g 5 and

PCV - 31.00 ± 1.00 %.

The enturis were normal in feeding after recovery.

Group II B

The date are presented in Tables 2, 5 and 8.

The average bodyweight of the animals in this group was 30.17 ± 2.39 kg. Triflupromesine hydromhioride solution at the rate of 0.2 mg per kg bodyweight followed by chloral hydrate at the rate of 1.5 ml per kg bodyweight as a six per cent solution was administered intraveneusly. The time

taken for the injection was 70.67 + 9.10 seconds.

The palpebral reflex disappeared by \$1.00 ± 0.52 minutes and reappeared by \$9.00 ± 14.10 minutes. Inscordination of movements was observed by 0.50 ± 0.32 minutes. All the animals, except two, (Nos. § & 6) became recumbent by 0.63 ± 0.38 minutes. Out of the four animals which became recumbent, three shaimals (Nos. 1, 2 and 3), assumed lateral recumbency following sternal recumbency and one animal (No.4), assumed lateral recumbency whome.

Salivation was observed in three numbers (Nos.1, 2 and 3), Inscordination of movements was not observed in one animal (No.5). The period of recumbency was persisted for 36.63 ± 8.75. The animals became apparently normal in gait by 50.50 ± 13.82.

The temperature, pulse and respiration recorded before the administration of the assessments were 102.32 \(\tilde{2} \) 0.22°F, 88.00 \(\tilde{2} \) 3.06 per minute and 18.33 \(\tilde{2} \) 2.29 per minute respectively.

The temperature during the 3th, 10th, 15th and 20th minute after the administration of the



anaesthetic was 101.90 ± 0.28°r, 101.47 ± 0.34°F, 101.13 ± 0.40°F, 101.10 ± 0.45°F respectively.

The pulse rate during the 5th, 10th, 15th and 20th minutes after the administration of the ensesthetic were 99.00 ± 6.28, 95.00 ± 7.44, 92.67 ± 5.97 and 93.33 ± 3.78 per minute respectively.

The respiration during the 5th, 10th, 15th and 20th minutes after the administration of assessmentic was 22.00 ± 2.31 , 21.17 ± 1.33 , 20.83 ± 1.60 and 22.67 ± 2.35 per minute respectively.

The hasmogram resorded before administration of the amosthetic was as follows:

RBC = $10.00 \pm 1.35 \text{ million/e.mm.}$

48C - 8050 ± 275.389/c.mm.,

Hb - 10.36 ± 0.46 g % and

PCV - 32.00 ± 0.86 %.

The haemogram recorded fifteen minutes after the administration of chloral hydrate was as follows:

RBC - 9.15 ± 1.17 million/c.mm.,

чес - 8400 ± 163.30/с.ша.,

Hb - 9.88 + 0.50 g % and

PCV - 29.50 ± 0.34 %.

All the animals started feeding immediately after recovery.

GROUP II C

The date are presented in Tables 3. 6 and 9.

The average body-weight of the animals in this group was 31.17 ± 2.48 kg. Triflupromasine hydro-onlorids solution at the rate of 0.2 kg per kg body-weight spllowed by shloral hydrate at the rate of 2 ml per kg bodyweight as a six per cast solution was admissistered intravenously. The time taken for the intravenous administration was 96.67 ± 12.02 seconds.

The pelpebral reflex disappeared by 0.67 ± 0.49 minutes. Incoordination of movements observed immediately following the injection. All the animals, except No.3, assumed recumbency immediately after the injection. While, subset No.3 assumed recumbent position only after one minute following the injection. Four animals (Nos.2, 3, 4 and 6), assumed lateral recumbency following sternal recumbency and the remaining two animals (Nos. 1 and 5) assumed lateral recumbency alone. Salivation and lacrimation were observed in two animals (Nos. 1 and 2). Lacrimetics alone was noticed in three animals (Nos. 4, 5 and 6).

The temperature, pulse and respiration recorded before the administration of the anaesthetic were $102.23 \pm 0.23^{\circ}$ F, 81.83 ± 4.43 per minute and 21.30 ± 1.50 per minute respectively.

The temperature during the 5th, 10th, 15th and 20th minute after the administration of the amaesthetic was 101.93 ± 0.36, 101.89 ± 0.19, 101.73 ± 0.22 and 101.37 ± 0.25 per minute respectively.

The pulse during the 5th, 10th, 15th and 20th minutes after the administration of the anaesthetic was 98.67 ± 3.79 , 91.33 ± 4.78 , 69.32 ± 3.82 and 91.00 ± 5.13 per minute respectively.

The respiration during the 5th, 10th, 15th and 20th minute after the administration of annesthetic was 21.17 \pm 2.35, 20.81 \pm 2.37, 20.17 \pm 1.47 and 20,67 \pm 1.67 per minute respectively.

The hemogram recorded before administration of the amenthetic was as follows:

RBC - 9.83 ± 1.22 million/e.mm.,

VBC - 8100.00 ± 89.44/e.mm.;

1fb = 10.43 ± 0.46 g % and

PCV - 31.50 ± 0.62 %.

The hasmogram recorded fifteen minutes after the administration of chloral hydrate was as follows:

RBC - 9.73 ± 1.22 million/c.mm.,

WBG - 8150.00 ± 131.02/e.mm.,

Hb - 9.88 ± 0.45 g % and

PCV - 29.83 ± 0.54 %.

All the enimals started feeding immediately after mesovery.

TABLES

Table 1. Ansesthetic effect after administration of (1) chloral hydrate 6% solution alone at the rate of 1 ml/kg bodyweight and (11) Siquil at the rate of 0.2 mg/kg bodyweight followed by chloral hydrate 6% solution, at the rate of 1 ml/kg bodyweight.

Animal	Age	Veight	Quan-	Quen-	Time		B TAKINI P			TON OF	TIME T	AKEN FOR
No.	(months)	of the animal (kg)	tity of siquil (mg)	tity of choloral hydrate (ml)	taken for	Disappear- ance of m palpebral reflex (mts.)	Incoor- disation (mts.)	Recum- bency (mts.)	Absence of palpebral reflex (mts.)	Recumbency (ste.)	Mormal gait to resppear (mts.)	**
I-A-1	22v	24.0	•	24.0	35.0	2.5	2.0	N41	12,5		15+0	**
2	22	27.0	•	27.0	42.0	4.0	NAI	nil	10.0	**	•	**
3	29	38.0	*	36.0	50.0	3.0	1,0	NAI	8,0	. A second	11.0	••
4	27	36.5	•	36.5	48.0	Nil	1.8	11.0	Nil	4.0	14.0	
5	19	24.0	-	24.0	45.0	7.0	2.0	NAL	2.0	···	8.0	••
6	30	36.0	•	36.0	85.0	nii ·	MTT	NAL	NAL	€		••
	24,83	30.98		30.92	50.83	4.12	1.50	11.0	8.13-	4,0	12.0	• •
	2 1,82	± 2.70		± 2.70	± 7.16	± 1.01	± 0.29	\$ 0	£ 2,24	2	1.58	
-A-1	29	39.5	7.9	39.5	61.0	4.0	2.0	•	13.0	*	20.0	
2	19	25.0	5.0	25.0	25.0	2.0	3.0	•	18.0	*	19.0	•
3	22	23+0	4.6	23,0	25.0	2.0	10.0	10.0	5.0	1.0	3.0	
4	25	24.5	4.9	24.5	45.0	5.0	9.0	•	11.0	٠	8.0	•
. 5	22	20.0	6.0	30.0	90.0	2.0	1.5	. .	9.0		17.0	
6	17	22.0		22.0	90.0	2.5	1.5		9.0		18,0	00
68	22.33	27.33	5.47	27.33	56.00	2.92	4,50	10.0	10.83	1.0	14,17	\$ 3.75
,	± 1.75	*	± 0.54	2.68	± 12,08	\$ 0•52	± 1,60	.± 0.0	± 1.80	± 0.0	2.65	

Table 2. Assesthetic effect after administration of (1) chloral hydrate 6% solution alone at the rate of 1.5 ml/kg bodyveight and (11) Siguil at the rate of 0.2 mg/kg bodyveight followed by chloral hydrate 6% solution, at the rate of 1.5 ml/kg bodyweight.

A A	A	Weight	Quan-	Quan-	Time		HE TAKEN	POR	DUBA	tion of	TIME TAKEN FOR
Actual Ro.	Age (months)		of siquil (ag)	of chlorel hydrate (ml)	taken for injection (sec.)	palpebrak reflex (mts.)	Incoopedination (ats.)	Recuse beney (mts.)	Absence of palpebrel reflex (mts.)	(atr.)	Rormal gait to reappear (ats,)
I-B-1	22	24.5	**	36.75	35	2.0	*	*	13.0	•	•
2	23	25.5	•	38,25	45	2,0	1.0	2.0	12.0	930	18,0
3	19	24,0		36.0	40	1.0	1.0	1.0	18.0	22.0	22,0
4	22	27.5	••• ·	41.25	45	0.5	0.5	0,5	18.5	20 08.	20.5
5	27	37.0	•	55.5	70	1.0	1.0	1.0	19.0	14,0	23.0
6	29	39.0		58.5	75	•	*	*	•		
	24.00	29.58	*	44.03	55.00	1.30	0,68	1.13	14,90	19.68	20.48
	.# 1.51	4 2.72	•	± 4,09	4 5.92	& 0,30	0.13	± 0.22	1.38	3 4,52	1.09
I-D-1	25	25.0	5.0	37.50	55	Innedist.	Immediate	Zomediste	104.0	60.0	\$60 . 0
48	30	35.0	7.0	58.50	55	Innedia to	Immediate	1.0	65.0	19.0	60.0
3	22	27.0	5.4	40.50	100	Zamedio 60	Investinte	Immediate	\$0.0	, gg ,0	31.0
Δ	32	32.5	6.5	48.75	90	2.0	1.5	103	28.0	, g8.5	35.5
5	38	23.5	4.70	35.25	60	3.0	*	· •	7.0	.	e
6	89	39.0	7.60	57.00	100	1.0	1.0		60.0		C 1833.0
	25,17	30.17	6,07	45.25	76,67	1.00	0.50	0.63	49.00	36.63	ab- # 30 - 80
And a financial section of the secti	2	\$	*		4	\$		\$. 3	etic con anticipi
	1.45		0.48	3.38	9.10	0.58	0.58	0.38	14,10	0.19	. 15.08

Table 3. Anaesthetic effect after administration of (i) chlorel hydrate 6% solution alone at the rate of 2.0 ml/kg bedyweight and (ii) Siquil at the rate of 0.2 mg/kg bedyweight followed by chloral hydrate 6% solution, at the rate of 2.0 ml/kg bedyweight.

			Quen-	Quan-	Time	TIM	e taken po	R	DURATOR	I OP	TIME TAKEN FOR
Anima)	(months)	· (res)	eity of eiquil (mg)	of chloral hydrate (ml)	taken for injection (sec.)	Disappear- ance of palpebral roflex (mts.)	Incoor- dination (mts.)	Recumbency (mts.)	Absence of palpebral reflex (mts.)	(mts.)	Normal gait to reappear (mts.)
Z-C-1	19	25.0		50	75	2.0	1.0	2.0	3.0	2. 0	5.0
2	27	36.0	- L	72	80	5.0	9.0	•	10.0	•	97.0
3	22	26.5	· ••	53	120	5.0	Immediate	1.0	2.0	7.0	50.0
4	22	23.5		47	62	1.0	Immediate	Damedia te	24.0	38.0	32.0
	30	34.5	•	69	68	5.0	2.0	5.0	15.0	10.0	53.0
6	25	25.5	, *	51	65	5.0	2.0	A.	4.0		50.0
Menn	24.16	30.17	,	57.00	78,33	3.83	1.93	2,00	9.67	12.75	47.83
	2	*	r	2	*	*	*	*	. '*	\$	
5 2	1.62	2,28	° 🖛	4.36	8.75	0.75	0.49	1,08	. 9.31	6.63	12,41
CleCe)	27	36.0	7.2	72	120	James din t	• Immediate	. Immediate	85.0	70.0	87.0
à	29	38.0	7.6	76	120	Immediat	· Immediate	• Immediate	42.0	40.0	57.0
, <u>3</u>	19	25,0	5.0	50	120	1.0	Immediat	e 5 ₄ 0	53.20	20.0	99.0
4	25	25.0	5.0	50	60	Imediat	o Inmediat	· Immediate	53.00	30.0	66.0
5	30	35.5	7.1	71	100	Implica	e Immediat	· Immediate	70.00	35.0	70.0
6	22	27.5	5.5	55	60		•	· Immediate	95+0	78,0	93,0
Keén		Activities and the Activities of the Control of the		62.33	96.67	0.67	0.0	0.69	66,33	47.00	78.67
	25.33 2 1.73	4	6.23 ± 0.49	12,02	± 12.02	± 0.49	*	± 0.83	8.43	9.88	6,82

Table 4. Temperature, pulse and respiration before and efter administration of

(i) chieval hydrate of solution at the rate of 1 mi/kg bodyweight clone and

(ii) Siguil 0.2 mg/kg bodyweight followed by chieval hydrate of solution
at the rate of 1 mi/kg bodyweight.

	Before	,	TEMPTH	ATURE	•	Before			Pulse edukaleti	ntion	Before After admin				nelon_
Animal No.	edminist retion of encesther	510	After admit	(°r)	(°F)	administ- ration of anaesthetic		1010	156h	2011	Son.	54h	10th		201h
Zoho!	103.0	102.2	102.0	102,0	102.0	90	100	100	90	90	20	22	24	20	20
2	102.2	101.2	102.0	102.0	102.0	82	68	84	84	60	20	20	22	23	20
3	109.8	109.8	103.4	103.4	103.4	72	72	80	80	70	16	16	20	16	16
4	103.8	103.6	103.6	102.6	103.2	80	108	96	88		18	28	80	24	18
5	103,2	103.2	102.6	103.0	103.0	88	100	100	90		16	80	20	80	18
6		102.0	101.8	102.0	101.8	88	65	68	96	96	80	50	28	. 20	18
Mann.	\$02,07	102,83	102,57	102,67	102,60	89.33	92,67	91.33	88,00	86.00	18,33	81.33	21.33	20,67	10.33
6E	& Q.30	± 0.81	2	2 0.35	± 0.31	2.77	± 8.77	± 3,49	± 8,25	3.45	0.80	0,80	1,43		0,61
II-b-1	102.8	102.8	102.2	102.0	101.0	88	100	104	80	603	20	10	N	18	16
á	101.8	101.8	101.4	101.0	101.8	74	100	100	82	98 (1)	24	20	10	18	
3	102-0	102.6	103.0	102,2	102,2	88	108	110	66	85	80	28		20	18
*	103.3	103.6	103,6	103.0	103.0	88	80	100	160	100	17	24	. 20	24	80
5	101.2	101.4	101.8	101.0	101.0	100	102	110	110	100	30	50	3 0	30	80
6	101.4	101.6	4.101	101.0	101.2	70	63	100	104	90	33	36	30	JP)	
	102.07	108-33		101,83	101.70	_	98,00	104,00	_	91.00 L	\$2.50 £	±		23.33 &	e erod L
	2 0.38	± 0.33	2 0.40	£ 0.31	± 0.3	, 4,46	2.06	2,00	5.19 °	3.04	1,86	1.75	e,W	2,37	1,66

Table 5. Temperature, pulse and respiration before and after administration of

(1) chloral hydrate 6% solution at the rate of 1.5 ml/kg bodyweight alone and

(11) Signil 0.2 mg/kg bodyweight followed by chloral hydrate 6% solution
at the rate of 1.5 ml/kg bodyweight.

animi	Before			Perature Ploiotro	iton	Defore edminist-			Pulse Moiet	ition	Batars Schoolst-	Afte	nesy:	RATTON destab	
No.	ention of encesthetic (°2)	5th (**)	(°F)	15th (\$r)	204h (°F)	ration of Gundathetio	546	foth	15tb	2013	e tion		TOEB	1306	20th
Z-D-1	161.8	102,0	101.8	102,2	202.2	72	96	90	100	100	20	18	20	18	16
3	102,2	101.8	101.8	101.6	102.2	80	100	104	104	110	16	20	18	18	34
3	102.2	103.2	108.8	102.0	102.0	GG.	108	104	110	106	16	25	24	25	24
•	103.4	103.4	103.4	102.2	102.2	80	92	88	92	92	16	24	20	20	18
3	102.2	102.4	102.2	102,8	102.6	60	96	100	98	98	16	23	18	20	20
6	193.0	103.0	102.8	103.0	103.0	72	96	92	88	88	16	18	20	16	16
Hand	102,47	108.77	102.47	102,30	102.37	78.67	90.00	96.33	98,67	95.00	16,67	21.50	20,00	20.67	19.61
6 1	4 0,25	± 0.33	± 0.87	4 0.21	£ 0.15	2 . 2 . 46	2 2,25	2 3.93),25	3.34	0.67	1.34	0,90	1,58	1.4
Z-B-1	102.0	102,2	102.9	101.8	101.6	26	(IO	72	80	85	98	16	16	24	80
.	101.6	101.6	100.4	100.2	100.5	78	100	100	28	100	46	20	8 \$	24	30
. 3	102.0	102.0	101.2	101.0	100.8	92	112	96	100	96	19	20 :	80	18	30
4	102,0	101.0	4.101	101.2	100.0	100	100	106	96	90	, 50	30	25	83	30
\$	102.8	103.0	102.8	102.6	103.0	30	.112	120	116	106	16	20	23	16	18
6	102,5	101.6	101.0	100.0	101.0	84	66	76	76	80		18			10
Man	102.32	101.90	101.47	101.13	101.10		99.00	95.00 £	92.67 £	93+33 ±	18,89	22.00 ±	21.17	. 4	4
8B	0.22 4		2 0.34	2 0,40	£ 0,45	3,06	\$ 6,85	7.4	5.97	2,78	3,1)	8.31	1.53	1.60	2.9

Table 6. Temperature pulse and respiration before and after administration of

(i) chloral hydrate 6% solution at the rate of 2.0 ml/kg bodyweight sione and

(ii) Siquil 0.2 mg/kg bodyweight followed by chloral hydrate 6% solution
at the rate of 2.0 ml/kg bodyweight.

Animal	Before administ-		TD) fter admi	PERATURE		Before			lse		Before administa		After administration		
No.	ration of amesthetic (°P)	5th (°p)	10th (°F)	15th (°r)	20th (°y)	ration of apacsthetic	5th	10th	nietratio 15th	20 th	, 136a	56h	10th	13th	20 th
1+G-1	102.0	102,4	102.2	103-2	103.0	88	120	100	100	98	18	25	20	24	55
2	101-8	101.8	102,0	101.8	101.8	86	120	86	86	80	. 18	24	24	20	20
. 3	101.8	102.0	101.8	101,4	102,0	80	100	100	84	100	80	18	20	16	16
4	101.8	102.0	101,8.	101,8	101.6	84	104	96	96	94	50	30	20	18	16
5	102.8	102.4	102.4	101.8	102+0	65	80	84	100	86	(9)	20	18	16	16
6	101.8	102.0	101.0	101.8	102,4	80	100	102	96	82	10	18	22	18	20
Hean SE	102.00 ± 0.16	102,10 ± 0,10	101.67 ± 0.20	101.97 ± 0.25	102.1: ± 0.28	4	104,00 ± 6,12	94.67 ± 3.17	93.67 ± 2.85	90.00 ± 3.46	*** 20.33 \$ 1.97	22.50 ± 1.93	20.68 3 0.84	19.67 2 0.68	18.67 ± 0.99
EZ-G-1	103.0	103.0	102,6	102.6	101.6	86	96	86	96	86	23.	14	16	20	20 /4 m
2	102.2	102.0	102.0	101.4	101.4	76	100	90	80	70	, 16	17	45	16	17
3	101.8	101,8	102.0	101.8	101.4	78	100	100	92	108	22	28	28	22	22
4	102.0	102.0	101.6	101,6	102.0	88	82	84	76	96	(1)	20	20	17	17
5	101.8	101.2	101.3	101.0	100.2	64	106	110	100	92		25	 	20	80
6	102.8	101.6	101.8	102.0	101,6	94	108	78	92	94	85	50	186	26	28
yesn 93	102,23 ± 0,23	101.93 ± 0,36	101.87 ± 0.19	101.73 ± 0.22	101.3 ± 0.2		98.67 # 3.79	91.33 ± 4.78	69.33 ± 3.82	91.00 ± • 5.13	21/59 1/50	21,17 2 2,35		20.17	4

Table 7. Hasmatological values before administration and fifteen minutes after administration of (1) chloral hydrate 6% solution at the rate of 1 ml/kg bodyweight and (11) Siquil 0.2 mg/kg bodyweight followed by chloral hydrate 6% solution at the rate of 1 ml/kg bodyweight.

	BEF	ORE ADMINIST	RATION	r .	15 MINUTES AFTER AIMINISTRATION						
Animal No *	RBC m1111on/e-ms	VEC/	Bb 6 \$	PGV	RBC million/c.mm	VBC/ C+III	# \$ 	P			
Z-A-1	12,00	9000	10.5	32	11.8	9000	10+5	30			
2	11.00	7300	11.0	28	11.0	8000	11.0	30			
3	6,80	6950	9.4	39	5.9	8100	8.2	93			
4	6.98	6650	9,6	40	5.8	8000	8.0	31			
	11.00	8000	10.0	30	10.0	9800	10.0	50			
6	6.10	7800	8.5	31	5.12	7950	8,5,	29			
Head SD	8.98 1.07	7650.00 340.59	9.83 0.35	33.33 2.03	8,27 1,22	8475,00 310,85	9.31 8.53	30,50 0.56			
2 3 4 5 6	7.50 11.80 12.00 6.20 12.00	10000,00 8000,00 8000,00 8000 8000	10.0 10.0 10.0 9.8 10.0	35 30 30 34 33 28	7.25 11.5 11.8 5.2 11.8 11.0	8500 9000 8000 8500 8500 7800	9.0' 10.0' 9.9' 9.0' 9.8'	90 90 90 36 30 90			
ions Is	10.08 ±	8333.33 ± 333.33	10.13 2 0.18	21.67 ± 1.12	9.76 ± 1.54	8383.33 ± 174.01	9.77	51.00 ± 1.00			

Table 8. Rematological values before administration and fifteen minutes after administration of (1) chloral hydrate 6% solution at the rate of 1.5 ml/kg bodyweight and (11) Siguil 0.2 mg/kg bodyweight followed by chloral hydrate 6% solution at the rate of 1.5 ml/kg body-weight.

الله المناسبة		erore amini			15 MINUTES AFTER ADMINISTRATION						
Animal No.	REC million/o,mm	WBC/ C+BB	#6 6 \$	*********	REC Militer/e.m.	VDC S.III	Bb A.L.	PEV			
X-B-1	4,8	9300	8.5	24	5+7	10000	8.0	35			
2	5.2	570 0	9,0	28	4.9	4800	7.4	24			
3	6.2	7900	10.0	32	6,0	7700	9.0	31			
4	6.0	7650	10.0	<i>3</i> 0	5.7	8200	9.0	29			
5	740	9200	9.0	34	. 3,9	9600	7.0	32			
6	6.8	11000	10.0	23	7.7	8500	9,0	50			
en E	6.80 # 0.35	6808.33 ± 1269.19	9.42 ± 0.27	30.5 6 ± 1.67	6,03 ± 0,39	8133.33 ± 754.39	8,20 ± 0,38	30.17 3 1.49			
i-B-1	6.0	8000	9.8	94	5.9	8000	9.0	30			
2	6.0	7800	8.5	21	5.2	7900	8.0				
3	11.0	8000	11.0	30 °	11.0	8500	10.8	50			
4	12.0	8000	10.0	35	11.8	8500	9,6	50 ()			
5	11.0	9000	11.0	30	11.0	8500	10.9	28 24 24			
6	14.0	7000	12.0	35	12.0	9000	11,0				
	10,00 2 1,35	8050 ± 275+38	10.38 ± 0.46	0.86 2	9.15 ± 1.17	8400 9 163.30	.68. <u>2</u> 2 0.50	29.50 2 0.34			

Table 9. Haematological values before administration and fifteen minutes after administration of (i) chloral hydrate 65 solution at the rate of 2.0 ml/kg bodyweight and (ii) Siquil 0.2 mg/kg bodyweight followed by chloral hydrate 65 solution at the rate of 2.9 ml/kg bodyweight.

	<u> </u>	EFORE ABUIN	ESTRATION			TISTRATION		
Animal No.	RBG million/c.um	VBC/ Opinio	HD 6 \$	PCV	RBC million/o.mm	VBC C	B) 6 \$	P C V
Z-6-1	6,6	6500	10	30	542	6000	8.0	25
2	7.0	8500	10	35	6.0	9500	8.5	32
3	6.4	8500	10	30	6.2	9500	9.0	32
4	5.8	9000	8,8	30	5.4	10000	8.0	28
5	6.5	9000	8.5	28	6.0	9500	8.0	26
6	6.2	8006	9.8	34	5.2	8000	9•0h	26
Keen	6.42	8250,00	9.52	29.50	5.67	8416,67	8,48	28,17
S B	± 0.16	2 381.86	± 0.28	± 1.45	± 0,18	± 624,72	2 0,20	土 1,28
II-C-1	12.0	8000	10.0	32.	11.8	7800	9.0	30
2	12.0	8000	12.0	32	12.0	8000	11.0	30
3	12.0	8000	11.0	30	11.8	8500	10.5	50 %/
4	6.0	8200	9.8	34	6.0	8600	9.0	32
.5	6.0	7900	8.8	31	5.8	8000	8,2	29 534
6	11.0	8000	11.0	30	11.0	8000	10.8	28
lean	9.833	8100	10.43	31.50	9.73	81,50,00	9.88	89. 03
	1.22	± 89.44	± 0.46	± 0,62	± 1,22	131.02	0,43	



DIBCUSSION

In the present study, freshly prepared chlorel hydrate six per cent solution was administered intraveneusly in Group I, while triflupromasine hydrochloride at the rate of 0.2 mg per kg bedyweight was also administered intravenously as the presediennt in Group II.

Palpebral Reflex

When chloral hydrate alone was administered, pulpobrul reflex disappeared by 4.13 \pm 1.01, 1.30 \pm 0.30 and 3.83 \pm 0.75 minutes and reappeared after 8.13 \pm 2.24, 14.90 \pm 1.38 and 9.67 \pm 3.51 minutes respectively in I-A, I-B and I-C. When triflupromatine was also administered, it disappeared by 2.92 \pm 0.52, 1.00 \pm 0.52 and 0.67 \pm 0.49 minutes and reappeared after 10.83 \pm 1.99, 49.00 \pm 14.10 and 66.33 \pm 8.43 minutes respectively in IX-A, IX-B and II-C.

When the dose of chloral hydrate was increased, disap, earnes of pelpebral reflex was quicker and the duration longer in both the groups.

It could be sen that disappearance of palpebral reflex took place quicker and the duration prolonged

when triflupromasine hydrochloride was used as the presedicant. In I-B, palpebral reflex disappeared quickest and persisted maximum.

Incoordination of Movements

when chlored hydrate alone was administered, incoordination of movements was observed by 1.50 ± 0.29, 0.88 ± 0.13 and 1.33 ± 0.59 minutes and these animals assumed normal gait by 12.00 ± 1.58, 20.88 ± 1.09 and 48.83 ± 12.41 minutes respectively in I-A, I-B and I-C. When triflupromasine hydrochloride was also administered, incoordination of movements was observed by 4.50 ± 1.60 and 0.50 ± 0.32 in II-A and II-B and was immutate in II-C, and these animals assumed normal mait after 14.17 ± 2.85, 50.50 ± 13.82 and 78.67 ± 0.82 minutes respectively in II-A, II-B and II-C.

when the dose of chloral hydrate was increased, setting in of incoordination of movements was quicker and the time for assuming the normal gait was longer. The administration of triflupromasine hydrochloride shortened the time for setting in of incoordination of movements except in II-A and prolonged the time for assuming normal gait.

Recumbency

In I-A, only one animal (No.4) assumed recumbest position at the 11th minute which persisted waly for four minutes. In I-B, all except two animals (Nos. 1 and 6), assumed recumbency in 1.13 ± 4.32 minutes which persisted for 14.88 ± 4.32 minutes. In I-C, two animals (Nee, 2 and 6) did not assume recumbency, while the rest four became recumbest within 2.00 + 1.08 minutes. Out of these four enimals, ene (No.4) became recombent immedia wely fellowing the injection. The duration of recumbercy in these enimals was 12.75 ± 6.63 migutes. It is interesting to note that the animal which become recumbent immediately after the injection had the longest duration of recumbency (32 minutes). In II-A only one snire! (No.3) assumed recumbency at the 10th minute which persisted only for one minute. In II-B, only four animals assumed recumbent position in 0.63 ± 0.38 minutes which persisted for 36.63 ± 8.75 minutes. Out of these, two animals (Nos.I and 3) assumed recumbency immediately after the injection. In II-C, all the animals, excepting

No.3 absumed recumbent position immediately after the injection. The animal No.3 assumed recumbency only after five minutes. In this subgroup, recumbency persisted for 47.00 ± 9.68 minutes.

The easet of recumbency was quicker and the duration more prolonged when triflupromasine bydrochloride was used as the premedicant.

revied of recumbency recorded in the present study is in agreement with that of Bols (1968) whereis, administration of 0.08 to 0.1 g per kg bodyweight produced narcosis lasting for about half as hour.

Temperature, Pulse and Respiration

recorded before and every five minutes after the administration of the ansesthatic up to 20 minutes.

The variation in temperature was within 1°F and was not of any olinical significance.

There was an increase in the rate of respiration and pulse within five minutes of

administration of the drug and it showed a gradual downward trend. This is in agreement with the observations of Chandra et al. (1970) in calves.

lin emogram

The blood values were estimated before and 15 minuses after the administration of the anaesthetie.

There was reduction in the erythrocyte count, has moglobin and packed well volume, while there was slight increase in the leucocyte count. Observations recorded in the present study are in agreement with that of Singh gi gl. (1975) in bovines using chloromag and of Major gi gl. (1978) in buffalo calves seing chloral hydrate.



SUMMARY

The present study was conducted on 36 apparently healthy, Alpine - Malabari cross-bred bucks, aged from 17 to 30 months and weighing from 22 to 39.5 kg. They were divided into two groups vis., Group I and II, each consisting of 18 emissis. Each of these groups was further divided into three subgroups vis.

A, B and C and the anisals were numbered serially from 1 to 6.

pressly prepared chloral hydrate six per cent solution was administered intravenously at the rate of 1.0, 1.5 and 2.0 ml per kg body-weight in Group I, while triflupromasine hydro-chloride at the rate of 0.2 mg per kg bodyweight was size administered intravenously as the pressdicent, in Group II.

When the dose of chloral hydrate was increased, disappearance of palpebral reflex was quicker and the duration longer in both the

groups. When triflupromasine hydrochloride was used as the premedicant, disappearance of palpebral reflex was more rapid and the duration further prolonged.

increased, setting in of incoordination of movements was quicker and the time for assuming the normal gait was longer. When triflupromasine bydrochloride was administered as the premediennt the time for setting in of incoordination of movements was shortened and it prolonged the time for assuming normal gait.

recumbency was not seen in Group I-A, excepting in one animal, whereas four animals each of the subgroups B and C, assumed recumbency. The onset of recumbency was quicker and the duration more prolonged when triflupromasine hydrochloride was used as the premedicant.

The temperature, pulse and respiration were

recorded before and every five minutes after the administration of the anaesthetic up to 20 minutes. The variation in temperature was within 1° r and was not of any significance. There was an increase in the rate of respiration and pulse within five minutes of administration of the drug and it showed a gradual downward trend.

The blood values were estimated before and 15 minutes after the administration of the anaesthetic. There was reduction in the crythrocyte count, as a moglobin and packed cell volume, while there was slight increase in the leucocyte count.

In goats, chloral hydrate six per cent solution at the sate of 2 ml per kg bodyweight may be administered intravenously for surgioni procedures of about 30 minutes duration. Premedication with triflupromaning hydrochloride at the rate of 0.2 mg per kg bodyweight would give better results.

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CHLORAL HYDRATE FOR GENERAL ANAESTHESIA IN GOATS

БΥ

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ABSTRACT OF A THESIS

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ARSTRACT

Thirtysix apparently healthy Alpine-Halabari cross-bred bucks aged from 17 to 30 months and weighing from 22 to 39.5 kg were used for the present study. They were divided into two groups vis.,

Group I and II, consisting of 16 animals in each.

Bach of these groups was further divided into three subgroups vis., A, B and C, consisting of six animals in each.

Freshly prepared chlorel hydrate six per cent solution was administered intravenously at the rate of 1.0, 1.5 and 2.0 ml per kg bedyweight, while triflugromagine hydrochloride (Siquil) at the rate of 0.2 mg per kg bodyweight was aims administered intravenously as the premedicant, in Group II.

Pellowing were the selient results obtained during the present study:

When the dose of chloral hydrate was increased,

- i disappears me of palpebral reflex was quicker and duration was lenger,
- ii) setting in of incoordination of movements was quicker and the time for assuming the normal gait was longer, and

iii) recumbency was seen only in higher doses.

Administration of triflupromasine hydrochloride

(Siquil) as promedicant, increased the efficiency
of analythetic effect in these animals.

Temperature variation consequent on the administration of chloral hydrate, with and without presedication, was not significant, the variation being less than 10 F.

There was an increase in the rate of respiration and pulse within five minutes of administration of the drug(s), which showed a gradual downward trand.

There was reduction in the erythrocyte count, has most obin and packed cell volume, while there was slight there as in the leucocyte count.

on the basis of the present study, intravenous administration of chloral hydrate, six per cent solution, at the rate of two ml per kg bodyweight, may be meconsended in gosts, for surgical procedures

of about half an hour duration. Premedication with triflupromasine hydrochloride at the rate of 0.2 mg per kg bodyweight would give better results: