

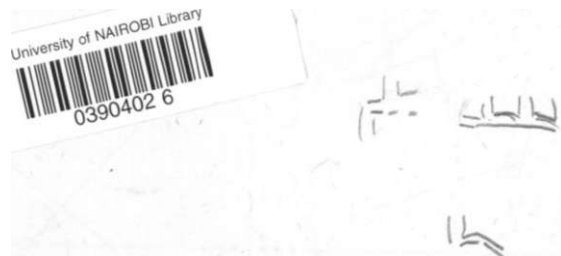
TITLE:

Intensive Care Management of Acute luryngotracheobronchitis
(LTB) patients admitted to Kenyatta National Hospital
Intensive Care Unit-1972 August to December 1984.

by

DR, MULATYA, JONAH LAZAROUS

A dissertation submitted in part fulfilment for the
Degree of Master of Medicine (Anaesthesia) of the
University of Nairobi.

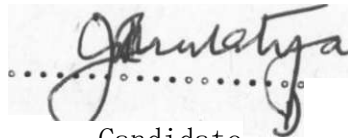


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DECLARATION:

This dissertation is *y origin work and has not been presented for any degree in any other University.

Signed;



Candidate /U C4 6

This dissertation has been submitted for examination with my approval

Signed:



Supervisor

V ^ W t t h h J)

A C K N O W L E D G E M E N T S

I wish to express my sincere thanks and gratitude to the following:-

Dr. SOK. Kahuho for his supervision and guidance in writing this dissertation.

Mr. Kimani, Nursing Officer I, IoC.Uo for making it possible for me to get the IoC.Uo Admission Registers.

All staff of Kenvatta National Hospital Records Department who made the patients files available to me.

The Director, Kenvatta National Hospital, for his permission to allow me to undertake this study.

My wife Lucy for her cooperation during the period of study.

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SUMMARY

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A retrospective study was carried out on 209 cases of laryngotracheal bronchitis (LTB) treated in the Intensive Care Unit (I.C.U) at the Kenyatta National Hospital (KNH) from August 1972 to December 1984. There were 134 males (64.1%) and 75 females (35.9%), and therefore, a Male:Female ratio of 1.8-1.

The average age of these patients was 3¹/₂ months, and the range was from 2 months to 13 years; about 80% of these patients were 2 years or less. About 65.6% of the patients were admitted to I.C.U. from the Paediatric Admission ward (P.O.Iv). Late admission of LTB patients to I.C.U. was noted in about 13.8% of the patients who were admitted in LTB Grade XV.

About 66% of the patients had post-measles LTB. Bronchopneumonia and congestive cardiac failure (CCF) were the main complications before and after admission on most of these patients. Clinical impression was the main criteria for the I.C.U. admission. Blood-gas analysis (BGA) on admission were done in only 24.4% of the patients. About 42.9% of these BGAs showed respiratory alkalosis. About 44% of these patients had 110 serial BGAs done.

The duration of stay in the Intensive Care Unit (I.C.U.) ranged from

one day to 183 days with an average of 11.2 days. Organisms commonly grown from tracheal aspirate cultures were Klebsiella, Pseudomonas and Staphylococcus aureus, These Organisms were resistant to commonly used antibiotics. Although most of the patients had ^I low packed cell volume (PCV) only 16 ^v patients had blood transfusions done.

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A. mortality rate of 15% was noted and this was high as compared to that found in other centres. Bronchopneumonia and CCF were the main causes of death. The high mortality rate appeared to correlate closely with late admission to I.C.U, complications on admission and other complications arising in I.C.U.

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Active immunization against measles has reduced the number of LTB patients admitted to I.C.U. progressively! over the last few years.

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INTRODUCTION AND REVIEW OF LITERATURE

Acute laryngotracheobronchitis (LTB), commonly known as croup, is an inflammatory disease of the upper respiratory tract, consisting mainly of subglottic oedema. It is a subacute viral illness (1-3) characterised by fever, barking cough and stridor. Etiologic agents in developed countries are mainly parainfluenza viruses 1 and 2 and adenoviruses (1, 31, 34, 39, 40, 51, 60, 62). In developing countries, measles is the commonest cause of LTB (50, 67).

The illness occurs mainly in cold months and usually lasts for 3-7 days. Males affected by the disease outnumber the females (5-8, 13, 25-26). Acute LTB affects children aged from 6 months to 14 years (15, 53). Its treatment has changed a lot in the last fifty years (15, 21, 32). Very mild cases are adequately treated as out-patients: moderate to severe cases are treated in General Paediatric wards, while very severe cases require Intensive Care Therapy (21, 30, 67). In the treatment of LTB oxygen is indicated because hypoxaemia can occur in patients with mild airway obstruction (37), and since the hypoxaemia associated with LTB corresponds with a raised respiratory rate (25, 27, 50). Humidification

of inspired gases used in the treatment of LTB helps to prevent drying up of secretions (22). Moist air

though commonly used, can cause water intoxication (41) and hyponatraemia and has not been shown to relieve symptoms, reduce obstruction or alter the clinical pattern of the illness (66). The use of saline and intermittent positive pressure breathing (IPPB) has also been shown to lead to clinical improvement in mild LTB patients (38, 65).

Racemic adrenaline has been used in the management of acute LTB. It has vasoactive action of adrenaline without rebound vasodilatation and cardiovascular side-effects of tachycardia and hypertension. (15)* The use of racemic adrenaline in treatment of acute LTB has been controversial over the past few decades and results on its use have led to varying conclusions: it can lead to avoided endotracheal intubation in only few patients (13, 50); IPPB and racemic adrenaline decreases the incidence of tracheostomies (10, 11, 15, 32); IPPB and nebulized racemic adrenaline is more effective than mist therapy alone (11); the use of racemic adrenaline gives short-lived beneficial effects (7, 11, 15), or it has no effect at all (38). Racemic adrenaline, however, has been shown to reduce airway oedema and obstruction in acute LTB as well as post-intubation croup (1, 4, 11, 15, 26, 32, 38, 42-43, 45, 61). The side-effects of racemic adrenaline are minimal, and it is useful in acute LTB since the disease usually-

runs its course before tolerance to its effects
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lias had time to develop. Initially treatments may
be repeated at frequent intervals. Lessening the
need for treatment with racemic adrenaline is a
clinical indicator for improvement in a patient's
condition (26).

An artificial airway is necessary in some
patients with severe LTB. Several factors point
to the need for airway support: patient fatigue,
decreasing response to racemic adrenaline, and toxicity
with evidence of bacterial superinfection. Tracheostomy
was for many years the procedure of choice as initial
studies suggested that nasotracheal intubation
might have serious airway injury(12,36,55). However,
recent studies show that carefully done nasotracheal
intubation provides effective treatment for LTB and
reduces morbidity when compared to tracheostomy (12,
28, 35, 40, 50, 67, 71, 72, 83-84).

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Endotracheal intubation has been used for a
long time and most of the problems associated with
it can be detected, and serious complications
avoided. The use of non-irritant materials and
methods of ensuring leak around the tubes have
reduced the incidence of subglottic stenosis (71,72).

Nasotracheal intubation is now preferred to tracheostomy in many units but in units where personnel are unfamiliar with nasotracheal intubation, tracheostomy is still a satisfactory alternative. However, where severe laryngeal obstruction is present and in older children likely to require artificial airway for a long period tracheostomy is the method of choice (67,72). The complications of intubation include pneumonia, atelectasis and subglottic stenosis (26,50). These complications are related to the duration of intubation (71), and any factor which increases tissue trauma and reduces tissue resistance to injury and infection following removal of the tube (71)* The incidence of intubation sequelae has not been consistently or clearly defined in terms of length of continuous intubation (58,71)* Longer periods of intubation may have very low rate of serious complications, associated with it (71). The complications arising from the conversion of nasotracheal intubation to a tracheostomy may be greater than those expected from increasing the length of endotracheal intubation (71). In developing countries, the duration of endotracheal intubation is longer and percentage of severe LTB patients is higher than in developed countries (13,50,67).

The greatest controversy in the treatment of acute LTB revolves around the use of Steroids (8, 16,

18, -21, 29, 50, 57-59, 65). Their known anti-inflammatory effects, particularly their actions on stabilizing cell membranes and to diminish capillary dilatation and permeability (33) have lead to wide applications and research, although methodologic problems have raised questions about validity of the results (1,6, 16). Many authors feel that steroids do not alter the clinical course of acute LTB. (1,7,8,58-59, 63); others maintain that steroids given in adequate doses provide an effective treatment (6, 19-21, 57, 64), while others feel that steroids may have beneficial effects in certain types of LTB (1,19). Random studies using low dose steroids¹ have shown that steroids have tracheostomy-sparing effects (6,20-21).

The use of steroids has also been shown to lessen the number of treatments with racemic adrenaline (9). It is now widely accepted that a single dose of steroids given early in the course of acute LTB provides most beneficial effects, and it is safe e.g. 1.0-1.5mg/kg of dexamethasone(9). Less than 0.3mg/kg of dexamethase can be considered an inadequate dose with which to treat acute LTB (16,17) while a dose of 0.3mg/kg, or greater, can be considered adequate (26).

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The use of hydrocortisone 100mg intramuscularly initially suggested by Davison (21) or its equivalent of dexamethasone or prednisone (21) is adequate in moderately severe LTB. The use of steroids appears contraindicated in immunologically depressed malnourished LTB patients (5*67). Although LTB is a viral infection, superinfection with streptococcus pyogenes, staphylococcus aureus, streptococcus pneumonia, Haemophilus influenza etc.,¹ does occur (1, 21). It has also been noted that LTB in under-privileged children rarely occurs alone. It is usually associated with lower respiratory tract (LRT) infection, pneumonia, anaemia, malnutrition and other complications of measles e.g. gastroenteritis and herpes stomatitis and cardiac failure (8,21,37, •46-47, 50, 67, **73**, 75-76, 78-81). Therefore, the routine use of antibiotics is widely accepted (1,10, 13,21, 30, 48, 50, 67) although some authors feel that antibiotics, though commonly used, are of doubtful value in uncomplicated LTB (52,54).

Acute LTB can progress rapidly to respiratory failure, and detection of the onset of respiratory-failure clinically is very subjective. Repeated blood gas analysis¹ is therefore very important.

BGA of these patients has shown that their arterial blood partial pressure of oxygen (PaO_2) is lower than the normal for the age group and that most of the patients have arterial blood partial pressure of carbon dioxide (PaCO_2) greater than normal values (25). There is a relationship between PaCO_2 and severity of airway obstruction (68) and an inverse relationship between PaCO_2 and respiratory rate (25).

LTB patients also frequently have gas exchange failure which does not correspond with the clinical picture. The presence of cyanosis in these patients indicates hypoxaemia, but the absence of cyanosis does not exclude moderate hypoxaemia (27). BGA in LTB patients has also shown that most mild to moderately obstructed patients tend to hyperventilate and therefore the PaCO_2 rise which accompanies decompensation may not be above the normal by the time intubation is needed (82). There is no relationship between arterial BGA results and the clinical picture and therefore BGA, cannot alone be used to assess severity of LTB. Furthermore, BGA has no value in assessing the need for intubation (1).

Therefore, the decision on when to intubate a LTB patient depends upon many factors; clinical grounds e.g. extreme respiratory distress, pneumofl

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Therefore, the decision on when to intubate a LTB patient depends upon many factors; clinical grounds e.g. extreme respiratory distress, pneumonia

cardiac failure; significant arterial hypoxaemia and hypercarbia (50) and scoring system (4[^]).

However, no absolute indications to intubation or extubation exist (67,50).

Mortality rate of severe LTB is higher in developing countries (7,39,50,67,7?) than in developed countries (10,13, 20, 53, 7[^]-75) where the duration of I.C.,U. stay of severe LTB patients is even shorter (10,13) than in developing countries (50). Long-term follow-up of LTB patients show that they have further airway involvement manifesting as recurrent LTB (?4) and clinical bronchial asthma (5,19,20, 25-26).

MMS AND OBJECTIVES

This study was undertaken to:

Evaluate the management of severe LTB patients
admitted to Kenvatta National Hospitalⁱ IoColJ.^{*}
between August 1972' and December 1984

To determine the complications of severe LTB and
their management.

Make suggestions on how to improve the management
of the patients.

M A T E R I A L ? A N D M E T H O D S

4. retrospective study of 209 acute LTB patients admitted to the I.C.U. (KNH) between August 1972 and December 1984 was done. From the I.C.U. admission registers, the names and other particulars were obtained and their case notes studied closely. Information relating to I.C.U. management, complications mortality etc was carefully studied and analysed. A data collection form was completed for each patient.

i

Most of the I.C.U. management of these patients is outlined in this study.- The management also included the following:

v

- 1). Vital signs monitoring: Pulse-rate by palpation of peripheral arterial pulses, one hourly; respiration rate by observing and counting of chest movements. One hourly; 4 hourly Temperature charting using mercury Thermometer 4 hourly blood pressure charting' using sphygmomanometer and arm cuff method.
- (2) Nursing Care: Bathing, Turning 4 hourly, suction (at least 4 hourly) Nasogastric feeding 4 hourly, Care of pressure areas weighing once daily.
Continuous Electrocardiography (ECG), central venous pressure (CVP) monitor, input-output charting, physiotherapy 4 hourly with endotracheal tube, tracheostomy

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- (2) Nursing Care: Bathing, Turning 4 hourly, suction (at least 4 hourly) Nasogastric tube feeding 4 hourly, Care of pressure areas and weighing once daily.
- (3) Continuous Electrocardiography (ECG), rarely central venous pressure (CVP) monitoring, input-output charting, physiotherapy at least 4 hourly with endotracheal tube, tracheobronchial

or/and oral suction.

laboratory investigations, including full
hemograms, culture and sensitivity of tracheal
aspirates twice weekly.

Radiological investigations, mainly chest X-Ray
films.

RESULTS :

The results of this study were as follows;

Out of the 209 patients studied 134 (64.1%) were males and 75 (35.9%) were females; giving a male: female ratio of 1.8:1.0 (Table I).

The age of the patients ranged from 2 months to 156 months, and the mean was 30.9 ± 4.9 months. (Figure 1 and Table 2). About 10.0% of the patients were 24 months (2 years) or less.

About 65.0% of the patients had post-measles LTB. There were no viral studies done on the 209 patients studied. The aetiological agents of the LTB were therefore, based on the clinical manifestations of the disease. Diagnosis of viral LTB (which was made when there was no clinical evidence of measles and it was therefore assumed that the LTB was caused by other viruses other than measles virus") see respiratory syncytial virus, parainfluenza etc. (Table 3).

The grading of LTB on admission to ICII was done on the following clinical grounds:

Grade I: LTB with hoarseness of voice and mild respiratory distress.

Grade II: LTB with moderate respiratory distress.

Grade III LTB with severe respiratory distress requiring endotracheal intubation.

Grade IV: LTB with severe respiratory distress requiring endotracheal intubation and with associated congestive cardiac failure (CCF).

About 13.9% of the patients were admitted to TCU in LTB grade IV (Table 4).

Only 24.4% of the patients had blood gas analysis (BGA) done on admission to TCU and about 12.9% of these BGAs showed respiratory alkalosis (Table 5).

About 41.0% of the patients had no serial BGAs done in ICU. A total of 589 blood samples were taken for BGA in about 66% of the patients admitted (Table 6).

Most of the LTB patients studied had one or more factors complicating their disease. Bronchopneumonia and congestive cardiac failure (CCF) were the main complicating factors occurring in 51.6% and 13.9% of the patients respectively. Most of these factors played an important role in the final prognosis of the LTB patients while others did not contribute to the ICU management and prognosis (Table 7).

Only about 12.0% of the patients needed no airway management in form of endotracheal intubation, tracheostomy or both. These patients were admitted in LTB Grade I and did not progress to LTB Grade II (Table 8). 86% of the patients had nasotracheal intubation done, the duration of intubation ranged from one day to 54 days, average 21.5 days (Table 9 and Figure 2).

Complications of nasotracheal intubation included laryngeal stenosis, laryngeal papilloma, blockage, removal or dislodging of the tube (Table 10). The duration of nasotracheal intubation in 6 patients who had laryngeal stenosis ranged from 7 to 31 days. No relationship was found between endotracheal intubation and development of laryngeal stenosis.

Indications for tracheostomy were failed intubation, long duration of intubation, laryngeal stenosis and obstructed type of breathing even after endotracheal intubation. Only 16 patients (7.7%) had tracheostomy done (Table 11).

Tracheal stenosis, Laryngeal papilloma surgical emphysema, bleeding and pneumothorax were some of the complications of tracheostomy (Table 12). Although nasotracheal intubation was done in about 86.61 of the patients compared to about of the patients who had tracheostomy done, only about 3.3% of intubated patients developed laryngeal stenosis while 25% of tracheostomy patients developed tracheal stenosis. Most of the patients developed various complications which influenced the ICU management and prognosis. Bronchopneumonia and were the main complications occurring in 21.8% and 22.51 of the patients respectively (Table 13).

All patients admitted to TCU were put on at least one systemic antibiotic. The decision on which antibiotics to put a patient on depended on many factors: (i) which year the child was admitted to ICU: in early 1970s most of the children were empirically started on Ampicillin and another broad-spectrum antibiotic. In 1980s the use of crystalline penicillin/Gentamycin combination has nearly become routine treatment.

(ii) Clinical judgement: If a patient was not responding to the antibiotics already started on in the wards admitting LTB patient to ICU the antibiotics were stopped and 3 broader and clinically more specific antibiotic started.

(iii) Culture and sensitivity tests of tracheal aspirate, blood samples etc.

(iv) Availability of the antibiotics eg. crystalline penicillin was used in 51.71 of the patients while a newly introduced antibiotic cefotaxime (Claforan) was used in 10 of the patients (Table 14).

Tetracycline eye ointment was the only onthalmic antiobioec used in 14.4% of the patients.

16% out of 209 patients (about 16%) were put on steroids . About 61.6% of the patients treated with steroids were treated with hydrocortisone (Table T51 .

The use of racemic adrenaline for treatment of acute IT? patients in our TCII has declined. In 1975 12 out of 52 patients admitted to Idtwith severe LTP were treated with racemic adrenaline using a nebulizer and intermittent positive pressure breathing (IPPB) while in 1984 35 patients were admitted to Id' with severe LTB, of which only 2 (5.7%) were treated with racemic adrenaline (Table 1b).

The range of other drugs used is shown in table I?. The drugs were used to (i) treat complications of LTP admission and those arising, in ICU eg CO₂, convulsions, hyperpyrexia etc.

- (ii) To facilitate artificial ventilation eg Diazepam.
- (iii) To facilitate removal of secretions eg Bromhexime - a mucolytic agent.
- (iv) For nutritional support eg parenterovite I and II, Trynhosan' and sorbitol.

Only 4 out of 209 (1.9%) patients had hypokalemia on admission which was treated with potassium chloride supplements. One patient had hyponatremia on admission which was corrected by half-strength darrows drip. Few patients had abnormal serum electrolyte results while in ICU (Table 18) .

The incidence of anemia was low . A total of 74 hematocrits were analysed. The average packed cell volume (PCV) was 32.5% .

All blood transfusions were done in patients with PCV less than

301

⁷ patient? were discharged alive from TCI' after nasotracheal intubation and successful extubation. The decision on when to extubate a patient was based mainly on clinical grounds (no sign of lower respiratory tract infection, acceptable PCV, good cough reflex) rarely supported by BGA results. Most of the patients had a successful first trial extubation (Table TP) .

A total of 124 tracheal aspirate cultures were done. The commonest organism cultured was Klebsiella (Table). The organisms cultured from tracheal aspirates were resistant to commonly used antibiotics. Klebsiella was mainly sensitive to Gentamycin, Kanamycin and Polymyxin B; Pseudomonas mainly sensitive to piperacillin, Gentamycin and polymyxin B, staphylococcus mainly sensitive to cephalosporins, cloxacillin and erythromycin and nocardia mainly sensitive to Gentamycin. Most of these organisms were resistant to crystalline penicillin and Ampicillin .

The duration of ICU stay ranged from 1 to 83 days with a mean of 11.2 ± 7.6 days. About 10.5% of the patients studied had one day stay in ICU (Table 21 and Figure 3). 94 out of the 209 patients admitted to ICU died. This gives a mortality rate of 45%. Bronchopneumonia and CCF were the main causes of death (Table 22).. The differences in prognosis of the LTB patients corresponding with LTB grade on admission were statistically significant. Late admission to ICU i.e. LTB. Grade IV was associated with poor prognosis (Table 23) .

DISCUSSION

Acute laryngotracheobronchitis (LTB), laryngotracheitis (LT) and croup are sometimes used to describe slightly different clinical conditions (2,16). In this study acute LTB, viral LT and croup were used synonymously.

Various male:female (M:F) ratios in acute LTB have been reported. Massicotte et al in 1973 gave M:F ratio of 2:1. The M:F ratio of 1.8:1 got in this study confirms the high occurrence of LTB in males. The widest age range given for LTR is 6 months to 14 years (13). The range in this study was 2 months to 156 months (13 years) which shows that LTB can occur at an earlier age. About 50% of the patients were 2 years or less. Therefore, the age distribution is as expected of a disease of early childhood. However, the average age in this study of (30.9 months) is higher than Dansky's figure of 17.7 months already thought to be high (50),

Mild to moderate LTB patients are treated in General paediatric wards. It is therefore, not surprising that most of the patients in the study were from Paediatric Observation ward (P.O.W) although rarely LTB can present with acute onset of symptoms mimicking a foreign body in the respiratory tract. Such patients are therefore, admitted to Ear, Nose and Throat (ENT) wards and may even have direct laryngoscopy or bronchoscopy done before diagnosis of LTB is made as seen in this study.

The decision as to when to transfer LTB patient to K. F. H. I. C. V. depended mainly on clinical grounds backed rarely by blood gas analysis (BGA) results. About 73.7% of the patients were admitted in LTB Grade III and were intubated or had tracheostomy done. About 12. and 13.9% were admitted in Grade II and IV respectively. This late admission to I. C. V., was perhaps a major contributing factor to poor prognosis.

The aetiology of LTB in developing countries is different from that in developed countries where parainfluenza viruses 1 and 2, Influenza A and adenovirus are the main agents (1, 31, 39, 40, 51, 60, 59). Dansky (1978) in his South African study of I. C. U. - treated LTB patients (10) had 135 postmeasles patients. Wesley (67) (South Africa) quotes 75% postmeasles LTB. In this study about 66% of the patients had postmeasles LTB which further confirms that measles is still the commonest cause of LTB in developing societies. There are many factors which adversely affect the survival of LTB in developing countries. Lower respiratory tract (LRT) involvement is common (about 73.7%) even in up to 66% of the patients (67). Congestive Cardiac

Failure (CCK) or symptomatic myocarditis, diarrhoea, malnutrition (50, 57-70) and iron deficiency anaemia, (H0, 8-11) are some of the common premonitory complications. Therefore, the occurrence of bronchopneumonia in 31.6% of the patients and CCF (13.9%) and other multiple complicating factors in this study reflect the typical picture of LTB expected in a developing country.

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Blood gas analysis (HCA) done on admission showed that about 19% of the patients had acceptable results, about 2.9% had respiratory alkalosis and only 7.9% showed respiratory acidosis. These findings throw some doubts on the usefulness of BGA in assessing severity of LTB or making a decision as to when to admit a patient to I.C.U. or intubate a severe LTB patient: this point has been stressed by various authors (3, 25, 50, 67-68, 82).

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Electrolyte imbalance in severe LTB patients on admission has not been documented, although it is known that diarrhoea and dehydration are frequent complications. Our patients were routinely put on half-strength Darrows drip in their admitting wards. These intravenous infusions were continued in I.C.U. and nasogastric tube feeds were started when it was thought the child could tolerate them. Therefore,

only 2.4; and 8.2% of the patients had significant electrolyte imbalances on admission to ICU and while in ICU respectively. The ICU management of severe LTP patients in KNH included respiratory support, Oxygen, steam, antibiotics, steroids, racemic adrenaline and management of complicating factors on admission and those arising in ICU.

All the patients in this study were given oxygen and about 22% required endotracheal intubation and T-piece oxygen. About 59.3% had endotracheal intubation, steam and oxygen tent. The use of oxygen therapy is much in keeping with the already known advantages of oxygen therapy in LTP (25, 37, 50, 57). The advantages of humidification of inspired gases in treatment of LTP and hyponatraemia and water intoxication (41, 54, 66) are known; however there was no association between hyponatremia and steam therapy in the 59.3% patients who were treated with steam. About 40.9% of the patients could not be treated with steam, due to temporary unavailability of nebulizers, mechanical failure of nebulizers and or unavailability of steam tents.

Airway management: 12% of the patients were admitted in LTP Grade II and needed no respiratory support. About 86.6% had nasotracheal intubation done while only 11 had tracheostomy alone done. About 6.7% had tracheostomy and endotracheal intubation.

Wesley et al (67), Xincan et al in 1984 (26), Thomson et al, (13) and Danskv et al (50) report intubation of 67%, 121, 18 and 721 respectively.

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Since the severity of upper airway obstruction in LTP can be estimated by the need for tracheostomy or intubation, the low intubation rate in developed countries (13,261 compared with the rates in developing countries (67,50"), and the rates in this study (Table 8) further confirm that LTB in a developing country represents a very different picture from that seen in developed countries.

Endotracheal intubation does not relieve airway obstruction in LTB with associated LRT disease. Such patients require assisted ventilation or continuous positive airway pressure (CPAP) for the management of their concomitant bronchopneumonia (67). Dansky et al (50) report 24% full ventilation, 18% CPAP and 42% bronchopneumonia rates. About 29.2% of patients in this study had assisted ventilation (IPPV) and 5.3% intermittent positive pressure ventilation (TPFV) and positive end expiratory pressure (PEEP). These rates of assisted ventilation compare well with the high rate of bronchopneumonia on admission (31.6%) and during ICU stay (27.8%).

The duration of endotracheal intubation depends on the severity of /LTF, complicating factors on admission and complications arising during the management in ICU. The high rates of bronchopneumonia, CCF, diarrhoea and other multiple complicating factors during admission and arising in ICU therefore prolonged the intubation period in this study to a mean of 21.5 days (and 1 to 54 days range) compared to Dansky's 5.5 - 6.5 days (50), Thompson's 40 hours (13), Shuller and Birck's 88 hours (28). However, the rates of bronchopneumonia, CCF and diarrhoea are similar to ones reported from the two south Africa

studies (50,61). Severe oral herpes reported by the two authors was not found in this study. In KMI.TCU nasotracheal intubation is not done unless tracheostomy was indicated after failure to intubate, long duration of intubation, laryngeal stenosis and prolonged breathing even after intubation. In the early 1970s the duration of nasotracheal intubation was considered longer, in our ICU, after 2 weeks, but in the 1980s no specific period of continuous intubation was considered long. The number of techniques in our unit had also declined with time from 1977 to 1984. Serious complications of nasotracheal intubation (laryngeal stenosis in 10% and laryngeal papilloma in 7% of the patients intubated) were lower than the complications of tracheostomy (in tracheal stenosis in 25% and laryngeal papilloma in 63% of the patients who had tracheostomies). These findings are similar to those pointed out on the advantages and disadvantages of tracheostomy and intubation outlined by Wesley et al (67) and are a further proof that endotracheal intubation is safer for relief of upper airway obstruction than tracheostomy as documented by many authors (12, 28, 35, 44, 50, 67, 70-72, 83-84).

Antibiotics were used in all patients in this study. Most of the patients had LRT disease, postmeasles immunosuppression and malnutrition. The combination of crystalline penicillin (51.9%) and Gentamicin (35.0% of patients) started on admission to TCU therefore appeared justified. The organisms grown from tracheal aspirate cultures were mainly Klebsiella and Pseudomonas which are more virulent and resistant to antibiotics than organisms isolated in other studies (1,2). The resistance of these organisms to commonly used antibiotics called for the use of newer antibiotics eg. claforan. Amoxicillin which is the popular antibiotic for complicated LTB (6%) was used in only

31.61 of the patients.

The use of antibiotics is less frequent in the treatment of uncomplicated LTB seen in developed countries eg in 50% of the patients (26) while rate of use of antibiotics is high in LTP in developing countries 100% in the Fouth African study (50) .

The use of steroids in LTP is accepted by many authors (6, TP-T2, 5, 64). In this study about 78.5% of the patients were put on steroids.

The doses of steroids used eg 50 mg hydrocortisone three times daily or 10 mg twice daily prednisone, were above the effective dose in LTB recommended by Davison (27). However the study showed that 66.01 of the patients had postmeasles LTP and 25.7% had Kwashiorkor. The use of steroids in 78.51 of the patients with such a high incidence of immunosuppressing factors is questionable (5, 67).

About 17.71 of the patients had racemic adrenaline and IPPB . The usefulness of racemic adrenaline in LTB is controversial (1, 4, 7, 10, 11, 13, 15, 26, 38, 45, 50, 42-43, 61). Considering that most of the patients had LRT disease and CCF with associated tachycardia, the use of racemic adrenaline in our unit may have had more disadvantages than advantages (ICU consultants personal communication) and hence the decline in its use in treatment of LTB in our unit.

Three patients were treated with subcutaneous adrenaline and two with nebulised isoprenaline. The usefulness of these drugs in LTP, or its complications, has not been established. "Sedation with a small dose of diazepam reduces unnecessary agitation and wasted energy and does not reduce respiratory effort or mask the physical signs monitored for the degree of obstruction (67)".

This fact was reflected in this study where about 49.31 of the patients were given diazepam mainly to facilitate artificial ventilation. Chest physiotherapy with pharyngeal suction though recommended at least twice daily (6") was done more frequently, even sometimes half hourly, in our unit. Most of the other drugs used in ICU were used to treat specific complication but the use of a few drugs cannot pass without challenge: Bromhexine (Bisolvon), a mucolytic agent was used in about 21.15 of the patients; terbutaline (Bricanyl) in about 13.9, and salbutamol (Ventolin) in about 12.0% of our patients. The advantages of using Bromhexine over the ensuring of adequate hydration of the patient and humidification of inspired gases have not been established (56). Salbutamol and Terbutaline are selective B-2 agonists with little action on B1 receptors. Considering that Bronchopneumonia and CCF with associated tachycardia were present in 59.4% and 36.4% of the patients in IGI respectively, the usefulness of these bronchodilators, which have not been shown to have any effect in LTB and which may even worsen the tachycardia, is doubtful. The use of propranolol to treat tachycardia caused by respiratory failure as was the case in 7 of our patients, should be discouraged. Lomotil and Kaolin sedative which were used a lot in the 1970s to treat diarrhoea have been banned due to their side-effects in children. Newer drugs like (Loperamide) Loperamide are now used to treat severe diarrhoea associated with postmeasles LTB, which is not adequately treated with intravenous infusions alone.

Pyrexia in LTP children is associated with increase in pulse and respiratory rate, cardiac output, gas exchange and oxygen consumption. Acetyl salicylic acid (Aspirin) was used a lot in early 1970s in community to control pyrexia, but due to increase in incidence of aspirin poisoning, Indomethacin (Indocin) suppositories, which are more convenient to use in children and is relatively safe, was widely used (in 46. of patients).

The decision as to when to extubate the patient has no hard and fast rules (50,67) though various suggestions have been made, based on clinical grounds, satisfactory BGA results and scoring system. Most of the patients in this study had Bronchopneumonia, CCF and other complications and therefore trial extubation after 2-4 days of intubation suggested for uncomplicated LTB by Wesley et al (67) and Schuller and Birck (28) could not be tried. About 91.81 of the patients in this study who had been intubated and were finally discharged from ICU, had first successful trial extubation. This gives a lower first extubation failure rate of 8.21 compared to Schuller and Birck's reintubation rate of 171 (28).

The duration of ICU stay probably reflects the severity of LTB and its associated complications. The duration is also shorter with endotracheal intubation than with tracheostomy (28). The duration of ICU stay was 6.7 ± 4.3 days in Dansky's South African study compared to 3.9 days in IXmeans USA study (28) and other studies 2.1 days (15) 34.4 ± 17.7 hours (9) 3.45 days (10) therefore the average duration of ICU stay of 11.2 days got in this study is, as expected, long for LTB ICU managed patients in a developing country.

The mortality rate of severe LTB is higher in developing countries (150, 47, 7%) than in developed societies whose mortality rate is less than 2% (12, 25, 26, 4%, 74). The highest mortality rate reported from a developing country is 161 (50). Therefore the mortality rate of 451 got in this study is extremely high. Bronchopneumonia and CCF, which were the main complicating factors on admission and also the main complicating factors arising in ICU were also the main causes of death. There was no death due to LTB alone. These findings are similar to the findings on studies done on severe LTB in developing countries (50, 67). All the findings in this study confirm Wesley's findings that LTB in developing countries is a more severe disease than that managed in developed countries.

OCCLUSION

Laryngotracheobronchitis patients formed the majority of admissions in ICU during the period of study, and their mortality was very high compared to similar studies done elsewhere. Many of these patients came to the hospital rather late for treatment. Many of these developed as a complication of measles and were associated with malnutrition and belonged to the low socio-economic group of our population.

It is therefore suggested that mass immunisation campaigns against measles be stepped up in all pre-school age children, and that general education on nutrition be encouraged. It is also suggested that the severe forms of this disease be identified and diagnosed early, for early admission to ICU and treatment, before complications arise.

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VABLE L

SEX DISTRIBUTION

SEX	NO. OF PATIENT	PERCENTAGE %
MALES	134	64.1
FEMALES	75	35.9
TOTAL	209	100

FIG, 1

AGE DISTRIBUTION

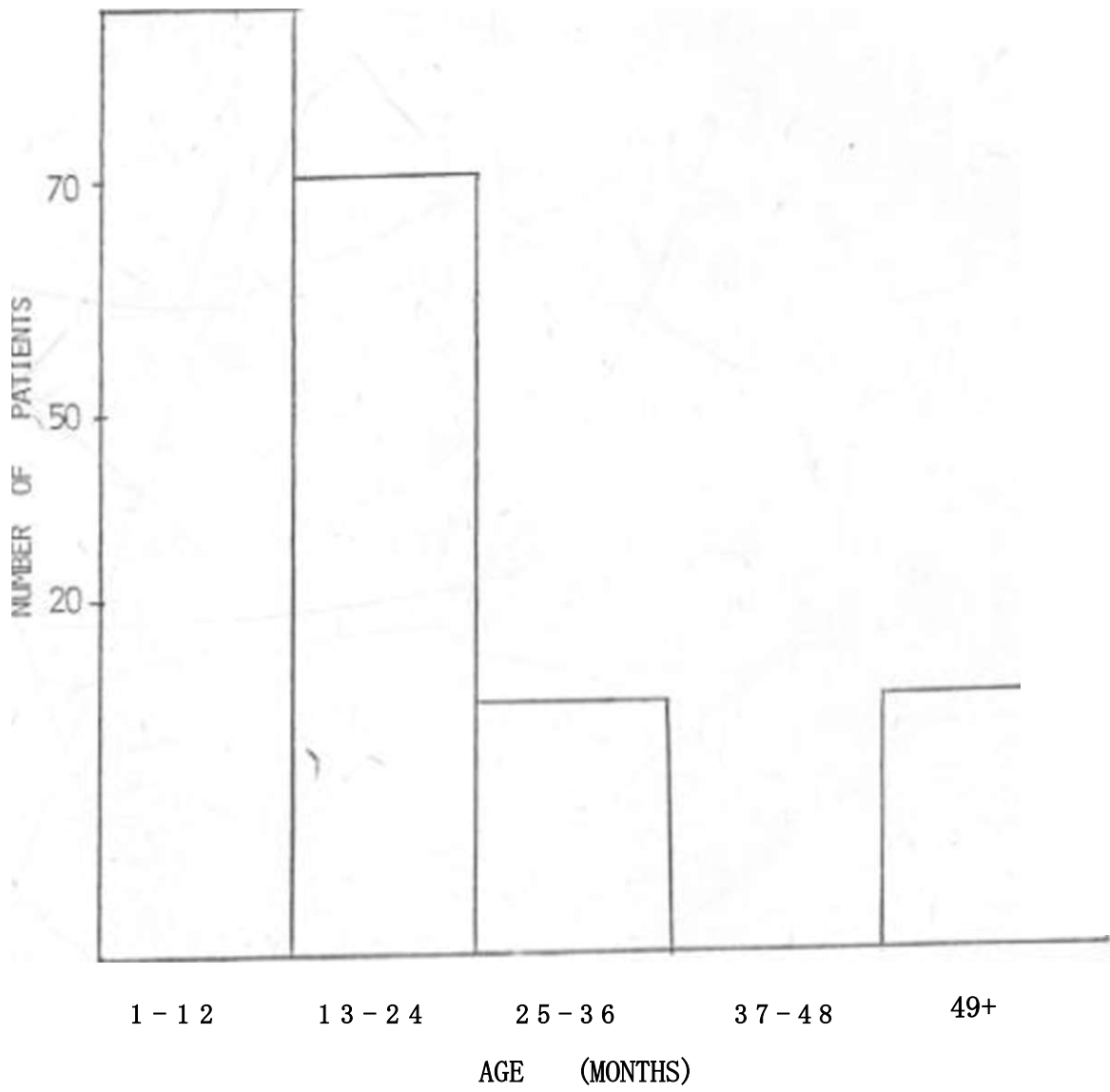


TABLE 2
THE AGE DISTRIBUTION

AGE ·MONTHS	NO. OF PATIENTS	PERCENTAGE (%)
1 - 12	92	IFL.0
13 - 21	75	35.9
25 - 36	21	10.0
37 - 48	6	2.9
	15	7.2
TOTAL	209	100%

TABLE 3

CAUSES OF LTB

LTB CAUSES	NO. OF PATIENTS	PERCENTAGE
POST-MEASLES	138	55.0
VIRAL	71	34.0
TOTAL	209	100%

%

TABLE ^

LTB GRADE ON ADMISSION

LTB GRADE	NO. OF PATIENTS	PERCENTAGE
I	0	0
II	25	12.4
III	154 <i>i</i>	73.7
IV	29	13.9
TOTAL	209	100%

TABLE 5.

RESULTS OF BGAs DONE ON ADMISSION

BGA RESULT	NO. OF PATIENTS	% OF BGAs (DONE ON' ADMISSION)
RESPIRATORY ALKALOSIS	27	42.9
METABOLIC ACIDOSIS	7	11.1
RESPIRATORY ACIDOSIS	5	7.9
HYPOXIA	6	9.5
ACCEPTABLE	19	30.2

Respiratory alkalosis i.e. $P_a CO_2$ less than

Respiratory acidosis i.e. CO_2 more than 40mmHg.

Metabolic alkalosis or acidosis depending on FH, Base Excess or Deficit, serum bicarbonate (HCO_3^-) levels.

Hypoxia: $P_a O_2$ less than 40mmHg with patient on ROOM air or less than 60mmHg with the patient on oxygen.



f

TABLE 6

RESULTS OF SERIAL BGA

BGA RESULT	NO. OF PATIENTS	% OF SERIAL BGAS DONE
RESPIRATORY ALKALOSIS	135	21.2
ACCEPTABLE	113	19.2
HYPOXIA	101	17.1
RESPIRATORY ACIDOSIS	60	10.2
METABOLIC ALKALOSIS	58	9.8
METABOLIC ACIDOSIS	40	6.8

TABLE 7

COMPLICATING FACTORS ON ADJUSTMENT TO 1, C. U.

COMPLICATION	NO. OF PATIENTS	% OF PATIENTS IN STUDY
BROCHOPNEUMONIA	66	31.6
CONGESTIVE CARDIAC FAILURE (CCH)	29	13.9
DIARRHOEA	21	10.0
DEHYDRATION	19	9.1
OTITIS MEDIA	12	5.7
KWASHIORKOR	11	5.3
STOMATITIS	9	4.3
ANAEMIA	6	2.9
MARASMS	5	2.4
CONVULSIONS	4	1.9
HYPERPYREXIA		
CARDIAC ARREST (ONE)	3	1.4
BURNS		
SEPTICAEMIA	2	1.0
OTHERS	1	0.5

TABLE 8

AIRWAY KANAGEKErT ATO RESPIRATORY SLP PORT

RESPIRATORY SUPPORT	NO. OF PATIENTS	PERCENTAGE OF PATIENTS STUDIED
NO INTUBATION OR TRACHEOSTOMY DONE	25	12.0
OROTRACHEAL INTUBATION	1	0.5
NASOTRACHEAL INTUBATION	181	86.6
TRACHEOSTOMY ALONE DONE	2	1.0
TRACHEOSTOMY AND NASOTRACHEAL INTUBATION	14	6.7
ENDOTRACHEAL INTUBATION AND T-PIECE OXYGEN	46	22.0
ENDOTRACHEAL INTUBATION/ STEAM AND OXYGEN TENT	124	59.3
MECHANICAL VENTILATION X IPPV	61	29.2
IPPV + PEEP	11	5.3

IPPV - Intermittent positive pressure ventilation

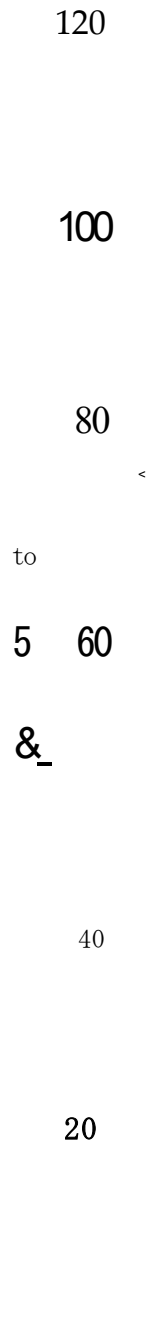
PEEP - Positive End Expiratory Pressure.

TABLE 9
DURATION OF NASOTRACHEAL INTUBATION

DURATION OF NASOTRACHEAL INTUBATION (DAYS)	NO. OF PATIENTS	% OF PATIENTS INTUBATED
1 - 5	76	42.0
6 - 10	53	29.0
11 - 20	40	22.1
21 - 30	9	5.0
31 - 40	2	1.1
41+	1	0.6
TOTAL	181	100%

FIG. 2

DURATION OF NASOTRACHEAL INTUBATION



DURATION OF NASOTRACHEAL I INTUBATION (DAYS)

TABLE 10.

COMPLICATIONS OF NASOTRACHEAL INTUBATION

COMPLICATION	NO. OF PATIENTS	% OF PATIENTS
LARYNGEAL STENOSIS REQUIRING DILATATIONS	6	3.3
LARYNGEAL PAPILLOMA	2	1.1
BLOCKAGE (AT LEAST ONCE)	44	24.3
REMOVAL OR DISLODGED TUBE (AT LEAST ONCE)	45	25.4

TABLE 11

INDICATIONS FOR TRACHEOSTOMY

INDICATION	NO, OF PATIENTS
FAILED INTUBATION	2
LONG DURATION OF INTUBATION	6
LARYNGEAL STENOSIS	<i>H</i>
OBSTRUCTED TYPE OF BREATHING EVEN AFTER ENDOTRACHEAL INTUBATION	<i>i</i> 3
? FOREIGN BODY	1

TABLE 12
r ;

COMPLICATION OF TRACHEOSTOMY

COMPLICATION	NO. OF PATIENTS	% OF PATIENTS WITH TRACHEOSTOMY
TRACHEAL STENOSIS	8	25 .0
LARYNGEAL PAPILLOMA	1	6.3
SURGICAL DEBRIDEMENT	1	6.3
BLEEDING	1	6.3
PNEUMOTHORAX DUE TO DISPLACED TRACHEOSTOMY TUBE	1 «	6.3
BLOCKAGE (AT LEAST ONCE)	1	6.3

TABLE 13

COMPLICATIONS ARISING IN I. C. U.

COT-PL I CATION	NO, OF PATIENTS	% OF PATIENTS IN STUDY
BRONCHOPNEUMONIA	58	27.8
CONGESTIVE CARDIA FAILURE	47	22.8
DIARRHOEA	20	9.6
COTMJSIONS	19	9.1
PERSISTENT FEVER	7	3.3
LUNG COLLAPSE	5	2.4
MENINGITIS, PNEUMOTHORAX	4	1.9
OTITIS MEDIA		
SEPTICAEMIA		
^r ENCEPHALITIS	> 2	1.0
ORAL THRUSH		

TABLE 1M

SYSTEMIC ANTIBIOTICS USED

ANTIBIOTIC	NO. OF PATIENTS	% OF PATIENTS IN THE STUDY
CRYSTALLINE PENICILLIN (CRYSTAPEN)	108	51.7
GENTAMYCIN	73	35.0
AMPICILLIN	66	31.6
CHLORAMPHENICOL	23	11.0
KAMAKYCIN	23	11.0
AF-PICLOX	21	10.0
SEPTRIN	18	8.6
CLOXACILLIN	11	6.7
CEPHALOSPORIN	8	3.8
AMIKACIN	7	3.3
ANTI TUBERCULOUS THERAPY (STREPTOMYCIN, THIAZINA, INH)	6	2.9
PYOPEN	<4	1.9
LINCOMYCIN	3	<i>IA</i>
ERYTHROMYCIN		1.0
CEFOTAXIME (CLAFORAN)		
CEFATREXYL (DALACIN C)		
FLUCLOXACILLIN SODIUM (FLOXAPEN)		0.5
PROCAINE PENICILLIN		
TRIPLOPEN		
AMOXYCILLIN		

TABLE 14

SYSTEMIC ANTIBIOTICS USED

ANTIBIOTIC	NO. OF PATIENTS	% OF PATIENTS IN THE STUDY
CRYSTALLINE PENICILLIN (CRYSTAPEN)	108	51,7
GENTAMICIN	73	35,0
AMPICILLIN	55	31,6
CHLORAMPHENICOL	23	11,0
KANAMYCIN	23	11,0
AMPICLOX	21	10,0
SEPTRIN	18	8,6
CLOXACILLIN	14	6,7
CEPHALOSPORIN	8	3,8
AMIKACIN	7	3,3
ANTI TUBERCULOUS THERAPY (STREPTOMYCIN, THIAZINA, INH)	6	2,9
PYOPEN	4	1,9
LINCOMYCIN	3	1,4
ERYTHROMYCIN		
CEFOTAXIME (CLAFORAN)	2	1,0
CEFATREXYL (DALACIN C)		
FLUCLOXACILLIN SODIUM (FLOXAPEN)		
PROCAINE PENICILLIN	1	0,5
TRIPLOPEN AMOXYCILLIN		

TABLE 15

STEROIDS USED

TYPE OF STEROID	NO. OF PATIENTS	% OF TOTAL PATIENTS OF J STEROIDS
HYDROCORTISONE	101	51.6
PREDNISONE	36	22.0
DEXWCTHASONE	24	14.6
SOLLJMEDRAL (KETHYIPREDNISOLCNE)	3	1.8
TOTAL	164	100%

TABLE 16

USE OF RACEMIC ADRENALINE

	NO I OF PATIENTS	* OF PA<I,J?^ IN THE *
RACEMIC ADRENALINE IPPB	37] JJ
SUBCUTANEOUS ADRENAL I ^		
NEBULIZED ISOPRENALINE		

TABLE 17

<u>OTHER DRUGS USED</u>		
DRUG	NO. OF PATIENTS	% OF PATIENTS IN STUDY*
		49.3
DIAZEPAM	103	31.1
DIGOXIN	65	22.0
INDOCID (SIPPOSITORIES)		21.1
BROWEXIME (BISOLVON)	^	
FRUSEMIDE (LASIX)	30	iu. »
PARENTENOVITE I AT ^o II	30	13.9
TERBUTALINE (BRICANYL)	v 29	12. "
CHLOROQUINE	26	12.0
SALBUTAMOL (VENTOLIN)	25	10.0
DIPHENOXYLARE + ATROPINE (LOMOTIL)	21	9.*
PRCMEVIAZINE (PHENERGAN)	20	8 ⁶
ACETYL SALICYLIC ACID (ASPIRIN)	18	11
MULTIVITE	16	

TABLE 17

OTHER DRUGS USED

DRUG	NO. OF PATIENTS	% OF PATIENTS IN STUDY
DIAZEPAM	103	49.3
DIGOXIN	65	31.1
INDOCID (SUPPOSITORIES)	46	22.0
BRONHEXIME (BISCLVON)	44	21.1
FRUSEMIDE (LASIX)	30	14.4
PARENTE NOVITE I AND II	30	14.4
TERBUTALINE (BRICANYL)	29	13.9
CHLOROQUINE	26	12.4
SALBUTAMOL (VENTOLIN)	25	12.0
DIPHENOXYLARE + ATROPINE (LOMOTIL)	21	10.0
PROMETHAZINE (PHENERSAN)	20	9.6
ACETYL SALICYLIC ACID (ASPIRIN)	18	8.6
MJLTIVITE	16	7.7

TABLE 18

RESULT	NO. OF PATIENTS	% OF PATIENTS IN STUDY
(< 130 MF-DL/L) HYPONATRAEMIA		1.0
KYPONATRAEMIA (>150 FTLCU L)		0.5
(< 3.0 MMOL/L) HYPOKALAEMIA		4.3
HYPERKALAEHIA (> 5.0 MMOL/L)		2.4
TOTAL	17	8.2

TABLE 19

SUCCESSFUL TRIAL EXTUBATION

SUCCESSFUL TRIAL EXTUBATION	NO- OF PATIENTS	% OF PATIENTS INTUBATED AND DISCHARGED ALIVE
FIRST	89	91.8
SECOND	3	3.1
THIRD	2	2.1
FOURTH	1	1.0
FIFTH	2	2.1
TOTAL	97	100



TABLE 20

RESULTS OF CULTURE OF TRACHEAL ASPIRATES

ORGANISM ISOLATED	no-	OF PATIENTS	% OF CULTURES
KLEBSIELLA		35	29.0
PSEUDOMONAS		27	21.8
STAPHYLOCOCCUS AUREUS		21	17.0
PROTEUS		12	9.7
CITRIBACTER		9	7.3
E. COLI		6	4.8
STAPHYLOCOCCUS ALBUS			3.2
ENTEROBACTER		3	2.4
ACINETOBACTERIA		2	1.6
ALKALIGENES FAEC/U-IS		1	0.8
PNEUMOCOCCI		1	0.8
P - HEAMOLYTIC		1	0.8
STREPTOCOCCUS		1	0.8

TABLE 21

DISTRIBUTION OF 1 C.U. STAY

STAY IN 1. C. U. (DAYS)	NO. OF PATIENTS	PERCENTAGE
1 - 5	74	35.4
6 - 10	58	27.8
11 - 20	53	25.4
21 - 30	14	6.7
31 - 40	3	2.9
41 - 50	0	0
51 +	4	1.9
TOTAL	209	100%

FIG. 3 MIRATIOK CP I. C. U. STAY

140

120

100-

80

I

60

40

20_

j _____ i

1 - 10

11 - 20

21 - 30

31 - 40

41 - 50

51+

I. C. U. STAY (DAYS)

I

I

TABLE 22

CAUSLS OP CEATH

CAUSE		NO. OF PATIENTS	% OF PATIENTS WHO DIED
BRONCHOPNEUMONIA AND CCF		86	91.5
HYPOXIC BRAIN DEATH	<i>I</i>	5	5.3
MARASMUS		2	2.1
OTHERS	/	1	1.1
TOTAL		94	103%

KANAFIEKgv'T OF LT? PATIENTS IN K.N.H. .1.C.C.

197? - 198^.

IATA COLLECTION FORM:

STUDY NO:

NAME I. P. NO

LTB GRADE ANT CAUSE:

A.IE ZZX WEIGHT

A. :TTL. FRO

ADMISSION D>TE

CISCRAJ^E/DEATH DJTE

DURATION OF I. C. U. STAT(DAYS)J

COMPLICATING /ACTORS (ON ADMISSION)

4

COMPLICATIONS IN I. C. U

1.1.V. I *N QEMEH1:

, -NTIBIOTICS

OK ADMISSION BGA:

PH

$p\text{CO}_2$

$P\text{O}_2$

SO_2

HCO_1

BE

SERIAL BOA:

DATE _____ RESULTS.

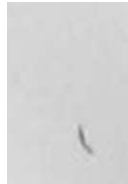
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OTHER COMMENTS: