

**NEUROLOGICAL STATUS OF
THE VERY LOW BIRTH WEIGHT
INFANTS AT SIX, NINE AND TWELVE
MONTHS OF AGE**

**A DISSERTATION PRESENTED IN PART FULFILMENT
FOR THE DEGREE OF
MASTER OF MEDICINE (PAEDIATRICS)**

AT THE UNIVERSITY OF NAIROBI

1990

BY

KIROS .M. MEKONNEN.



DEDICATION

TO MY MOTHER AND FATHER.

DECLARATION:

This is my original work and has not been presented for a degree in any other University.

SIGNED *[Signature]* Date *15th of June, 1990*.....

DR. KIROS.M MEKONNEN., M.D (ADDIS ABABA)

This Dissertation has been submitted for examination with our approval as University supervisors.

SIGNED... *[Signature]*

DR. A.O WASUNNA, M.B.Ch.B. M.MED (PAED), NEONATOLOGIST,
SENIOR LECTURER, DEPARTMENT OF PAEDIATRICS AND CHILD HEALTH,
UNIVERSITY OF NAIROBI.

DATE : *June 15 1990*

SIGNED *[Signature]*

DR. R.N. MUSOKE, M.B.Ch.B. M.MED (PAED), NEONATOLOGIST,
SENIOR LECTURER, DEPARTMENT OF PAEDIATRICS AND CHILD HEALTH,
UNIVERSITY OF NAIROBI.

DATE: *15th of June, 1990*.....

LIST OF TABLES

TABLE I.	AGE AND SEX DISTRIBUTION OF THE VLBW INFANTS VERSUS THE CONTROLS.	P. 10
TABLE II.	GESTATIONAL AGE DISTRIBUTION OF THE VLBW INFANTS	P. 12
TABLE III.	FIVE MINUTES APGAR SCORE DISTRIBUTION OF THE VLBW INFANTS	P. 13
TABLE IV.	DISTRIBUTION OF NEONATAL MORBIDITIES OF THE VLBW INFANTS	P. 14
TABLE V.	SUMMARY OF NEUROLOGICAL SEQUELAE OBSERVED	P. 16
TABLE VI.	SUMMARY OF THE VLBW INFANTS WITH NEUROLOGICAL SEQUELAE	P. 17
TABLE VII.	RISK FACTORS VERSUS NEUROLOGICAL SEQUELAE	P. 18
TABLE VIII.	MEAN WEIGHTS OF THE VLBW MALE INFANTS VERSUS THE FEMALES	P. 21
TABLE IX.	MEAN LENGTHS OF THE VLBW MALE INFANTS VERSUS THE FEMALES	P. 21
TABLE X.	MEAN WEIGHTS OF THE VLBW INFANTS VERSUS THE CONTROLS	P. 22
TABLE XI.	MEAN LENGTHS OF THE VLBW INFANTS VERSUS THE CONTROLS	P. 25
TABLE XII.	MEAN HEAD CIRCUMFERENCES OF THE VLBW INFANTS VERSUS THE CONTROLS	P. 28
TABLE XIII.	DISTRIBUTION OF MATERNAL AGE OF THE VLBW INFANTS VERSUS THE CONTROLS	P. 29
TABLE XIV.	MATERNAL DISTRIBUTION BY MARITAL STATUS	P. 30
TABLE XV.	DISTRIBUTION BY MATERNAL EDUCATION	P. 31
TABLE XVI.	DISTRIBUTION BY INCOME STATUS	P. 32
TABLE XVII.	DISTRIBUTION BY ANTENATAL CARE ATTENDANCE	P. 33
TABLE XVIII.	PATTERN OF OBSTETRIC RISK FACTORS	P. 34
TABLE XIX.	DISTRIBUTION BY PARITY OF THE MOTHERS OF THE VLBW INFANTS VERSUS THE CONTROLS.	P. 35

LIST OF FIGURES

FIG. 1 AGE AND SEX DISTRIBUTION OF THE VLBW INFANTS.....11

FIG. II MEAN WEIGHTS FOR AGE OF THE VLBW INFANTS VERSUS
THE CONTROLS.....20

FIG.III MEAN LENGTHS FOR AGE OF THE VLBW MALE INFANTS
VERSUS THE CONTROLS.....23

FIG. IV MEAN LENGTHS FOR AGE OF THE VLBW FEMALE INFANTS
VERSUS THE CONTROLS.....24

FIG. V MEAN HEAD CIRCUMFERENCES OF THE VLBW MALE INFANTS
VERSUS THE CONTROLS.....26

FIG. VI MEAN HEAD CIRCUMFERENCES OF THE VLBW FEMALE INFANTS
VERSUS THE CONTROLS.....27

ABBREVIATIONS

1. A.N.C Antenatal care
2. A.P.H Antepartum haemorrhage
3. B.W Birth weight
4. C/S Caeserian section
5. F.E.T Fisher's exact test
6. G.A Gestational age
7. K.N.H Kenyatta National Hospital
8. N.C.H.S National centre for health and statistics
9. P.R.O.M Prolonged rupture of membrane
10. P.E.T Preeclampsia toxemia
11. P.V/I.V.H Periventricular/Intra ventricular haemorrhage
12. R.R Relative risk analysis
13. R.D Respiratory distress
14. S.V.D Spontaneous vertex delivery
15. V.L.B.W Very low birth weight
16. W.H.O World health organization
17. 95% C.I 95% confidence interval

CONTENTS

DECLARATION.....	i
LIST OF TABLES.....	ii
LIST OF FIGURES.....	iii
LIST OF ABBREVIATIONS.....	iv
SUMMARY.....	vi
INTRODUCTION.....	1
STUDY OBJECTIVES.....	6
MATERIALS AND METHODS.....	6
RESULTS.....	10
DISCUSSION.....	36
CONCLUSIONS.....	42
RECOMMENDATIONS.....	43
ACKNOWLEDGEMENTS.....	44
REFERENCES.....	45
APPENDIX I	55
APPENDIX II.....	59
APPENDIX III.....	64
APPENDIX IV.....	65
APPENDIX V.....	66

SUMMARY

A cross sectional study on neurological status was conducted on 73 VLBW infants between January to December 1989 inclusive at the KNH. These were compared with age and sex matched normal term infants. The prevalence of neurological sequelae among the VLBW infants was 8.2%, and none of them had major sequelae. All the infants with neurological sequelae had one or more preconceptual, prenatal, perinatal and neonatal risk factors which might have been related to their neurological outcome. The prevalence of obstetric risk factors among the VLBW infants was 73.34%. PROM (39.7%) followed by history of abortion (32.9%) were the commonest factors encountered in this study group. None of the normal term infants included into the study as controls had neurological sequelae or obstetric risk factors.

INTRODUCTION

Very low birth weight infant is defined by W.H.O as an infant who weighs 1500 grams or less at birth (1). These infants are further subclassified into three categories, based on their birth weight and gestational age, as appropriate, small and large for their gestational age. (2). Very low birth weight infant is small because of a short period of gestation, intrauterine growth retardation or both. The importance of classifying these babies is because their neonatal morbidities, survival rate and subsequently their long term physical growth and neurobehavioural outcome are largely determined by their gestational age and birth weight (2-6).

The incidence of very low birth weight infants and low birth weight in general parallels, the socio economic status of a community, such that the lower the socio economic status, the higher the incidence of very low birth weight infants. In developed countries, the incidence of VLBW is reported between 1 to 1.5% of total births (6,7) while Lekha (1989), has reported from Kenyatta National hospital to be of 4.7% of total births (8). Very low birth weight infants comprise a high proportion of babies admitted to a newborn unit. Kasirye (1984), reported that 24.5% of infants admitted to the KNH newborn unit were VLBW. (9)

Infants of VLBW are at greatest risk of high morbidity and mortality from respiratory distress syndrome, recurrent apnoeic attacks, periventricular/Intraventricular haemorrhage, metabolic

derangements like hypoglycaemia and hyperbilirubinaemia, feeding intolerance, infections, circulatory and thermal instability related to their immaturity. They are also at greatest risk for long term neurobehavioral abnormalities. They tend to have more post neonatal medical problems, particularly recurrent respiratory infections with frequent hospitalization. (10) Therefore the cost towards the care of the VLBW infants from the beginning is enormous in terms of money, time, and skilled manpower.

Before the last three decades the overall survival rate of the VLBW babies was less than 50% with a mortality rate for those less 1000 grams being almost one hundred percent. During the pioneering of the neonatal intensive care units (1963-1970), the survival rate of the VLBW babies started increasing to about 50-60%. However, as the survival rate increased, there were alarming reports of a high prevalence (40-60%) of neurological sequelae among the survivors as evidenced by data on long term follow up (5,10,11). During the last two decades the survival rate of the VLBW infants has increased markedly, longitudinal studies also show that the proportion of survivors with long term major neurological sequelae has decreased from 40% to 60% to about 10-25% (4,11-17). Horbar et al. (18) in 1984 studied 1776 infants in eleven neonatal intensive care units in USA and United Kingdom and showed that the survival rate during the first 28 days of life was 85%, with a range of 80-95%.

In developed countries with neonatal intensive care centres, the current status of the VLBW infants is very encouraging. Even the extremely low birth weight babies have a survival rate of about 50-70% presently (11,13). This improvement in survival rate and lower prevalence of neurological sequelae among the survivors is not only due to an advanced obstetric and neonatal care, but also due to a change of socio-economic status, demographic shifts, better maternal education, improved antenatal care, and good nutritional status of the mothers (4,12,13,19,20,21,). At Kenyatta National hospital, Kasirye (9) reported in 1984 that 96.2% of 80 babies of less than 1000 grams and 51.3% of 150 infants with birth weight of 1000-1500 grams admitted to the newborn unit died during the neonatal period.

The overall survival rate was 33.4%. Mati et al. (1983) reported in the Nairobi birth survey that the twenty four hours perinatal mortality rate of the VLBW babies was 75%. (22). VLBW infants as mentioned earlier, are at greatest risk of having long term neurological sequelae due to various prenatal, perinatal, and neonatal risk factors. Precht1 (1967), in the United Kingdom reported that out of 102 VLBW infants without neurological sequelae during neonatal period, 80% were still normal on long term follow up, while out of 150 babies with neurological sequelae, 75% were still found with sequelae on long term follow up. (23). It would appear that most neurological sequelae seen in late infancy and childhood represent most likely

a continuum from the spectrum of prenatal, perinatal and neonatal events.

One of the most important factors for the high prevalence of neurological morbidities in the VLBW infants is due to the high incidence of intracranial haemorrhage, primarily the germinal layer matrix haemorrhage (periventricular/ intraventricular haemorrhage). A serial Computerized Tomography scan and/or cranial ultrasonography done in some centres on VLBW babies in the first week of life shows an incidence of periventricular/intraventricular haemorrhage (PV/IVH) in the range of 30-50% (24). This is because the vasculature of the germinal layer matrix of the premature, VLBW infants is anatomically immature, very fragile, and uniquely vulnerable to injury, mainly to circulatory fluctuations, repeated apnoeic spells, seizures, hypercapnia and metabolic acidosis which are all associated with hypoxic-ischaemic brain damage and subsequently to periventricular / intraventricular haemorrhage.

The outcome of the PV/IVH depends on the degree and severity of the haemorrhage . In grade I and II haemorrhages, the majority of them resolve or remain with minor neuromotor sequelae, while grade III and IV have 60-80% mortality rate while the survivors have a high prevalence of major neurological sequelae (24-29).

Like most organ systems, the brain of the preterm, VLBW baby is anatomically and physiologically immature. The neurologic

functions in the first six months of life are largely of subcortical level, ie, at the brain stem and spinal cord level. The cerebral hemispheres show poor differentiation of gray and white matter. Majority of the neurones are present, however, there is poor myelinization, and axonal and synaptic connections of neurones are poorly developed. Cortical functions cannot be assessed reliably during the first six months of age.

Neurological sequelae of cortical origin can subsequently be missed on assesement before the first six months of life.

Neurological assesement of the VLBW preterm infants beginning from six months of age is thus, much more reproducible and has more predictive value than of early assesement (30).

In developing countries, VLBW infants have received little attention in terms of their long term neurological outcome. Most studies reported so far have been concerned primarily with neonatal morbidity and mortality patterns of the VLBW infants.

There is no data on record concerning with their neurological outcome. The author was thus prompted to carry out a cross-sectional study on the neurological status of the VLBW infants at six, nine and twelve months of age, using a standard neurological test for each age group, in an attempt to understand the extent of the neurological morbidity among the VLBW infants cared for at the Kenyatta National hospital.

HYPOTHESIS:-

The proportion of VLBW preterm infant survivors with neurological sequelae is 20%.

OBJECTIVES:

1. To determine the neurological status of the VLBW preterm survivors at 6,9 and 12, months of age.
2. To relate some prenatal, perinatal and neonatal risk factors with neurological sequelae.

MATERIALS AND METHODS

STUDY DESIGN - A cross-sectional type of study.

SAMPLE SIZE:-

Was calculated to give a 95% confidence limit based on the formula-

$$M = \frac{Z^2 P(I-P)}{d^2}$$

where M = Minimum sample size which is 62.

Z^2 = Volume obtained from tables of standard normal distribution at 5% significance level = 1.96.

P = Anticipated prevalence of neurological sequelae.

This was taken from reports of other studies.
 d^2 = Absolute precision and anticipated on either

STUDY AREA:

Subject to approval of the protocol by the KNH research ethical committee, the study was carried out at the new born follow up clinic, Kenyatta National hospital, which is a referral centre as well as a university teaching hospital of Kenya. Infants who require follow up at the clinic are mainly those who have been admitted to the newborn nursery unit with birth weight of less than 2000 grams and some other high risk babies who require special care at the newborn nursery unit during neonatal period. The clinic is run on regular basis, once a week. An average of forty infants are attended to on every clinic day. They are reviewed at an interval of two to eight weeks as appropriate.

INCLUSION CRITERIA:

All VLBW infants who came to the clinic at ages of 6, 9 and 12 months plus or minus two weeks, corrected for gestation and who were born and had neonatal care at the Kenyatta National hospital.

EXCLUSION CRITERIA:

All VLBW infants with severe congenital malformations.

CONTROLS:

Were recruited from the KNH child welfare clinic. They were term, appropriate for gestational age infants who were born at KNH with no perinatal or neonatal morbidities and were matched for age and sex.

STUDY POPULATION:

All the VLBW infants recruited for neurological assessment.

SOURCE POPULATION:

All the VLBW infants attending the KNH, newborn follow up clinic.

REFERENCE POPULATION:

All the VLBW infants cared for at the KNH, newborn nursery unit.

The cases and controls were recruited on the same clinic days of the study period. In both groups, consecutive children fulfilling the inclusion criterias were selected.

The VLBW infants who were recruited into the study were identified with the help of their hospital files or hospital discharge summaries they came with to the clinic while the controls were identified from their growth charts they come with to the child welfare clinic. After an informed consent was obtained, the mothers were interviewed using a standard questionnaire (Appendix II). Their obstetric and /or neonatal files were also reviewed for any antenatal, perinatal or neonatal events.

The growth patterns of the cases and controls were assessed using anthropometric measurements (weight, length and head circumference).

- a) Weight- measurement was obtained using a Seca balance no 62076 made in West Germany which measures to the nearest 10 grams. Accuracy of the machine was ascertained by the

investigator before recruitment starts. All infants were weighed nude.

- b) Length- A crown to heel measurement was taken with the infant supine and both legs extended in a measuring device containing a built in centimeters rule.
- c) Head circumference: was taken using an ordinary tape measure, at the largest occipitofrontal measurement obtained from two trials and an average of the two was recorded.

Their growth curves for weight and length were compared with the standard growth charts of the NCHS (31), while for the head circumferences were compared with the international and interracial composite graph by Nellhaus G.(39).

Their neurological status was assessed using a standard neurological test, by Dubowitz and Griffiths (Appendix III and IV) (36,37).

STATISTICAL ANALYSIS

Results were analysed statistically using X^2 test, student's t-test, Fisher's exact test and relative risk analysis where applicable. Statistical significance was set at $P < 0.05$ with 95% confidence limit.

RESULTS

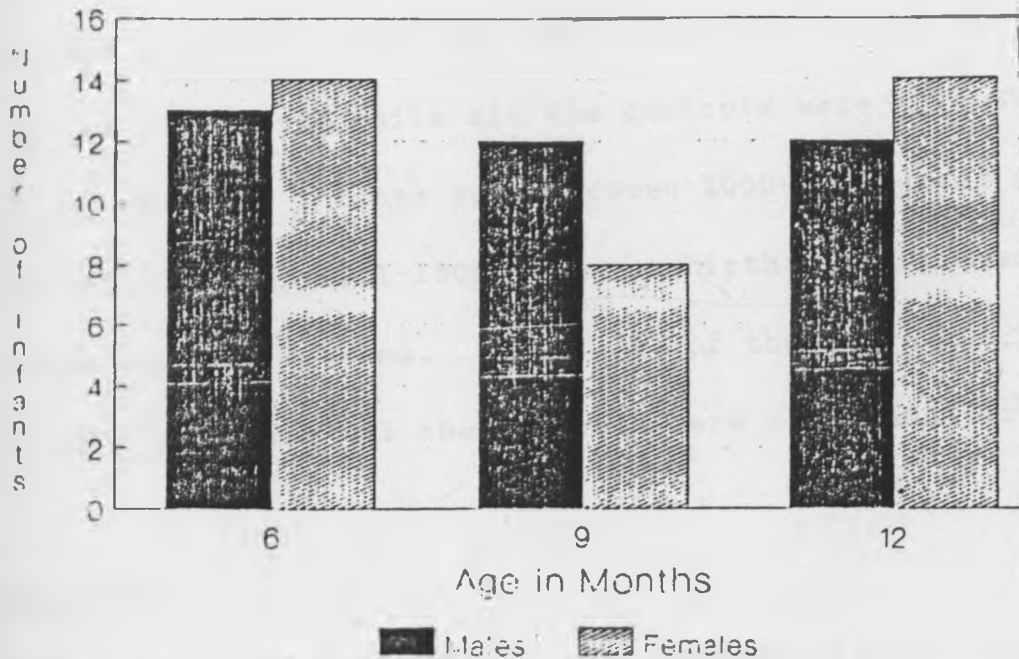
The study was carried out in the months of February to December 1989 inclusive. Seventy three of VLBW infants were recruited during the study period. These were compared with the same number of normal, term infants which were matched for age and sex. Table I and figure I show the age and sex distribution of the VLBW infants and the controls.

Table I

AGE AND SEX DISTRIBUTION OF THE VLBW INFANTS VERSUS THE CONTROLS.

AGE	M A L E S		F E M A L E S		TOTAL
	VLBW	CONTROLS	VLBW	CONTROLS	
6MO	13	13	14	14	54
9MO	12	12	8	8	40
12MO	12	12	14	14	52
TOTAL	37	37	36	36	146

FIGURE I AGE AND SEX DISTRIBUTION OF THE VLBW INFANTS.



MODE OF DELIVERY AND BIRTH WEIGHT DISTRIBUTION:

Among the VLBW infants, 50 (68.5%) were born SVD, 12 (16.4%) by C/S and 11 (15.1%) breech while all the controls were born SVD. 21 (28.7%) of the VLBW infants were between 1000-1250 grams and 52 (71.7%) were between 1251-1500 grams at birth. Their mean birth weight was 1356.2 grams. 64 (87.7%) of them were AGA, and 9 (12.3%) were SGA while all the controls were term, AGA infants.

GESTATIONAL AGE

Gestational age of the VLBW infants was assessed within the first 24 hours of birth, using the Dubowitz scoring system and it ranged from 28-36 weeks with a mean G.A score of 30.57 weeks. Sixty two (84.90%) of them were born at a gestational age of ≤ 32 weeks.

Table II.GESTATIONAL AGE DISTRIBUTION OF VLBW INFANTS

G.A (weeks)	NO	%
28-30	35	47.9
31-32	27	37.0
33-35	8	11.0
36-37	3	4.1
TOTAL	73	100

APGAR SCORE:

Sixty three of the VLBW infants had their 5 minutes apgar score recorded. Among these, 14 (22.2%) had perinatal asphyxia. This could not be compared with the controls because their 5 minutes apgar score could not be obtained. However, they all cried immediately after delivery, did not require resuscitation and / or admission to nursery and were all discharged from the maternity unit within 24 hours.

Table III.

5 MINUTES APGAR SCORE DISTRIBUTION OF THE VLBW
INFANTS

Score	Number	%
< = 6	14	22.2
> = 7	49	77.8
TOTAL	63	100

Among the VLBW infants 69 (94.5%) had one or more neonatal morbidities while only 4 (5.5%) of them had no problems at all. Respiratory distress followed by jaundice were the commonest problems encountered in this study group. Clinically suspected neonatal sepsis, patent ductus arteriosus, anaemia which required blood transfusion and repeated apnoeic attacks were also common problems. None of the controls had neonatal morbidities.

TABLE IV

DISTRIBUTION OF NEONATAL MORBIDITIES OF THE VLBW INFANTS

MORBIDITY	NO	%
RESPIRATORY DISTRESS	55	75.3
JAUNDICE	42	57.5
SUSPECTED NEONATAL SEPSIS	17	23.3
PATENT DUCTUS ARTERIOSUS	9	12.3
ANAEMIA	6	8.2
REPEATED APNOEIC ATTACKS	5	6.8
NONE	4	5.5

The prevalence of neurological sequelae among the VLBW infants was 8.2% which disagrees with the hypothesis. Gross motor (66.6%) followed by fine motor (22.2%) were the commonest neuromotor delays observed in this study group. Others were psychosocial (5.6%) and convulsive disorders (5.6%). All the neuromotor delays observed in this study group are of the minor types (Table V). None of the control groups had any neurological sequelae.

TABLE V

SUMMARY OF NEUROLOGICAL SEQUELAE OBSERVED

INFANT	GROSS MOTOR	FINE MOTOR	PSYCH. SOCIAL	SEISURES.
K.C	NO ATTEMPT TO ROLL BUT LIFT HIS HEAD.	DOES NOT TRANSFER OBJECT FROM HAND TO HAND.	NO SOCIAL RESPONSE TO A STRANGER.	-
6MO	DOES NOT SIT WITH TRUNK SUPPORT, TENDS TO FALL BACKWARDS. PARTIAL WT BEARING.	DOES NOT BRING OBJECT FROM HAND TO MOUTH FOR ORAL EXAM.		
9MO	DOES NOT CRAWL.	PARTIAL PINCER GRASP.	-	-
2.A.N				
3.J.W 9MO	CANNOT CRAWL CANNOT SIT WITHOUT SUPPORT PARTIAL WT BEARING	PARTIAL PINCER GRASP		OCCASIONAL
4.S.G 9MO	DOES NOT CRAWL.	-	-	-
5.L.A 9MO	DOES NOT CRAWL. PARTIAL WT BEARING (INTERMITTENT STANDUNG).	PARTIAL PINCER GRASP	-	-
12MO	NOT ABLE TO STAND			
6.R.A	NOT ABLE TO CRAWL	-	-	-

TABLE VI: SUMMARY OF THE VLBW INFANTS WITH NEUROLOGICAL SEQUELAE

	AGE (MO)	IP.NO.	SEX	MAT. AGE	EDUC. STATUS	MARITAL STATUS	ANC	OBST. RISK	MODE OF DELIVERY	B.W.	G.A.	APGAR SCORE	NEONATAL PROBLEMS
1.	6/mo	928224	M	20	Form IV	Single	NO	-PROM	SVD	1060	27	8	- R.D.
2.	9/mo	916275	M	24	Form IV	Married	YES	-PROM Abort. TWIN	SVD	1450 (SGA)	34	10	-
3.	9/mo	903681	F	21	Form IV	Single	NO	-PROM -APH	SVD	1400	30	5	-
4.	9/mo	898549	F	19	Form IV	Single	YES	-PROM	BRECH	1480	30	-	- R.D. Jaundice
5.	9/mo	892633	M	25	-	Single	NO	-	SVD	1200	28	10	- R.D.
6.	12/mo	897674	F	20	St.6	Married	YES	-	SVD	1200	31	8	- R.D. Jaundice -PDA

Table VI Shows that all the VLBW infants with neurological sequelae had one or more preconceptual, prenatal, perinatal and/or neonatal risk factors which might have been directly or indirectly related to their neurological outcome.

TABLE VII RISK FACTORS VERSUS NEUROLOGICAL SEQUELAE

		VLBW WITH SEQUEL	NORMAL VLBW	F.E.T P.V.	R.R	95% C.I.
1. MODE OF DEL.	SVD	5	45	0.71	1.18	0.14<RR<8.51
	BREECH	1	10			
2. BWT	1000-1250	3	18	0.23	2.48	0.54<RR<11.3
	1251-1500	3	49			
3. NEONATAL MORBID.	YES	4	65	0.03	0.12	0.03<RR<045
	NO	2	2			
4. APGAR SCORE	< = 6	1	13	0.69	0.88	0.11<RR<7.21
	> = 7	4	45			
5. MARITAL STATUS	SINGLE	4	13	0.23	6.6	1.32<RR<32.9
	MARRIED	2	54			
6. ANTENATAL CARE	NO	3	18	0.23	2.48	0.54<RR<11.3
	YES	3	49			
7. GEST. AGE (WEEKS)	28-30	4	31	0.03	2.17	0.42<RR<11.1
	> = 31	2	36			
8. PRIMARY CHILD'S CARE TAKER	MOTHER	4	44	0.67	1.84	0.2<RR<5.3
	OTHERS	2	23			

Table VII shows that neonatal morbidities and marital status are significantly related to the neurological sequelae observed in this study group.

NUTRITIONAL STATUS:

Nutritional assessment was done on both the VLBW infants and the normal term infants using anthropometric measurements of weight, length and head circumference. These were compared with the standard growth curves for each sex and age group.

Figures II to VI show that growth patterns of both the VLBW infants and the normal term infants lie within the normal range of the standard growth curves, except for the length of the VLBW male infants which is just below the normal range but parallels the standard curve. The VLBW infants were generally smaller for their ages than their counterparts of normal term infants.

FIGURE II: MEAN WEIGHTS FOR AGE OF THE VLBW INFANTS VERSUS CONTROLS.

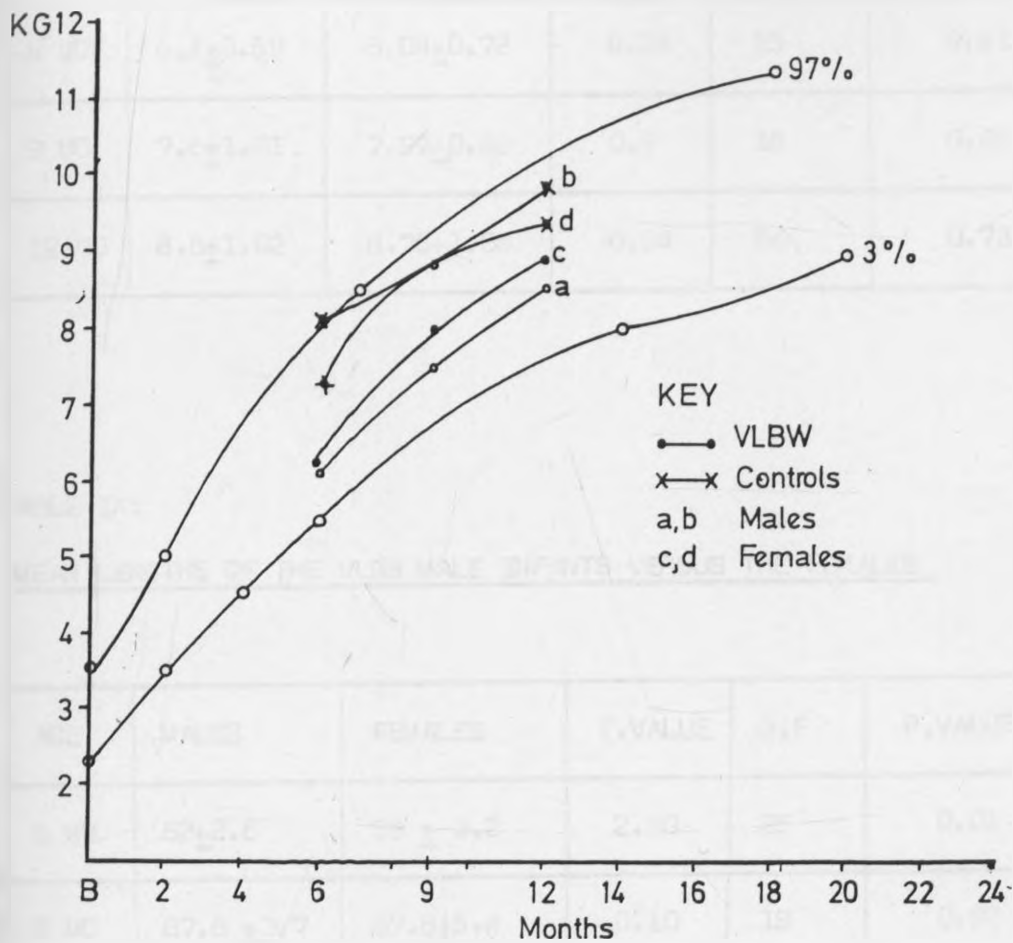


TABLE VIII:

MEAN WEIGHT (KG) OF THE VLBW MALE INFANTS VERSUS THE FEMALES

AGE	MALES	FEMALES	T.VALUE	D.F	P.VALUE
6 MO	6.1 \pm 0.59	6.04 \pm 0.72	0.24	25	0.81
9 MO	7.6 \pm 1.01	7.97 \pm 0.82	0.9	18	0.48
12 MO	8.6 \pm 1.02	8.76 \pm 1.36	0.34	24	0.73

TABLE IX:

MEAN LENGTHS OF THE VLBW MALE INFANTS VERSUS THE FEMALES

AGE	MALES	FEMALES	T.VALUE	D.F	P.VALUE
6 MO	62 \pm 2.6	59 \pm 3.2	2.70	25	0.01
9 MO	67.6 \pm 3.7	67.8 \pm 5.4	0.10	18	0.92
12 MO	70.7 \pm 3.6	70.9 \pm 3.8	0.14	24	0.88

Fig II, Tables VIII and IX show that among the VLBW infants, females are taller and heavier than the males at the ages of 9 and 12 months but there was no statistical significance.

TABLE X: MEAN WEIGHTS (KG) OF THE VLBW INFANTS VERSUS THE CONTROLS

MALES						FEMALES				
AGE	VLBW	NORMAL	t value	d.f.	p value	VLBW	NORMAL	t value	d.f.	p value
6 mo	6.05 ⁺ -0.59	8.15 ⁺ -1.15	5.86	24	0.0001	6.04 ⁺ -0.72	7.4 ⁺ -0.64	5.32	26	0.0001
9 mo	7.6 ⁺ -1.01	8.8 ⁺ -1.3	2.01	22	0.057	7.97 ⁺ -0.82	8.77 ⁺ -0.98	1.77	14	0.098
12 mo	8.61 ⁺ -1.02	9.75 ⁺ -0.9	2.90	22	0.008	8.76 ⁺ -1.36	9.3 ⁺ -0.99	1.20	26	0.24

Fig II and Table X Show that the VLBW male infants weigh less than the normal term, male infants and it was statistically significant at the ages of 6 and 12 months. The VLBW female infants also weigh less than their female counterparts but was only statistically significant at 6 months of age.

FIG. III MEAN LENGTHS FOR AGE OF THE VLBW MALE INFANTS VERSUS THE CONTROLS

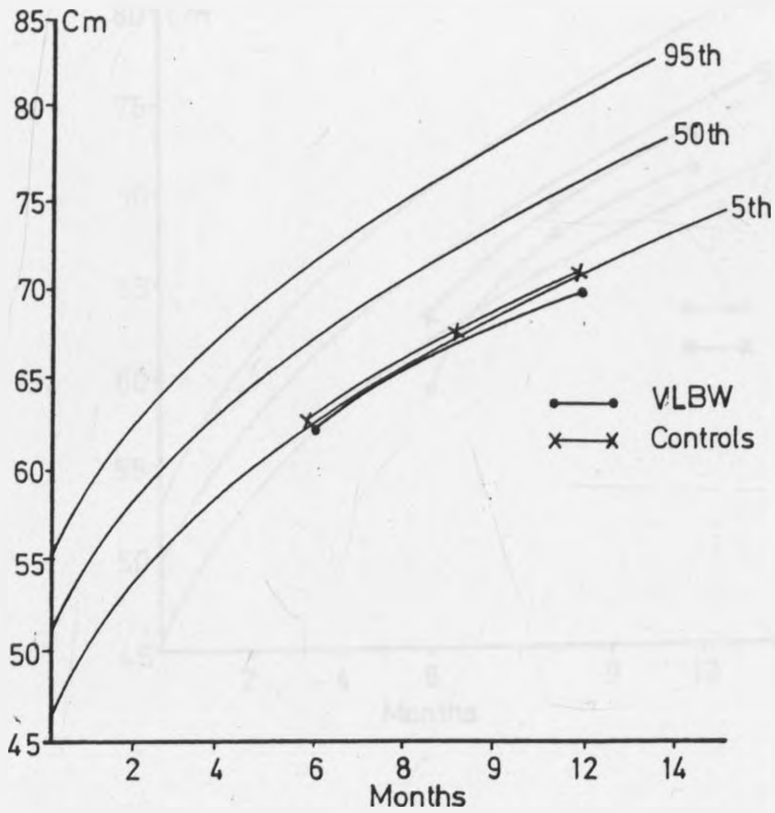


FIG. IV MEAN LENGTHS FOR AGE OF THE VLBW FEMALE INFANTS VERSUS THE CONTROLS

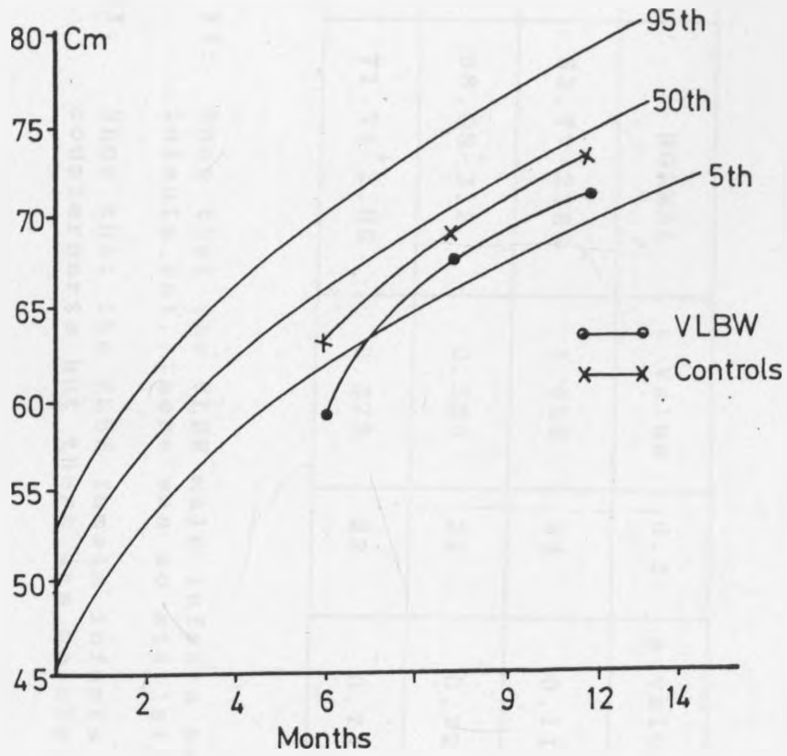


TABLE XI: MEAN LENGTHS (CM) OF THE VLBW INFANTS VERSUS THE CONTROLS

MALES						FEMALES				
AGE	VLBW	NORMAL	t Value	d.f.	p value	VLBW	NORMAL	t-value	d.f	p Value
6 mo	62 ⁺ -2.58	63.77 ⁺ -2.86	1.656	24	0.11	59.18 ⁺ -3.21	62.64 ⁺ -1.82	3.51	26	0.001
9 mo	67.58 ⁺ -3.65	68.08 ⁺ -3.2	0.356	22	0.72	67.75 ⁺ -5.4	69.31 ⁺ -3.65	0.67	14	0.15
12 mo	70.66 ⁺ -3.58	71.14 ⁺ -2.66	0.373	22	0.71	70.93 ⁺ -3.38	73.03 ⁺ -2.83	1.78	26	0.086

Fig. III and Table XI: Show that the VLBW male infants mean lengths are smaller than the normal term, male infants but, there was no statistical significance.

Fig. IV and Table XI: Show that the VLBW female infants mean lengths are also smaller than their female counterparts but there was no statistical significance, except at 6 months of age.

FIG. V MEAN HEAD CIRCUMFERENCES OF THE VLBW MALE INFANTS VERSUS THE CONTROLS

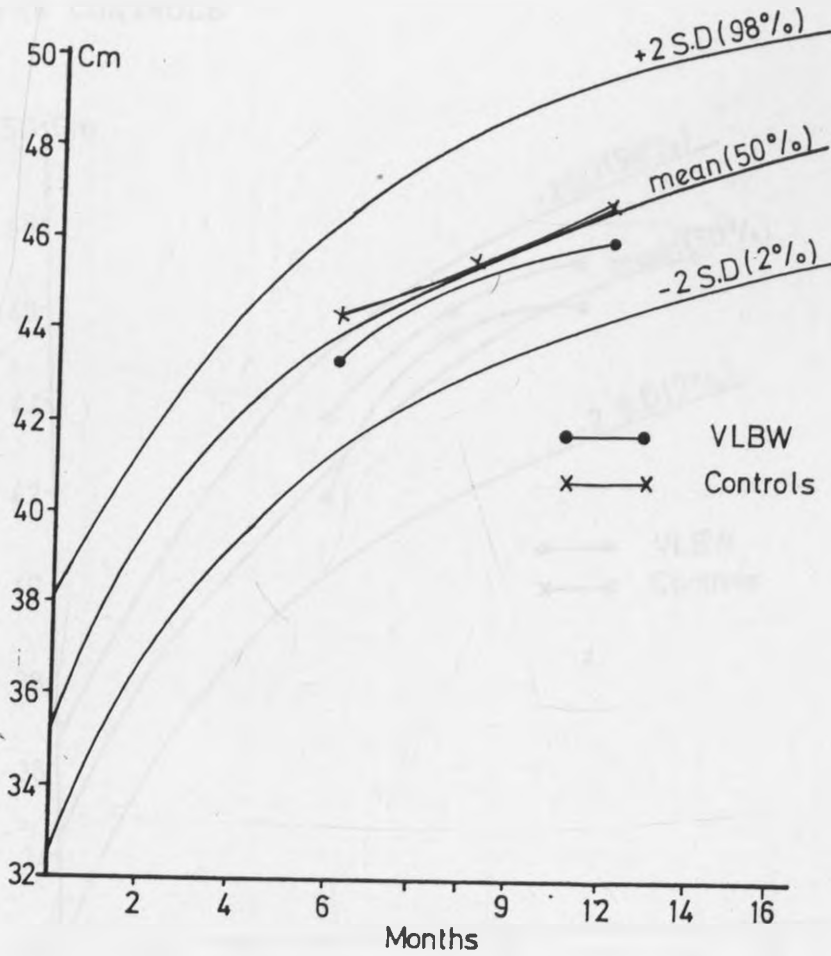


FIG. VI MEAN HEAD CIRCUMFERENCES OF THE VLBW FEMALE INFANTS VERSUS THE CONTROLS

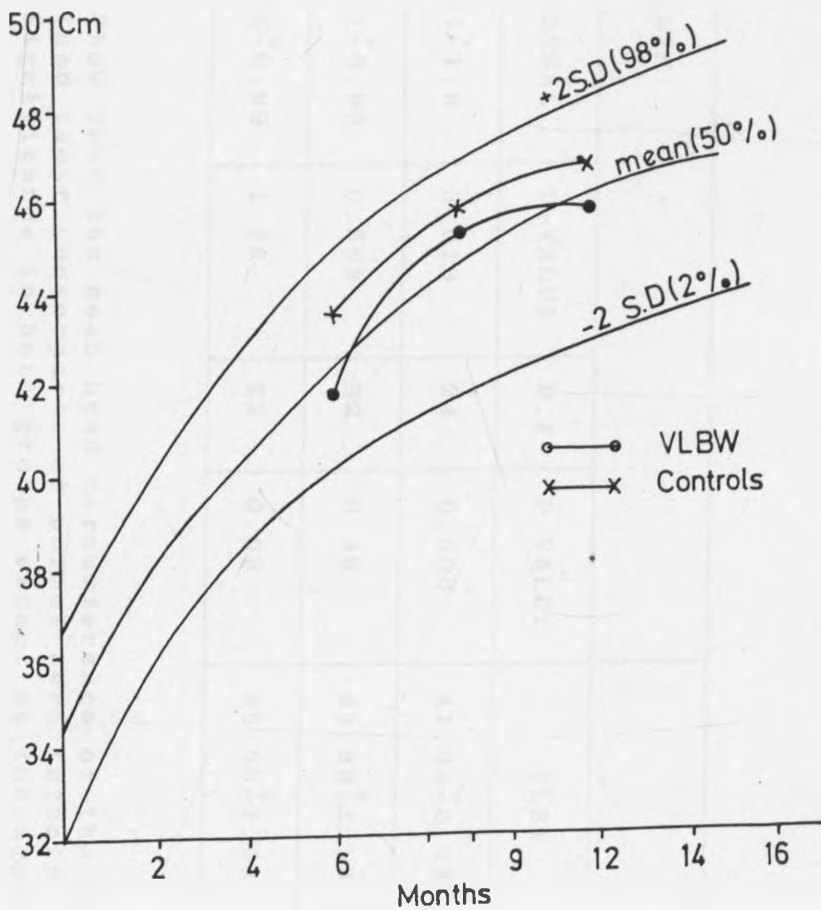


TABLE XII: MEAN HEAD CIRCUMFERENCE(CM) OF THE VLBW INFANTS VERSUS THE CONTROLS

MALES						FEMALES				
AGE	VLBW	NORMAL	T-VALUE	D.F	P.VALUE	VLBW	NORMAL	T-VALUE	D.F	P VALUE
6 mo	43.16 ⁺ -1.17	44.54 ⁺ -1.6	3.019	24	0.006	41.94 ⁺ -0.78	43.43 ⁺ -1.02	4.34	26	0.0001
9 mo	45.42 ⁺ -0.9	45.71 ⁺ -0.98	0.699	22	0.49	45.58 ⁺ -1.15	45.6 ⁺ -1.22	0.186	14	0.85
12mo	46.1 ⁺ -1.36	46.94 ⁺ -0.99	1.76	22	0.09	45.96 ⁺ -1.38	46.9 ⁺ -1.32	1.53	26	0.139

Fig V, VI and Table XII Show that the mean head circumference of the VLBW male and female infants are smaller than their counterparts of normal term infants. However, there was no statistical Significance in both groups except at the age of 6 months.

MATERNAL AGE:

Maternal age for the VLBW infants ranged from 17 - 44 years with a mean of 24.85 years while for the term infants ranged from 19 - 39 years with a mean value of 26.68 years.

Chi square trend shows that mothers of VLBW infants tend to be younger than the mothers of the term babies.

TABLE XIII:

DISTRIBUTIONN OF MATERNAL AGE OF THE VLBW INFANTS
VERSUS THE CONTROLS.

AGE (YRS)	VLBW	NORMAL	TOTAL
<= 20	12	2	14
21-26	42	38	80
27-32	17	26	43
>= 33	2	7	9
TOTAL	73	73	146

X^2 TREND = 9.55 P = 0.001

MARITAL STATUS:

Table XIV shows that there were significantly more single mothers among the VLBW infants than the mothers of the normal term infants.

TABLE XIV: MATERNAL DISTRIBUTION BY MARITAL STATUS

	VLBW	NORMAL	TOTAL
SINGLE	17	5	22
MARRIED	56	68	124
TOTAL	73	73	146

$$\chi^2 = 6.48 \quad P = 0.01$$

$$R.R = 1.71$$

$$95\% \text{ C.I} = 1.27 < R.R < 2.31$$

MATERNAL EDUCATION:

Table XV shows a trend that mothers of the VLBW infants tend to have less educational status at a secondary and college / university level than those of the normal infants.

TABLE XV: DISTRIBUTION BY MATERNAL EDUCATION

LEVEL OF EDUCATION	VLBW	NORMAL	TOTAL
NIL	4	2	6
PRIMARY	24	11	35
SECONDARY	44	51	95
COLLEGE/ UNIVERSITY	1	9	10
TOTAL	73	73	146

$$\chi^2 \text{ Trend} = 10.4$$

$$P = 0.001$$

INCOME:

Table XVI shows that more parents of the VLBW infants have no regular income than of the normal term infants.

TABLE XVI: DISTRIBUTION BY INCOME STATUS

REGULAR INCOME	VLBW	NORMAL	TOTAL
NO	9	1	10
YES	64	72	136
TOTAL	73	73	146

Fisher's exact test, $p = 0.008$

ANTENATAL CARE ATTENDANCE:

Table XVII shows that there is a significantly poor antenatal care attendance among mothers of the VLBW infants. All the mothers of the normal term infants had regular antenatal care attendance.

TABLE XVII: DISTRIBUTION BY ANTENATAL CARE ATTENDANCE

ATTENDED	VLBW	NORMAL	TOTAL
NO	21	-	21
YES	52	73	125
TOTAL	73	73	146

$$X^2 = 22.4 \quad D.F = 1 \quad P = 0.0001$$

$$R.R = 2.40$$

$$95\%/C.I = 1.95 < R.R < 2.96$$

OBSTETRIC RISK FACTORS:

Fifty five (73.34%) mothers of the VLBW infants had one or more obstetric risk factors. PROM followed by history of abortion were the commonest risk factors encountered. Antepartum haemorrhage (APH), preeclampsic toxemia (PET) and twin pregnancy were also major risk factors in this study group. None of the mothers of the normal infants had any risk factor.

TABLE XVIII: PATTERN OF OBSTETRIC RISK FACTORS

RISK FACTORS	NO	%
PROM	29	39.73
HISTORY OF ABORTION	24	32.88
APH DURING 2nd & 3rd TRIMESTER	15	20.60
TWIN PREGNANCY	13	17.80
PET	13	17.80
OTHERS	9	12.30

PARITY

Table XIX shows that there was no statistical difference between parity of the mothers of the VLBW infants and those of the normal term infants.

TABLE XIX: DISTRIBUTION BY PARITY OF THE MOTHERS OF THE VLBW INFANTS VERSUS THE CONTROLS

PARITY	VLBW	NORMAL	TOTAL
0 - 4	64	66	130
> 4	9	7	16
TOTAL	73	73	146

$$\chi^2 = 10.3 \quad P = 0.1$$

$$R.R = 0.88$$

$$95\% \text{ C.I} = 0.55 < R.R < 1.39$$

DISCUSSION

Fourteen (22%) of the VLBW infants with apgar score recorded had perinatal asphyxia that is an apgar score of 6 or less at 5 minutes. A 10 or 20 minutes apgar score would have been more useful as the perinatal asphyxia at 10 or 20 minutes is highly associated with neurological sequelae than the 5 minutes score. (4) However, these were not recorded in our study group as it is not a common practise in our maternity unit. Lekha (1989) from KNH reported that perinatal asphyxia at 5 minutes among those with birth weight of less or equal to 2000 grams was 37.5% (8). Kasirye (1984), from KNH has reported that perinatal asphyxia has been associated with 57% and 36% of neonatal mortality of infants with birth weight of less than 1000 grams and 1000 - 1500 grams - respectively (9). Therefore perinatal asphyxia is a major obstetric risk factor associated with a high perinatal and neonatal mortality rate in our set up.

94.5% of the VLBW infants had one or more neonatal morbidities. Respiratory distress (75%) and jaundice (57.5%) which required phototherapy were the main problems encountered. Worthington et al has also reported that respiratory distress (62%) was the commonest problem seen in the VLBW infants (20). The incidence of PV/IVH among the VLBW infants, its evolution and its association with specific perinatal and /or neonatal events is not known in our set up.

However, other studies indicate that PV/IVH is the most important risk factor related to neurological sequelae among the VLBW infants (4,12,13,14).

The prevalence of neurological sequelae among the VLBW infants was 8.2% and none of them had a major neurological sequelae. This is in contrast to reports from developed countries with neonatal intensive care centres which ranges from 10-25% only with major neurological sequelae. (4,12-17). However, it might be difficult to compare ours with other reports due to the following reasons:-

- a) This study was cross-sectional while others was a long term follow up. It is known that some of the infants with minor sequelae can progress into major ones or even those without sequelae during infancy can manifest at later age of childhood. Minimum of three years and five to ten years is required to detect major and minor neuropsychiatric abnormalities respectively. (7,17).
- b) Psychometric assessment was part of the long term follow up in other studies, but it was not done in this study because it is not a reliable predictor of neurological sequelae in infancy. (7,11)
- c) The survival rate of the VLBW infants in our set up ranges between 25-50% (9,22). This is very low as compared to developed countries which ranges between 80-95% (18). Even the extremely low birth weight infants who have the

highest prevalence of neurological sequelae have a survival rate of 60-70% in some neonatal intensive care centres (11,13), while in our setup is between 0-5% (9,22).

Therefore, the high survival rate in developed countries might be the reason for the high proportion of VLBW infant survivors with neurological sequelae as opposed to our setup in which most of those with neurological complications might have died during neonatal period.

- d) The author has observed that between December 1987 to May 1989 inclusive, 585 VLBW infants were admitted to the newborn nursery unit. (35) Out of these (313 (52.6%) survived and were discharged home. All of these were expected for neurological assessment during the study period. However only 73 (23%) turned up at their expected date for neurological assessment. It is not known how many of the defaulters survived after discharge home. Therefore, the low prevalence rate of neurological sequelae as opposed to other reports could have also been influenced by a selection bias.

All the VLBW infants with minor neurological sequelae had one or more preconceptual, prenatal, perinatal and neonatal risk factors which might have directly or indirectly been related to the neurological outcome. An attempt to relate these risk factors to neurological sequelae is made using Fisher's Exact test and relative risk analysis. The only significant factors related were neonatal morbidities and marital status .

This could be due to the high incidence of respiratory distress and Jaundice encountered in this group which agrees with Stewart et al that hypoxia due mainly to respiratory distress, and jaundice were the most common risk factors related to the long term neurological sequelae (12). marital status was significantly related to the neurological outcome which agrees with others that marital status especially with the young unmarried mothers is indirectly associated with high incidence of LBW and subsequently with their neurological outcome (16,38).

However, the number of VLBW infants with neuromotor delay in this study group are too small for a meaningful conclusions to be drawn about the cause of neuromotor delays. In addition it is known that in most instances, the neurological sequelae is an additive process of multiple factors which when statistical analysis is attempted for each risk factor, significant result might not come out especially when the numbers are small like in this study group.

Nutritional assessment using anthropometric measurements of weight, length and head circumference was conducted on both the VLBW and the normal term infants, as malnutrition is one of the confounding factors for neurological assessment. The growth patterns of both groups lie within the normal range of the standard growth curves except for the length of the VLBW male infants which lie below but parallels the standard curve. The VLBW infants in this study group were generally smaller for their age than the normal term infants. This agrees with Binkin et al

(1988) who has studied a group of children up to the age of five years with different birth weights and observed that infants with LBW are likely to remain smaller for their age than the normal or large birth weights (32). This has also been observed by Babson (33). Among the VLBW infants females were heavier and taller than the males at the ages of 9 and 12 months which is in contrary to findings by Drillien and Fitzhardinge (10,34). At the moment there is no explanation for these findings. However, these were not statistically significant.

Mothers of the VLBW infants were significantly younger and less educated. The VLBW infants also tend to come from the lower income class than those of the normal term infants. 28% of the mothers of the VLBW infants did not have antenatal care, while all of those of the normal term infants had one or more antenatal clinic visits. Mati et al (1980) in the Nairobi birth survey has observed that 40% of mothers of the LBW infants did not attend antenatal clinic (29). Lekha (1989) from K.N.H reported that 40% of mothers of babies with birth weight of less or equal to 2000 grams did not have antenatal care (8). These show that inadequate or lack of antenatal care is still a major problem in our setup which is one of the main contributing factors for the high incidence of LBW and prematurity. There were also more single mothers in the VLBW infants than their counterparts ($p < 0.01$). All these findings agree with others that young maternal age, single mothers, low socio-economic and educational status and poor antenatal care increase the incidence of LBW and

prematurity and subsequently the neonatal morbidity by two to four fold (1,8,9,16,22).

Fifty five (73.34%) of the mothers of the VLBW infants had one or more obstetric risk factors while none of the mothers of the control group had any risk factor. PROM (37.7%) followed by a previous history of abortion was the commonest risk factor observed. This agrees with others that PROM is the commonest risk factor associated with prematurity. It has been observed in the United States that among the VLBW infants 75%, have history of PROM (7). Mati et al in the Nairobi birth survey reported that the incidence of PROM among all pregnant mothers was found to be 32.5% (22). This figure might even have been higher had the observation been only for the VLBW infants. Antepartum haemorrhage, preeclampsia toxemia, and twin pregnancy were also major risk factors encountered in this group. Others were previous history of still birth, neonatal death, or prematurity and febrile illnesses.

CONCLUSIONS

1. The prevalence of neurological sequelae among the VLBW infants was 8.2% and none of them had major sequelae.
2. All the infants with neurological sequelae had one or more preconceptual, prenatal, perinatal and neonatal risk factors which might have been related to their neurological outcome.
3. The prevalence of obstetric risk factors among the VLBW infants was 73.34%. PROM (39.7%) followed by history of abortion (32.9%) were the commonest factors encountered.
4. The VLBW infants were smaller for their ages than the normal term infants, in terms of their weights, lengths and head circumferences. However, between the male groups, the weights were statistically significant at the age of 6 and 12 months while their head circumference only at the age of 6 months. There was no statistical significance between their mean lengths. Between the female groups, there were statistical differences in their weights, lengths and head circumferences only at the age of 6 months.

RECOMMENDATIONS

1. A long term follow up of the VLBW infants is advocated to assess their neurological and psychometric outcome.
2. A study on the incidence of PV/IVH is desirable in our set up.
3. There is a need for all pregnant mothers to have adequate antenatal care.

ACKNOWLEDGEMENTS

I would like to express my sincere appreciation to:

1. My Supervisors:

Dr, A.O. WASUNNA and Dr. R.N. MUSOKE, for their constant and tireless help and guidance throughout the various stages of the preparation of this work.

2. Prof. J. BRADY, for her valuable criticisms.

3. The German Academic Exchange Service (DAAD), for awarding me the Scholarship through the University of Nairobi.

4. I would also like to thank the Dean, Faculty of medicine, Prof. Pamba and the then Chairman of the Department of paediatrics Prof. Meme, for accepting me in the Training programme.

5. Finally my sincere appreciation goes to all those Mothers who allowed me their Children and themselves to be included in the study.

REFERENCES

1. W.H.O Bulletin 65: no 5, 575-768, 1987.
2. Battaglia, F.C and Lubchenco, L.O.
A practical classification of Newborn infants by birth weight and gestational age.
J. Pediatr. 71; no 2, 159-163, 1967.
3. Lubchenco L.O, Searls, D,T.
Neonatal morbidity rate : Relationship to birth weight and gestational age.
The J. Pediatr. 81; no 4, 814-822, 1972.
4. Yu., V.Y.H, Nong, P.Y, Bajuk, B, et al.
Outcome of extremely low birth weight infants.
Br. J. Obstet and Gynaecol. 93; 162-170, 1986.
5. Commey, J.O.O., and Fitzharding, P.M.
Handicap in the preterm, SGA infant.
J. Pediatr.94; no 5, 779-789, 1979.
6. Philip, A.G.S, Little, G.A, Polivy, D.R., et al.
Neonatal morbidity risk for the eighties:
The importance of birth weight/gestational age groups.
Pediatr. 68; no 1, 122-130, 1981.

7. Klaus, S.A and Fanaroff, A.A.
Care of the high risk neonate,
3rd edition, W.B. Saunders, Philadelphia, 1986.
8. Lekha Dissanayake, M.MED Dissertation (1989).
Obstetric variables determining survival of the ≤ 2000
grams babies during the first seven days of life at Kenyatta
National hospital.
9. Kasirye, E., M.MED dissertation (1984).
Neonatal morbidity and mortality at Kenyatta National
hospital.
10. Cecil M. Drillien.
Growth and development in a group of children of VLBW
infants.
Arch. Dis. Child, 10-18, 1957.
11. The fate of the baby under 1501 grams at birth. Lancet,
461-462, 1980.
12. Ann. Stewart, and Reynolds, E.O.R.
Improved prognosis for infants of VLBW.
Pediatr. 54: 724-734, 1974.

13. Hoskins, E.M., Elliot, E., Shennan, A.T., et al.
Outcome of the VLBW born at perinatal centre.
Am. J. Obstet., Gynaecol. 145; 135-139, 1983.
14. Ann Stewart, Turcan, D.M., Rawlings, G., and Reynolds, E.O.R.
Prognosis for infants weighing 1000 grams or less at birth.
Arch. Dis. Child 52; 97-103, 1977.
15. Paul, R.H., Koh, K.S. and Monfared, D.M. et al.
Obstetric factors influencing the outcome of infants
weighing 1001 to 1500 grams.
Am. J. obstet, Gynaecol. 133: 503-507, 1979.
16. Hilda Knobloch, Malone, A., Ellison, P.H., et al.
Considerations in evaluating changes in outcome for infants
weighing less than 1501 grams.
Pediatr. 69: No 3; 285-295, 1982.
17. Krishna-Moorthy, K.S., Shannon, D.C., Delong, G.R. et al.
Neurological sequelae in the survivors of neonatal IVH.
Pediatr. 64; 233-237, 1979.
18. Horbar, J.D., McAuliffe, T.L., Adler, S.M. et al.
Variability in 28 days outcome for VLBW infants. An
analysis of eleven neonatal intermine care units.
Pediatr. 82; no 4, 554-559, 1988.

19. Maureen Hack, Merkatz, I.R, Jones, P.K, Fanaroff, A.A.
Changing trends of neonatal and post deaths in VLBW infants.
Am. J. obstet. Gynaecol., 137: 797-800, 1980.
20. Dennis Worthinton, Davis, L.E, Grausz, J.P, and Sobocinski, K.
Factors influencing survival and morbidity with VLBW
delivery.
Obstet and Gynaecol 62: 550-554, 1983.
21. Infants of VLBW: A 15 years analysis.
Lancet, 1332-1335, 1979.
22. Mati, J.K.G, Aggarwal, V.P., Lucas.S., et al.
Early perinatal mortality rate:
A Nairobi birth survey IV.
J. obstet, Gynaecol. East, Cent. Africa. No 2: 129-133,
1983.
23. Prechtl, H.F.R,
Neurological sequelae of prenatal and perinatal
complication.
Br. Med. J. 4: 763-767, 1967.
24. Tarby, T.J. and Volpe, J.J.
Intraventricular haemorrhage in the premature infant.
Pediatr. Clinics of N. America 29: no 5, 1077-1100, 1982.

25. Garecia et al.
Behavioural responsiveness in pre term infants with IVH.
Pediatr. 81; no 3, 1988.
26. Intraventricular haemorrhage in the newborn Beoge puppy: A limited model of I.V.H in the premature infant.
Pediatr. neurosci. 13: 78-83, 1987.
27. Amato et al.
Ultra sonographic and Biochemical correlate of periventricular leucomalasia in VLB infants.
Pediatr. neuro sci. 13: 84-89, 1987.
28. Cole, U.A, Durbin, G.M, Olaffson, A. et al.
Pathogenesis of I.V.H in newborn infants.
Arch. Dis. Child 49: 722-727, 1974.
29. Papile, L.A, Brustein, J., Brustein, R. and Koffler, H.
Incidence and evaluation of subependynal and I.V.H: A study of infants of birth weight less than 1500 grams.
J. Pediatr. 92: 529-533, 1978.
30. Hilda Knobloch, Stevens, F., Malone, A., Ellison, P. et al.
Validity of parental reporting of infant development.
Pediatr. 63: no 6, 1979.

31. Nelson, W.E. Vaughan U.C. Behrman R.E (editors), Nelson text book of paediatrics, 13th edition, W.B. Saunders. Philadelphia, 1987.
32. Einkin, N.J, Yip, R., Fleshood, L and Trowbridge, F.L. Birth weight and childhood growth. *Pediatr.* 82: no 6, 1988.
33. Babson, S.G. Growth of the LBW infants. *J. Pediatr.* 77: no 1, 11-18, 1970.
34. Fitzhardinge, P.M. Early growth and development in VLBW infants, following treatment at intensive care nursery. *Pediatr.* 56: 162-171, 1975.
35. KNH, Newborn nursery unit admission and discharges register book, December, 1987 to May 1989 inclusive.
36. Clinics in developmental Medicine No. 79, the Neurological assessment of the Preterm and full term infants, by L. Dubowitz and V. Dubowitz, 1981, Spastics international medicine publications.

37. Record book for use with Griffiths mental developmental scales for testing babies from birth to eight years of age.
38. The LBW infant, perspectives of Paediatric nutrition, 1981.
39. International and interracial composite graph by Nelihaus .G Paediatrics, 1968.
40. Njuki, H,M.MED Dissertation (1983).
LBW infants at Machakos Provincial hospital:
Incidence, etiology and mortality.
41. Harvey, D., Prince, J, Bunton, J., Parkinson, C., et al.
Abilities of children who were SGA babies:
Pediatr. 69: no 3, 296-300, 1982.
42. Improved outcome in VLBW infants, correspondence.
Am. J. obstet, Gynaecol, 136: no 84, 1080-1081, 1980.
43. Intensive care and the VLBW infants.
Lancet pp 362-363, 1979.
44. Connel, J., Vries, L.D., Oozeer, R., et al.
Predictive value of early continuous electroencephalogram monitoring in ventilated preterm infants with IVH.
Pediatr. 82: no 3, 337-343, 1988.

45. Epstein, M.F., Leviton, A., Kuban, K.C.K, et al.
Bilirubin, IVH and phenobarbital in VLBW babies.
Pediatr. 82: no 3, 350-354, 1988.
46. Dimesony et al.
Intracranial haemorrhage in SGA neonates:
Comparison with weight matched AGA infants.
Clinical Pediatr. 27:no 1, 1988.
47. Shennan, A.T., and Milligan, J.E.
The growth and development of infants weighing 1000-2000
grams at birth and delivered in a perinatal unit.
Am. J. obstet, Gynaecol. 136: 273-275, 1980.
48. Howard, J, Pamelee, A.H, Kopp, C.B and Littman, B.
A neurological comparison of preterm and full term infants
at term conceptual age.
The J. Pediatr. 88: no 6, 995-1001, 1976.
49. Littmann, B and Pamelee, A.H.
Medical correlates of infant development.
Pediatr. 61: no 3, 470-474, 1978.

50. Meme, J and Hillman, D.
LBW infants at Kenyatta National hospital.
E, Africa Med. J. 54: 27-30, 1977.
51. Is intensive care justified for infants weighing less than
800 grams at birth?
J.Pediatr. 99: 1981.
52. Frankenburg, W.K, Goldstein, A.D and Camp, B.W.
The Revised Denver developmental screening test:
Its accuracy as a screening test.
J. Pediatr. 79: no 6, 988-995, 1971.
53. Dubowitz, L.M.S, Dubowitz, U and Goldberg, C.
Clinical assessment of gestational age in the newborn
infant.
J. Pediatr. 77: no 1, 1-10, 1970.
54. Boules, W.A.
Fetal heart monitoring in premature infants weighing 1500
grams or less.
Am. J. obstet. Gynaecol. 137: 1980.

55. Kitchen, W, Ford, G.W., Doyle, L.W., et al.
Caesarian section or vaginal delivery at 24-28 weeks of gestation:
A comparison of survival and neonatal and two years morbidity.
Obstet. Gynaecol. 66: no. 2 149-156, 1985.
56. Hilda Knobloch, Pasamanick, B and Sherald, E.S.
Development screening inventory for infants.
Pediatr. 38: no 6, 1095-1104, 1966.
57. Mortality and morbidity in VLBW infants 500-1500 grams.
Pediatr. 69: no 5, 1982.
58. Tison, C.A.
Neurological evaluation of the maturity of newborn infants.
Arch. Dis. Child. 43: 89-93, 1968.
59. Avery, M.E, Tooley, W.H, Keller, J.B. et al.
Is chronic lung disease in VLBW infants preventable?
A survey of eight centres.
Pediatr. 79: no 1, 26-30, 1987.
60. Karlberg, P. and Ericson, A.
Perinatal mortality in Sweden:
Analysis with international aspects.
Acta Pediatr. scand supp 275, 28: 28-34, 1979.

APPENDIX I

DEFINITIONS

1. Very low birth weight (VLBW):- Infants who weigh ≤ 1500 grams at birth. (1)
2. Extreme low birth weight (ELBW):- Infants with birth weight of ≤ 1000 grams.
3. Low birth weight (LBW):- Infants with birth weight ≤ 2500 grams.
4. Preterm:- Infants born before 37 completed weeks of gestation from the first day of last menstrual period.
6. Appropriate for gestation age (AGA):- Birth weight between 10th and 90th percentile of Lubchenco's norms (2).
7. Small for gestational age (SGA):- Birth weight below 10th percentile of Lubchenco's norms.
8. Large for gestational age (LGA):- Birth weight above 90th percentile of Lubchenco's norms.
9. Respiratory distress (RD):- Expiratory grunting, nasal, flaring, cyanosis in room air, tachypnea ($RR > 50/\text{min}$), chest in drawing. (7)

10. Premature rupture of membrane (PROM):- When membrane ruptures \geq 24 hours before labour starts. (7).
11. Preeclampsia toxemia:- Pregnancy induced hypertension with BP \geq 140/90 mmhg + oedema + proteinuria.
12. Apnoea:- Cessation of breathing for more than 20 seconds or long enough to produce cyanosis or bradycardia. (7,31)
13. Perinatal Asphyxia:- Infants with Apgar score of \leq 6, five minutes after birth. (7,31)
14. INCOME:- Regular earning of the mother or father or both not less than Ksh 500 a month.
15. Neurological Sequelae:
 - i) Minor:- hypotonia, hypertonia, or generally mild distortion of gross and/or fine motor integration, delay in neuro motor development, IQ 70-85 or occasional attacks of seizures. (16)
 - ii) Major:-
 - cerebral palsy of any type.
 - severe visual and hearing impairment.
 - IQ $<$ 70.
 - hydrocephalus.

- Frequent attacks of seizures. (16)

16. HANDICAP:-

i) Minor:-

-disability that does not or is unlikely to prevent the child from going to a normal school, or to interfere with normal life in society. (14)

ii) Major:-

- Disability of body, intellect, or personality that is significantly severe to prevent or is likely to prevent the child from going to normal school or causes a serious interference with normal function in his/her society. (14)

17. IQ (Intelligent quotient):- Ratio of mental age to -chronological age multiplied by 100. (31)

18. Mental retardation:- If the IQ is more than two standard deviation below the mean for his/her age on a standard psychometric test for IQ. (31)

a) 70-85 = Borderline

b) 55-70 = mild

c) 35-55 = moderate

d) 20-35 = Severe

e) < 20 profound.

19. PV/IVH (periventricular/intraventricular haemorrhage)

Intracranial haemorrhage of subependymal matrix origin. a)

grade I - subependymal haemorrhage only

b) grade II - IVH without ventricular dilatation

c) grade III - IVH with ventricular dilatation

d) grade IV - IVH with parenchymal involvement and ventricular enlargement with or without periventricular cyst

(24).

Appendix II.

STUDY FORMAT NO..... DATE

Mother's Name IP.NO..... Age

Infant's Name..... IP.NO..... Age Sex

Address

1) MATERNAL VARIABLES

Social and Family history

i) marital status - single / married

ii) parity -

iii) level of maternal education -

illiterate/primary/secondary/college or university

iv) Avearge income per month

a) mother Ksh -

b) father Ksh -

2) Chronic maternal disease

- Diabetes mellitus Yes/No
- Hypertension Yes/No
- Renal problem Yes/No
- Epilepsy Yes/No
- Chronic cough Yes/No
- Chronic diarrhoea Yes/No
- Weight loss Yes/No
- Others (specify)

5) Resuscitative measures:

- Suction and /or bagging only
- I.V adrenaline
- Intubation
- Extra cardiac massage
- Adrenaline
- I.V Na HCO₃
- I.V destrose
- Others specify

6) Neonatal and / or postneonatal morbidities

- RD Yes/No
- Repeated apnoeic attacks Yes/No
- Infections Yes/No

if Yes, Specify

- Anaemia which required blood transfusion Yes/No
- Hypoglycaemia cerca Yes/No
- Seizure disorder Yes/No
- Birth injuries Yes/No

if Yes, specify

- Others, specify

7) Nutritional Assessment:-

Wt.....length HC MAC

Growth faltering Yes/No

8) Neurological and Milestone assessment:-

a) History of convulsion Yes/No

b) Milestones assessment

i) at 6 months Age:

- Social response to a strange Yes/No.

- In prone position, raises head and chest off couch Yes/No.

- Rolls from prone to supine position and vice versa Yes/No.

- Head support without lag Yes/No.

- Sits with trunk support Yes/No.

- Orientates entire body towards desired object Yes/No.

- Brings object to mouth for oral examination Yes/No.

- Has palmar grasp Yes/No.

- Laughs at pleasurable social contacts Yes/No.

- Supine position = can lift head up spontaneously Yes/No.

- On standing position = can bear full weight on the leg Yes/No.

- Can transfer object from hand to hand Yes/No.

ii) At 9 months Age:

- Can crawl Yes/No.

- Stands with hand held Yes/No.

- Sits without support Yes/No.
- Transfer object from hand to hand Yes/No.
- Waves bye-bye (imitate) Yes/No.
- Says ba-ba, ma-ma, da-da Yes/No.
- Has Radial - Palmar grasp Yes/No.
- Turn head consistently to locate source of sound
Yes/No.
- Begins to go for object with index finger.
Yes/NO.

iii) At 12 months of Age:

- Release object on demand Yes/No.
- Has pincer grasp Yes/No.
- Walks with one hand supported Yes/No.
- Plays simple games a toy and immitates games
Yes/No.
- Obeys simple requests "Give me shoes" etc
Yes/No.
- Says three clear words Yes/No.

c) Neurological Assessment

See attached format by L. Dubowitz , V. Dubowitz and Griffiths (Appendix III and IV).

APPENDIX III
6 MONTH NEUROLOGICAL

No	Description	1 - 2		3 - 7		Score
		A	B	C	D	
8	Eye Movements	No following	Follows horizontally only	Follows vertically only	Follows in arc	Sustained upwards gaze
9	Visual acuity	No focus	Strabismus	Questionable	Normal with convergence	
10	Eye appearance	Nerve palsy + pupil inequality	Strabismus	Questionable	Normal with convergence	
11	Grasp of brick or rattle	No sustained grasp	2 handed grasp	1 handed grasp	Object in each hand	Transfers object
12	Hearing with bell	No response	Listens-no head turning	Hearing-slow localisation	Immediate response	
13	Movements: Arm	Abnormal (specify)	Questionable, ie clumsy, over-pronates		Normal	
14	Leg		Jerky, very slow		Normal	
15	Resistance to passive movements	Elbow extension < 100°		Up to 160° tight	Up to 140° easily	
16		Hip abduction < 70°	70° + spasm	Up to 70° floppy	Up to 70° easily	
17		Popliteal angle < 100°		Up to 100° tight	Up to 140° easily	
18		Foot dorsiflexion < 90°		Up to 90° tight	Up to 90° easily	
19	Reflexes: Biceps	Def. absent exag. unequal		7ble/mildly unequal	Normal	
20	Adductor	Def. absent exag. unequal		7ble/mildly unequal	Normal	
21	Knee	Def. absent exag. unequal		7ble/mildly unequal	Normal	
22	Clonus	Present			Normal	
23	Supine (pass by report)	Immobile no head lifting	Lifts head - no attempt to roll	Lifts head - attempts to roll	Rolls one way	Rolls both ways
24			Cannot get toes near head	Does not play with toes	Plays with toes	Toes to mouth
25	Pull to sitting a Head	Head lag or wobble			Normal	
26	b Body	No traction response	40 Falls to stand or poor tract resp	70	Normal	
27	Sitting	Inability to sit supported	Jackknives backwards on strng legs	Tendency to fall backwards	Sits well supported	Sits with support
28	Protective Reflexes	Absent		Forwards present	Forwards + sideways	All present
29	Weight bearing	Absent	Prolonged straightening	Intermittently standing	Good weight bearing	Stands holding on
30	Prone (pushing up on arms)	No chest supt		Head + chest up	Sustained chest support	Pushes up and on to knees
31	Tone	Abnormal		75 Questionable	30 Normal	
32	Laterality	Marked		77 Minimal	40 None	
33	Hands	Always fistd	Mostly fistd	78	50 Open	
34	Rapport (with examiner)	Cplt apathetic	Unresponsive	79 Short atten-span	60 Alert & respse	80 Gd atten-span
35	Behaviour (by report)		Inconsolable	80 Irritable/jittery	Normal	
36	Language	Mute/monotone		81 Poor sleeper		
37	Anticipation	None		Antcpts ferds	Antcpts being lifted	
PROFILE TOTALS		38-39	40-41	42-43	44-45	46-47

Mobility Absent/Present (nought)
If abnormal test primary reflexes
1 2 3 4
A B I O
Abn 7Abn 7N N

COMMENT (U) if anything of note

ALL TOTALS 48-50
[] [] []
[] []

9 - 12 MONTH NEUROLOGICAL

Serial No

Identification No

--	--

--	--	--

--	--

1 - 2

3 - 7

COL SCORE

9	Eye Appearance	A Nerve palsy	0	9 Strabismus	1	52 Questionable	2	84 Normal	3	4
10	Visual fields	B Definite unilateral defect	0	10 Doubtful	1	53	2	85 Full	3	4
10	Visual acuity	C No regard	0	11 Regards brick	1	54	2	86 Regards minute object	3	4
11	Hearing (localization)	D No localization	0	12 Vague localization	1	55	2	87 Localizes in directly above ear level	3	10+
12	Grasp of raisin	E No grasp	0	13 Partial Pincer	1	56 True Pincer	2	88 Pointing	3	4
11	Manipulation	F Cannot hold object in each hand	0	14	1	57	2	89 Picks up two bricks	3	11+
14	Movements of arm	G Abnormal	0	15 Over pronation	1	58 Clumsy	2	90 Normal	3	4
15	trunk & leg	H Abnormal	0	16	1	59 Awkward	2	91 Normal	3	4
16	Balance	I Needs support sitting	0	17 Balance poor sitting	1	60	2	92 Sits & turns round for objects	3	12+ support
17	Upward progression	J Does not get to sitting	0	18 Gets to sitting	1	61	2	93 Pulls up to stand	3	13+
18	Weight bearing	K No weight bearing	0	19 Some weight bearing	1	62	2	94 Stands alone momentarily	3	14+
19	Mobility	L Immobile	0	20 Purposeful with no achievement	1	63	2	95 Purposeful with achievement	3	4
20	Walking	M No stopping reaction	0	21	1	64	2	96 Walks round furniture	3	15+
21	Prone	N Head up only	0	22 Arm/chest support	1	65	2	97 Up on to knees	3	4
22	Resistance to passive movements	O Arm extension < 90°	0	23	1	66	2	98 To 180° easily	3	4
23		P Hip abduction < 70°	0	24	1	67	2	99 To 70° easily	3	4
24		Q Popliteal angle < 100°	0	25	1	68	2	100 To 100° easily	3	4
25		R Foot dorsiflexn < 90°	0	26	1	69	2	101 To 90° easily	3	4
26	Reflexes: Diceps	S No response	0	27	1	70	2	102 Normal	3	4
27	Adductor	T No response	0	28	1	71	2	103 Normal	3	4
28	Knee	U No response	0	29	1	72	2	104 Normal	3	4
29	Cleonus	V Sustained	0	30 Unsustained	1	73	2	105 Absent	3	4
30	Hyperaemia	W Def. present	0	31	1	74	2	106 None	3	4
31	Response (with examiner)	X Completely apathetic	0	32 Unresponsive	1	75	2	107 Alert and responsive	3	4
32	Response to dressing	Y No help	0	33	1	76	2	108 Great attention span	3	4
33	Language verbal	Z Does not say mama/dada	0	34	1	77	2	109 Multi-syllabic babble	3	4
34	Understanding	AA No anticipation	0	35 Obvious anticipation	1	78	2	110 Understands NO	3	4
35	Non-verbal	AB Mute/monotone	0	36	1	79	2	111 Indicates wants	3	4
36	Laterality	AC Marked	0	37	1	80	2	112 Shakes head NO	3	4
37	Sleep	AD Waken regularly inconsolable	0	38 Waken regularly easily settled	1	81	2	113 Sleeps through night	3	4
GRAND TOTALS		38-39	40-41	42-43	44-45	46-47				

Abnormality: Quiet, undemanding, whining/demanding, alert/active, overactive

If abnormality, test: Plantars Parachute

Response: (Punch 0 if abnormal)

NORMAL	ABNORMAL
--------	----------

COMMENT (U) if anything of note

ALL TOTALS 48-50

COL 78

A	B	C	D	E	F	G	H	I	J	K	L	M	N
1	2	3	4	5	6	7	8	9	10	11	12	13	14
Abn	Abn	7N	N	65	65-69	70-74	75-79	80-84	85-89	90-94	95-99	100-104	105-109

APPENDIX VMEAN WEIGHTS (KG)

AGE	MALES		FEMALES	
	VLBW	CONTROLS	VLBW	CONTROLS
6 MO	6.1	8.2	6.04	7.4
9 MO	7.6	8.8	7.97	8.8
12 MO	8.6	9.7	8.8	9.3

MEAN LENGTHS (CM)

AGE	MALES		FEMALES	
	VLBW	CONTROLS	VLBW	CONTROLS
6 MO	62	63.8	59.3	62.6
9 MO	67.6	68.1	67.8	69.3
12 MO	70.7	71.1	70.9	73.0

MEAN HEAD CIRCUMFERENCES (CM)

AGE	MALES		FEMALES	
	VLBW	CONTROLS	VLBW	CONTROLS
6 MO	43.2	44.5	41.9	43.4
9 MO	45.4	45.7	45.6	45.7
12 MO	46.1	46.9	46.0	46.9