## FLUID INTAKE AND WEIGHT LOSS DURING THE FIRST TEN DAYS OF LIFE AND RISK OF BRONCHOPULMONARY DYSPLASIA IN (VLBW) INFANTS

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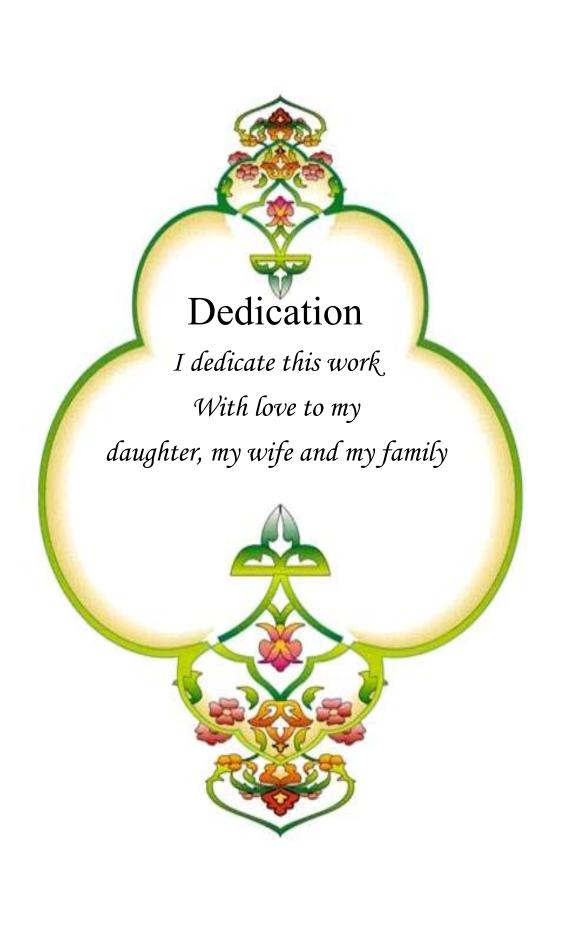
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#### **LIST OF ABBREVIATIONS**

**ABG** Arterial blood gasses.

**AGA** Appropriate for gestational age.

**APH** Antepartum hemorrhage.

**B.W** Birth weight

**BPD** Broncho pulmonary dysphasia.

CBC Complete blood count.CI Confidence interval.CLD Chronic lung disease.

**CPAP** Continue positive airway pressure.

**CRP** C–reactive protein.

**D.M** Diabetes mellitus.

 $\mathbf{D}_1$ . **W** Dextrose  $\mathbf{V}_1$  % in water

**D.W** Dextrose % in water

**ELBW** Extremely low birth weight.

FIO, Fraction of inspiratory oxygen.

**G.A** Gestational age.

**H.S** Highly significant.

**ICH** Intra cranial hemorrhage.

**IUGR** Intra uterine growth retardation.

**IVH** Intraventricular hemorrhage.

L/S ratio Lecithin to sphingomilin ratio.

**LBW** Low birth weight.

**LGA** Large for gestational age.

MOD Mode of delivery .

N.S Non significant .

**NBW** Normal birth weight.

**NEC** Necrotizing enterocolitis.

**NICU** Neonatal intensive care unit.

**P** Probability.

Pa O<sub>Y</sub> Pressure o<sup>Y</sup>.

PCO<sub>Y</sub> Pressure co<sub>Y</sub>.

**PDA** Patent ductus arteriosus.

**PEEP** Peak end expiratory pressure.

**PIP** Peak inspiratory pressure.

**PPV** Positive pressure ventilation.

**PROM** Premature rupture of membrane.

**PT** Preterm.

**RDS** Respiratory distress syndrome.

**ROP** Retinopathy of prematurity.

S Significant.

**SD** Standerd deviation.

**SGA** Small for gestational age.

SOY Oxygen saturation.
SOD Super oxide dismutase.

**SRT** Surfactant replacement therapy.

**t** Student t test

**VEGF** Vascular endothelial growth factor

**VLBW** Very low birth weight.

X' Chi square test.

### **INTRODUCTION**

Infants were defined as suffering from Bronchopulmonary dysplasia (BPD) if they were on oxygen support  $\geq \text{TA}$  days. These newborns were then reassessed when they reached TA weeks corrected GA (if GA < TY weeks) or at hospital discharge (*Ehrenkranz et al.*, Y···o).

Those who were room air at the time of reavaluation were classified as having mild BPD. Those receiving less than  $\checkmark \cdot \%$  fraction of inspired oxygen (FiO $\checkmark$ ) were classified as having moderate BPD and those on FiO $\checkmark > \checkmark \cdot \%$  and/or continous positive airway pressure (CPAP) and/or mechanical ventilation were classified as having severe BPD (*Ehrenkranz et al.*,  $\checkmark \cdot \cdot \circ$ ).

BPD is still the most common cause morbidity among (VLBW) newborns, although the incidence, risk factors and severity of the disease have changed substantially since the introduction of new treatments and mechanical ventilation techniques (*Monte et al.*, \*\*...\*\*).

Pathogenesis of BPD is multifactorial, including immaturity, barotrauma or volutrauma and oxygen toxicity (*Bancalari et al.*, \*\*.\*\*\*\*).

Excessive ingestion of liquids and sodium in these high risk neonates during the early postnatal period has been suggested as an additional risk factor for the development of BPD (*Hartnoll et al.*, \*\*.\*.).

Body water content is very high in VLBW infant and a large proportion of the body water is in the extracellular fluid (ECF) compartment. During the first week of life, there is a physiologic contraction of the ECF which is associated with weight loss during the early neonatal period. This is achieved by fluid intake that is less than the amount of water excreted through the kidney in the form of postnatal diuresis and via insensible water loss. It is postulated that this physiologic process of ECF contraction may not occur if excessive fluid and/or sodium is given during the critical period (*Oh et al.*, \*\*.\*\*\*).

High fluid intake with persistent expanded ECF is associated with a higher incidence of symptomatic patent ductus arteriosus (PDA) and necrotizing enterocolitis (NEC), also there is suggestive evidence that PDA is associated with an increased incidence of (BPD) (*Oh et al.*, \*\*\(\mathcal{T}\cdot\)\).

Strategies and interventions that might reduce the incidence of BPD have been widely investigated. Recently, on a multicenter study, it was reported that administration of caffeine during the first days of life was capable of reducing (BPD) incidence in a population of VLBW newborns (*Schmidt et al.*, \*\*••\*\*\*).

## **AIM OF THE WORK**

The aim of this work is to demonstrate the association between fluid intake and weight loss during the first ' days of life and the risk of bronchopulmonary dysplasia (BPD) in VLBW infants.

## Chapter (1)

# PREMATURE AND LOW BIRTH WEIGHT INFANTS

#### **Definitions:**

The World Health Organization defined **preterm infants**, as live born infants delivered before  $\[mathbb{r}\]$  weeks from the first day of last menstrual period. Birth weight is governed by two major considerations, the duration of gestation and the intrauterine growth rate. **Low birth weight (LBW)** that is birth weight equal to or less than  $\[mathbb{r}\]$  or may be caused by a short period of gestation, intrauterine growth retardation (IUGR) or both (*Graham*,  $\[mathbb{r}\]$ .

Very low birth weight (VLBW) infants are infants who weight less than 'o'' gm, while extremely low birth weight infants (ELBW) are infants who weigh less than ''' gm. Both of them are predominantely premature, but some are small for date at a later gestation (*Stoll and Kleigman*, '''').

LBW infants can be grouped into three categories, firstly, infants with LBW who are prematurely delivered (before TV weeks) and are appropriate for gestational age (Preterm- AGA). Secondly, infants with LBW who are born at TV weeks or later, and inappropriately small for gestational age (SGA). Lastly, infants who are prematurely

delivered but whose weight is still inappropriately small for gestation age (Preterm -SGA) (Lee and Cloherty,  $r \cdot \cdot \lambda$ ).

In developing countries, approximately  $\checkmark \cdot \%$  of LBW have intrauterine growth retardation, while in developed countries  $\checkmark \cdot \%$  of LBW have IUGR. Intrauterine growth retarded infant (I U G R), have a greater morbidity and mortality than appropriately grown gestational age infants (*Stoll and Kleigman*,  $\checkmark \cdot \cdot \land$ ).

#### **Incidence:**

The incidence of prematurity is very difficult to be determined, until recently, all low birth weight babies were lumped together as being premature. The incidence of hospital births below Your gm, is approximately Y.7% in the United States, 7.0% in Great Britain, 0.0% in Sweden, Y% in France, and You in Japan. The low birth weight rate has increased because of an increased number of preterm births. You of LBW infants in the United States have IUGR and were born after YV weeks. VLBW infants weigh less than Your gm and are predominately premature. In the United States, the VLBW rate is approximately Y.5% and their survival is directly related to birth weight (Horber et al., Your).

In order to have the incidence of prematurity in Egypt, we should have national or semi-national survey and this