# **Original Research Paper**

# **Determination of Serum Zinc Concentration in SGA Neonates as Compared to AGA Neonates**

# Singh J\* Kaur M\*\* Gupta M\*\*\*, Kaur D\*\*\*\*

\* Professor & Head \*\*\* Junior Resident, Deptt of Pediatrics, Govt. Medical College, Patiala \*\* Professor Deptt of Biochemistry, Govt. Medical College, Patiala \*\*\*\*MBBS Student

## **Corresponding Author**

Dr. Maninder Kaur, Professor Department of Biochemistry, Govt. Medical College, Patiala H. No. 53, B/2, Model town, Patiala (Pb) Phone: +91 98159-04935 E-mail: mkaur68@yahoo.com

### Article History

Received Oct 25, 2018 (Received in revised form Nov 15, 2018) Accepted on Nov 16, 2018

**Abstract:** Birth weight is the single most important determinant of infant survival in developing countries. Low birth weight is associated with poor subsequent growth in infancy and childhood with increased morbidity from infectious diseases and compromised congnitive and behavioural development.

Zinc is required to maintain the normal structure and function of multiple enzymes including those that are involved in transcription and translation of genetic material and cell division and growth and development. In our study Serum Zinc levels were compared in small for gestational age babies with respect to appropriate for gestational age babies. It was found that maternal Zinc levels affect the weight of the baby. So Zinc supplements during pregnancy lead to normal birth weight babies.

**Keywords-** Zinc, SGA Babies, Birth Weight, Enzymes, Growth and Development.

### © 2018 JCGMCP. All rights reserved

### Introduction

determinant of infant survival in developing countries. It is estimated to be responsible for >70% perinatal deaths, 90% of neonatal deaths deficiency leads to decreased foetal growth & and 50% of infant deaths. <sup>[1]</sup> Low birth weight is associated with poor subsequent growth in infancy and childhood with increased morbidity from infectious diseases and compromised cognitive and behavioural development.<sup>[2]</sup>

structure and function of multiple enzymes retain micronutrients and increased requirement including those that are involved in transcription for catch up growth, thus making them vulnerable and translation of genetic material and cell for zinc deficiency.<sup>[6]</sup> This contributes to divison.<sup>[3]</sup> Due to effect on DNA and RNA synthesis. zinc helps in cell replication and differentiation of ophthalmologic morbidity. chondrocytes, osteoblasts, fibroblasts, and in cell Aims and Objectives transcription and synthesis of somatomedin C, osteocalcin, alkaline phosphatase and Gestational Age babies with respect to metabolism of carbohydrates, proteins and fats. Zinc helps in hormonal mediation by growth tertiary care hospital whether term or preterm. hormone synthesis, secretion and action on Materials and Methods somatomedin C in liver cells and activation of

somatomedin C in bone cartilage. It helps in Birth weight is the single most important synthesis of Insulin, thyroid hormone and Vitamin D, all of which are required for growth.<sup>[4]</sup>

There is evidence that maternal zinc development that can cause foetal distress leading to assisted or operative delivery which can lead to preterm delivery, neonatal sepsis, birth asphyxia & even stillbirth.<sup>[5]</sup> Small for gestational (SGA) born babies have small liver and inadequate Zinc is required to maintain the normal stores of zinc, limited capacity to absorb and substantial neuro-cognitive, pulmonary and

To compare serum zinc levels in Small for Appropriate for Gestational Age babies in a

This study was conducted on 100 newborns

with birth weight small for gestational age as outother causes that can affect zinclevels in babies. study group which were delivered in Deptt. of Inclusion criteria: Obstetrics and Gynaecology, Govt. Medical College, Patiala and admitted to neonatal section weeks) or preterm (<37 weeks) were included in of Deptt. of Paediatrics, Govt. Medical College study group and AGA babies both term and Patiala. 100 newborn term AGA and preterm AGA preterm were taken as controls. babies were taken as control group. Approval of Exclusion criteria: All newborns whose mothers ethics committee was taken along with written were having intrauterine infections, toxaemia of consent from the parents.

A detailed antenatal and clinical history of the and all birth-asphyxiated babies. mother covering personal history, past history of **Results** any medical illness like pregnancy induced hypertension, diabetes mellitus etc., previous in the study group were 55% and 45% and in the obstetrical history like previous number of control group were 56% and 44% respectively. childbirths, abortions, perinatal loss, mode of The mean gestational age in the study group was delivery, family history of tuberculosis/diabetes 37.13± 2.33 weeks and 36.82 ± 2.50 weeks in the mellitus/hypertension etc., socio-economic control group. The maximum number of cases in status and present complaints were taken to rule both the groups were in the 39<sup>th</sup> week of gestation.

All SGA babies whether term (37-41

pregnancy, diabetes mellitus, hepatitis, smokers

The male and female percentage of babies

Group	No.	Mean ±SD (µg/dl)	't' value	'p' value	Sig.
Study	100	56.8 ±40.6	-6.1	<0.001	HS
Control	100	107.4±72			

The mean ( $\pm$ SD) serum zinc levels of the study and the control groups were 56.8  $\pm$  40.6  $\mu$ g/dl and  $107.4 \pm 72 \,\mu\text{g/dl}$  respectively. Statistically highly significant difference was found in the mean serum zinc levels between the two groups. (Normal zinc levels =  $60-120 \mu g/dl$ ).

Group	No.	Mean ±SD (µg/dl)	't' value	'p' value	Sig.
Preterm SGA	38	46.26± 22.54	2.07	<0.05	S.
Term SGA	62	63.35± 47.47			

The mean serum zinc levels of the preterm SGA group and term SGA group were 46.26 ±22.54  $\mu$ g/dl and 63.35±47.47  $\mu$ g/dl respectively. Statistically significant difference was found in the mean serum zinc levels between the two groups.

Group	No.	Mean ±SD (µg/dl)	't' value	'p' value	Sig.
Term	120	81.13 ± 52.25	-0.277	>0.05	NS
Preterm	80	83.68 ± 77.88			

The mean (±SD) serum zinclevels of the term and preterm group whether they are SGA born or AGA born babies are  $81.13 \pm 52.25 \,\mu\text{g/dl}$  and  $83.68 \pm 77.88 \,\mu\text{g/dl}$  respectively. A statistically non significant difference was found in the mean serum zinc levels between the two groups (p> 0.05, as computed from the SEDM and the t-test).

# **Discussion:**

serum zinc levels of the study and the control morbidity and mortality depends on the birth groups were 56.8  $\pm$  40.6  $\mu$ g/dl and 107.4  $\pm$  72 weight. Therefore zinc supplementation to µg/dl respectively. Statistically highly significant pregnant mothers could prevent preterm and difference was found in the mean serum zinc LBW babies. levels between the two groups. A similar study **Conflict of Interest: None** done by Elizabeth et al (2007) showed the same observation in SGA babies taken in study group and term normal weight babies taken in control group. Serum zinc levels were  $70.25 \pm 24.5$ , 78.09  $\pm$  18.39 and 92.24  $\pm$  19.4 µg/dl respectively. The 2. difference between study and control group was statistically significant.<sup>[7]</sup> Another study by Akram et al (2011) compared serum zinc levels in babies born SGA with the babies born large for gestational age (LGA). Zinc levels were 78 µg/dl in the SGA group and 92  $\mu$ g/dl in the LGA group. The 4. difference was statistically significant in the two groups.<sup>[8]</sup> A study was done by Ozdemir et al (2007) on the estimation of zinc levels from cord blood in SGA, AGA and LGA babies. Zinc levels were below 100  $\mu$ g/dl in the SGA group and 150 µg/dl in the AGA group. It showed significantly lower zinc levels in the SGA group. The method used for estimation was atomic absorption spectrophotometery.<sup>[9]</sup>

# **Conclusion:**

The present study demonstrates that maternal zinc level affects the weight of the baby but not the length or head circumference of the baby. Zinc deficiency in mothers during pregnancy has adverse outcome on foetus. Therefore preventing zinc deficiency in mother leads to improvement in prenatal growth which translates into improvement in postnatal growth, improvement in immune functions & decreased

risk of morbidity in infancy. Prognosis of the baby In the present study, the mean (±SD) in neonatal period and infancy regarding

### **References:**

- Smith Vincent C. The High Risk Newborn: Anticipation, 1. Evaluation, Management and Outcome. Manual of Neonatal Care, edition 7; chapter 7:74-90
- Osrin David, Costello Anthony M.de L. Maternal nutrition and fetal growth: practical issues in international health. Semin Neonatol 2000;5:209-219
- Bernstein Ira M, Horbar Jeffrey D, Badger Gary J, Ohlsson Arne, Golan Agneta. Morbidity and mortality among low-birth-weight neonates with intrauterine growth restriction. Am J Obstet Gynecol 2000;182:198-206
- Merialdi Mario, Caufield Laura E, Zavaleta Nelly, Figueroa Alberto, Dominici Francesca. Randomized controlled trial of prenatal zinc supplementation and development of fetal heart rate. American Journal of Obstetrics and Gynecology 2004;190:1106-12
- Janet C. King. Effect of Zinc Supplementation on 5. Pregnancy and Infant Outcomes: A Systematic Review. Paediatr Perinat Epidemiol. 2012; 26(1): 118-137
- 6.. Bhatnagar Shinjini, Natchu Uma Chandra Mouli. Zinc in Child Health and Disease. Indian J Pediatr 2004;71(11):991-995
- Elizabeth K.E, Krishnan Viji. Umbilical cord blood 7. nutrients in low birth weight babies in relation to birth weight and gestational age. Indian J Med Res 2008;128:128-133
- 8. Akram Shahzad K. Placental IGF-1, IGFBP-1, zinc and iron and maternal and infant anthropometry at birth. Acta Paediatrica 2011;100:1504-1509
- Ozdemir U. Correlation between birth weight , leptin and copper levels in maternal and cord blood. J Physiol Biochem2007;63(2):121-128