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Research Article

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Role of zinc in febrile seizures

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ABSTRACT

Background: Febrile convulsion is the most common type of seizures in childhood. Zinc has been studied in the context of pathophysiology of febrile seizure. The objective of the present study was to compare the serum zinc levels in children with febrile seizure, acute febrile illness without seizure and in healthy children.

Methods: The present study was a cross sectional study. A total of 150 infants and children aged between 6 months to 5 years fulfilling the inclusion criteria were included. The study comprised of 3 groups - Group A: Children with febrile seizures (50 cases), Group B: Children with only fever, but no seizures (50 cases) and Group C: Healthy children (50 cases). Serum zinc was measured by colorimetric method and compared among the groups using statistical methods.

Results: There was no significant difference in sex, age, weight, height and head circumference between the three groups (P >0.05). There was no statistically significant difference between the groups regarding temperature at admission. Mean serum zinc level was $50.49 \pm 15.17 \ \mu gm/dl$, $61.53 \pm 16.87 \ \mu gm/dl$ and 85.70 ± 20.76 in Group A, Group B and Group C respectively. It was significantly lower in Group A compared to Group B, it was significantly lower in Group B compared to Group C. Serum zinc level was less than 65 $\mu gm/dl$ in 65% children in Group A and 35% in Group B.

Conclusions: Serum zinc level was lower in children with simple febrile seizures as compared to children with acute febrile illness and healthy children which were found to be statistically significant.

Keywords: Febrile seizure, Serum zinc level, Fever

INTRODUCTION

Febrile Seizure (FS) is one of the most common neurological conditions in childhood. Although febrile seizure are generally of a benign nature, they remain a serious condition currently due to the recurrence rates seen in some cases and the slight risk they carry of developing into epileptic attacks.

The etiology of febrile seizure is still not clear. Various factors have been described in the pathophysiology of febrile seizures like bacterial and viral infections, susceptibility of the immature brain to temperature,

association with interleukins, circulating toxins, trace element deficiency and iron deficiency. Role of trace elements like selenium, magnesium, copper and zinc have been described in association with febrile seizures. Trace elements appear to play a role by their ability to modulate neurotransmission by acting on ion channels and their coenzyme activity.¹⁻⁶

Zinc is an important element in growth, development and normal brain function. It is also an important cofactor for different enzymes, and is involved in cellular growth and differentiation, enzymatic activity of different organs, proteins and cellular metabolism. In brain, zinc is present

in synaptic vesicles in subgroup of glutaminergic neurons. In this form it can be released by electrical stimulation and may serve to modulate responses at receptors for number of different neurotransmitters. These include both excitatory and inhibiting receptors particularly NMDA (N-methyl-D aspartate) and GABA (Gamma aminobutyric acid) receptors respectively.^{7,8}

Decreased zinc levels modulate the activity of glutamic acid decarboxylase, the rate limiting enzyme in the synthesis of GABA, which is a major inhibitory neurotransmitter. Any abnormalities of GABAergic function, including synthesis, synaptic release, receptor composition, trafficking or binding, and metabolism, can each lead to a hyperexcitable, epileptic state. Pinc has an inhibitory effect on NMDA receptors which is responsible for excitatory phenomenon after binding with glutamate. Thus decreased zinc levels may play a role in pathogenesis of febrile seizures.

This study aims to determine zinc levels in children who have febrile convulsions and thus help to reveal possible associations between zinc deficiency and febrile convulsions.

METHODS

The present study was a cross sectional study conducted at Basaveshwara medical college and hospital, Chitradurga during the period January 2014 to January 2015. A total of 150 infants and children aged between 6 months to 5 years fulfilling the inclusion criteria were included. The study comprised of 3 groups - Group A: Children with febrile seizures (50 cases), Group B: Children with only fever, but no seizures (50 cases) and Group C: Healthy children (50 cases). Inclusion criteria were children with simple febrile seizure and acute febrile illness without seizures and normal development. Febrile seizures were defined as seizures accompanied with fever (≥38°C) without Central Nervous System (CNS) infection. Children with diarrhea, pneumonia, protein energy malnutrition, developmental delay and or

neurological deficit and children on zinc therapy were excluded from the study. A written informed consent was obtained from parents/guardians of all the children after fully explaining the study procedure. The Institutional ethical committee of our hospital approved the study. Socio-demographic data, seizure details, nature of febrile illness, family history of epilepsy/febrile seizures, temperature at admission, nutritional status and vital signs namely heart rate, respiratory rate and blood pressure were measured. The axillary temperature was recorded in all children with mercury thermometer placed in axilla for three minutes, followed by general examination and systemic examination in detail. Taking aseptic precaution, 2 ml of blood from venipuncture using 22 gauge sterile needle, was collected in morning. non-fasting state within 24 hours of contact of patient in all the 3 groups. The sample was centrifuged for 3-4 minutes at 3000-4000 rpm; serum thus obtained is collected and preserved at 2-8°C in sterile deionized plain vials. Estimation of serum zinc was done within 6 hours of collection. Method used was based on colorimetric test kit, reagent used was 2-(5-bromo-2-pyridylazo)-5-(Npropyl-N-sulphopropylamino)-phenol. Zinc forms a red chelate with it. Increase in the absorbance of wavelength 560 nm can be measured and is proportional to concentration of the zinc.

In the present study serum zinc level less than 65 µgm/dl was taken as cut off for zinc deficiency. ¹¹ The three groups included in the study were compared with respect to serum zinc level.

Statistical methods used were descriptive statistics, contingency table analysis, independent sample t test, 2 way ANOVA. All the statistical calculations were done through SPSS 16.0 for windows.

RESULTS

There was no significant difference regarding sex, age, weight, height, head circumference temperature at admission between the 3 groups.

Table 1: Baseline characteristics, anthropometric parameters, temperature at admission and mean serum zinc levels in 3 groups.

Variables	Febrile seizure group (N=50)		Febrile group (N=50)	Normal children (N=50)
Sex	Male	35 (70%)	31 (62%)	34 (68%)
	Female	15 (30%)	19 (38%)	16 (32%)
Age (months)	27.13 ± 15.73		28.49 ± 16.51	28.86 ± 16.51
Mean weight (kg)	11.62 ± 2.61		12.02 ± 2.58	12.24 ± 2.58
Mean height (cm)	81.96 ± 10.91		85.62 ± 14.16	85.92 ± 14.16
Mean head circumference (cm)	48.6 ± 2.44		48.19 ± 2.16	48.80 ± 2.50
Mean temperature at admission (°C)	38.65		38.64	
Mean serum zinc (µgm/dl)	50.49 ± 15.17		61.53 ± 16.87	85.70 ± 20.76

The mean serum zinc was significantly decreased in febrile seizure group compared to febrile illness group and normal children (P <0.01). There was no statistically significant difference in the serum zinc level in relation to age and sex between the two groups. 65% of Group A and 35% of Group B had serum zinc <65 μ gm/dl. There was a statistically significant difference in serum zinc between the groups (P <0.001).

DISCUSSION

The results of this study revealed that the mean serum zinc level in children affected with febrile seizure is lower than in febrile group and normal healthy children and the difference is statistically significant.

Limited numbers of studies have been conducted regarding the role of zinc in occurrence of febrile seizures. Burhanoğlu M et al. reported that the average level of serum zinc in children affected with febrile seizure was less than control group. 12 Ehsani F et al. carried out study on 34 children with febrile seizure and 58 healthy children revealed that the serum zinc level in children with febrile seizure was lower than those in control group and the difference was significant, statistically. 13 Tütüncüoğlu S et al. reported that the serum zinc level among children with febrile seizure was considerably lower than those in control group. 14 In a study by Hamed SA et al., it was shown that the trace elements such as zinc have crucial role in pathogenesis of seizures.¹⁵ The study of Gündüz Z et al. on 102 children with febrile seizures indicated that the serum zinc level in the group affected with febrile seizures was significantly lower than those in control group. 16 In a very latest study by Mishra OP et al. on 20 children with febrile seizures and 48 children as control group, it was reported that the serum zinc level in children affected with febrile seizure was lower than those in control group, and the difference was significant.¹⁷ In contrast to our study, Kafadar I et al. found no significant difference in serum Zinc concentration in children with febrile convulsion and other two control groups. This may be due to the smaller sample size in their study.¹⁸

The reason for reduction of serum zinc level in patients affected with febrile seizure is not clear. However, fever and acute infection may have some roles in developing such condition. ¹⁹ It is believed that the release of Tumor Necrosis Factor (TNF) and interleukin (IL) during fever or tissue injury may result in reduction of serum zinc level. ¹³ Izumi Y et al. proposed that the hypozincemia during fever trigger the NMDA receptor, one of the members of glutamate family of receptors, which may play an important role in the initiation of epileptic discharge during febrile seizures. ²⁰

The role of zinc in nervous system function has been broadly discussed in literature. ^{21,22} Brain contains an abundant value of zinc, especially in hippocampus region. Five to fifteen percent of zinc is concentrated as

vesicle zinc in glutamatergic synapses.²¹ Zinc acts as a neurotransmitter and improves the communicating and locomotive function, and also evolution of neurological system.²³ Zinc deficiency diminishes hippocampal zinc and leads to seizure discharge.²² However, the mechanism of this altered serum zinc level in the febrile seizure group is poorly understood. Further studies are needed to identify the cause of this observation. Serum zinc levels are influenced by the time of day, the specific disease, or the presence of other trace elements.²⁴ Therefore, a study design considering these effects is needed to further explain hypozincemia in febrile seizure children.

Limitations of the study were zinc estimation in the present study was done by colorimetric method. Atomic Absorption Spectrometry (AAS) is a more accurate method to measure serum zinc level. Though AAS was not done, there is a good correlation between AAS and colorimetric method which is used in the present study. The same children in simple febrile seizure group should have had a follow up serum zinc estimation when healthy, which would have shown the baseline serum zinc status. Further studies are required in this aspect.

This study showed that serum zinc level decreased during infection and that decrease was more significant in patients with febrile seizures.

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Ethical approval: The study was approved by the

institutional ethics committee

REFERENCES

- 1. Millichap JG, Millichap JJ. Role of viral infections in the etiology of febrile seizures. Pediatr Neurol. 2006 Sep;35(3):165-72.
- Holtzman D, Obana K, Olson J. Hyperthermiainduced seizures in the rat pup: a model for febrile convulsions in children. Science. 1981;213:1034-6.
- Tsai FJ, Hsieh YY, Chang CC, Lin CC, Tsai CH. Polymorphisms for interleukin 1 beta exon 5 and interleukin 1 receptor antagonist in Taiwanese children with febrile convulsions. Arch Pediatr Adolesc Med. 2002;156:545-8.
- 4. Virta M, Hurme M, Helminen M. Increased plasma levels of pro- and anti-inflammatory cytokines in patients with febrile seizures. Epilepsia. 2002;43:920-3.
- 5. Amiri M, Farzin L, Moassesi ME, Sajadi F. Serum trace element levels in febrile convulsion. Biol Trace Elem Res. 2010;135(1):38-44.
- Kumari PL, Nair MK, Nair SM, Kailas L, Geetha S. Iron deficiency as a risk factor for simple febrile seizures - a case control study. Indian Pediatr. 2011;49:17-9.
- 7. Ebadi M, Wilt S, Ramaley R. The role of zinc and zinc-binding proteins in regulation of glutamic acid

- decarboxylase in brain. In: Ebadi M, Wilt S, Ramaley R, eds. Chemical and Biological Aspects of Vitamin B6, Catalysis. 1st ed. New York: Alan R Liss; 1984 (Part A): 255-275.
- 8. Cossart R, Bernard C, Ben-Ari Y. Multiple facets of GABAergic neurons and synapses: multiple fates of GABA signalling in epilepsies. Trends Neurosci. 2005;28:108-15.
- 9. Macdonald RL, Kang JQ. Molecular pathology of genetic epilepsies associated with GABAA receptor subunit mutations. Epilepsy Curr. 2009;9:18-23.
- Peters S, Koh J, Choi W. Zinc selectively blocks the action of NMDA on cortical neurons. Science. 1987;236:589-93.
- 11. Report of WHO/UNICEF/IAEA/IZiNCG Interagency Meeting on Zinc Status Indicators. Executive summary. Recommendations for indicators of population zinc status. Food Nutr Bull. 2007;28:S399-400.
- 12. Burhanoğlu M, Tütüncüoğlu S, Coker C, Tekgül H, Ozgür T. Hypozincaemia in febrile convulsion. Eur J Pediatr. 1996 Jun;155(6):498-501.
- 13. Ehsani F, Vahid-Harandi M, Kany K. Determination of serum zinc in children affected by febrile convulsion and comparison with control group. J Iranian Medi Sci Univ. 2006;12:219-76.
- Tütüncüoğlu S, Kütükçüler N, Kepe L, Coker C, Berdeli A, Tekgül H. Proinflammatory cytokines, prostaglandins and zinc in febrile convulsions. Pediatr Int. 2001 Jun;43(3):235-9.
- 15. Hamed SA, Abdellah MM. Trace elements and electrolytes homeostasis and their relation to antioxidant enzyme activity in brain hyperexcitability of epileptic patients. J Pharmacol Sci. 2004 Dec;96(4):349-59.

- Gündüz Z, Yavuz I, Koparal M, Kumandaş S, Saraymen R. Serum and cerebrospinal fluid zinc levels in children with febrile convulsions. Acta Paediatr Jpn. 1996 Jun;38(3):237-41.
- 17. Mishra OP, Singhal D, Upadhyay RS, Prasad R, Atri D. Cerebrospinal fluid zinc, magnesium, copper and gamma-aminobutyric acid levels in febrile seizures. J Pediatr Neurol. 2007;5:39-44.
- 18. Kafadar I, Akinci AB, Pekun F, Adal E. The role of serum zinc level in febrile convulsion etiology. J Pediatr Inf. 2012;6:90-3.
- Garty BZ, Olomucki R, Lerman-Sagie T, Nitzan M. Cerebrospinal fluid zinc concentrations in febrile convulsions. Arch Dis Child. 1995 Oct;73(4):338-41
- Izumi Y, Ishii K, Akiba K, Hayashi T. Hypozincemia during fever may trigger febrile convulsion. Med Hypotheses. 1990 May;32(1):77-80
- 21. Cole TB, Robbins CA, Wenzel HJ, Schwartzkroin PA, Palmiter RD. Seizures and neuronal damage in mice lacking vesicular zinc. Epilepsy Res. 2000;39(2):153-69.
- Tapiero H, Tew KD. Trace elements in human physiology and pathology: zinc and metallothioneins. Biomed Pharmacother. 2003 Nov;57(9):399-411.
- 23. Mahyar A. The preventive role of zinc from communicable and non-communicable diseases in children. NCD Malaysia. 2005;4:21-5.
- 24. Maret W, Sandstead HH. Zinc requirements and the risks and benefits of zinc supplementation. J Trace Elem Med Biol. 2006;20:3-18.

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