



Serum zinc, copper, selenium, and lead levels in children with chronic renal failure

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ARTICLE INFO	ABSTRACT
<p>Article type Review article</p> <p>Article history Received: 14 Dec 2013 Revised: 24 Dec 2013 Accepted: 9 Jan 2014</p> <p>Keywords Chronic kidney disease Renal failure Trace elements</p>	<p>In the last two centuries, chronic kidney disease (CKD) and renal failure (RF) have been considered as the main medical problems which were fatal in many cases. Trace elements such as selenium, copper, and zinc are the components of biological enzymes which have a crucial role in decreasing reactive oxygen. The levels of these trace elements reduce in CKD patients. Close and careful nutritional support of children with CKD, particularly in the early stages of the disease, is necessary.</p>

Please cite this paper as:

Rakhshanizadeh F, Esmaeeli M. Serum zinc, copper, selenium, and lead levels in children with chronic renal failure. *Rev Clin Med.* 2014;1(1):21-24.

Introduction

CKD is defined as Glomerular filtration rate (GFR) less than 60 ml/min per 1.73 m² (1). This definition is not convenient for children younger than 2 years, because their GFR level is lower than older children and adults (2). Some of end-stage renal disease (ESRD) causes are listed as below: autoimmune disorders, congenital renal disease (such as polycystic

kidney disease), renal toxic drugs or chemicals, glomerulonephritis, renal blind or penetrating trauma, kidney stones and infection, reflux nephropathy, and other kidney diseases (3). Two important complications of childhood CKD are retarded growth and renal osteodystrophy (ROD), all of which happened due to growth hormone axis abnormalities, vitamin D

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insufficiency, nutritional deficiency, drug toxicity, etc. Other hormonal abnormalities might occur in CKD patients such as hyperparathyroidism and hypogonadism (4).

In recent years, there is some evidence supported the theory of oxidative stress in chronic renal failure, especially in hemodialysis patients. Trace elements such as selenium, copper, and zinc are the components of biologic enzymes which have a crucial role in decreasing reactive oxygen (2).

On the other hand, changes in mineral metabolism happen with chronic kidney disease progression (5). Trace element elimination in CKD results from reduction in renal function and proteinuria which leads to (a) losing protein-bound elements (b) changes in gastrointestinal absorption due to vitamin D metabolism, and (c) the dialysis procedure. CKD-related complications such as fluid and electrolyte disturbances, renal osteodystrophy, anemia, hypertension, dyslipidemia, endocrine abnormalities, etc become worse when CKD progresses into stage G3a and G3b (6).

In Table1 the factors affected trace element concentrations have been shown.

Table1. Factors affect trace element concentration

Nutritional	Malabsorption	Distribution disorders
Inadequate intake (low income diet, malnutrition)	Gastrointestinal dysfunction	Transport disorders
Alcoholism	—	Storage inability
Increased metabolism or requirement	—	Stress conditions such as MI, infections, etc

In the first step, early and proper diagnosis of the underlying disease which leads to chronic kidney disease (CKD) is important.

Choosing the best treatment method with fewer complications is the next step. The progression of the disease could delay or halt by these strategies (7). The aim of this study was to evaluate serum concentration of trace elements in children with CKD.

Trace element concentrations might be affected by renal failure intensity and renal replacement therapy methods. In CKD, some trace elements such as cobalt, arsenic, mercury, and silicon tend to rise. Some other elements including zinc, brom, and selenium might decrease. These trends might differ in hemodialysis and peritoneal dialysis (8,9). Different studies also reported various levels of trace element concentrations (Table2). In Table2 trace element levels have been shown in different studies.

Discussion

It seems that mean blood concentrations of trace elements have a remarkable difference in hemodialysis patients. Serum trace element levels in CKD patients had a great impact on patient's general condition and might lead to cardiovascular diseases, carotid atherosclerosis, cancers, decline of renal function, renal osteodystrophy, insulin resistant, and anemia (19,20).

Daily need of zinc in children with CRF varies from 4 to 9.5 mg in different childhood stages. These ranges for copper are about 0.2 to 1 microgram (10). Although in general population zinc supplement could decrease all cause death and the risk of infection, there are few studies supported the theory of trace element impact on the CKD patients' outcome (2-6).

Malnourished children have a lower body mass index (BMI) level and are at risk of infection which is associated with a worse prognosis (8).

Zinc is required to maintain the structures and functions of more than 200 enzymes,

Table 2. Serum zinc, copper, selenium, and lead levels in patients with chronic renal failure

Author	Publication year	Patients' age	Zinc	Copper	Lead	Selenium
Tayeb (10)	2009	9-17	14.56±6.09 micromol/ dl	—	—	—
Esfahani (11)	2013	3.64 ± 11.75	0.35 ±0.077 ppm	0.533±0.231 ppm	—	—
Abiri (12)	1994	—	0.806±0.177 ppm	—	—	—
Marjani (13)	2003	50.02±16.08	78.35 ±37.46 microgram/dl	—	89.92±32.54 microgram/dl	—
Mishra (14)	2008	0-10	1.07±0.46 mg/dl	1.56±0.48 mg/dl	—	—
Ari (15)	2011	38 ±8	0.53±0.3 microgram/dl	1.54±0.34 microgram/dl	—	—
Apostolidis (16)	2002	15 ± 54	—	—	—	44±9 nanogram/dl
Guo (17)	2013	6 ± 53	7.7 ± 49.2 microgram/dl	—	—	10.2 ± 46.1 nanogram/dl
Palaneeswari (18)	2013	52.5	—	—	1.67 ± 07.32 microgram/dl	—

such as enzymes needed in transcription and translation of genetic material and cell division (11).

Lower zinc level in CKD patients might happen due to the increased urinary zinc excretion and the decreased intestinal zinc absorption. Hypozincemia is a cause of redistribution of total body zinc because liver tissue zinc had been increased in autopsy (21).

On the other hand, decrease in zinc and copper serum concentrations happen in ESRD due to the excretion of protein and albumin in urine (2).

In conclusion, close and careful nutritional support in children with CKD, particularly in the early stages of the disease, is necessary.

Acknowledgement

We would like to thank Clinical Research Development Center of Ghaem Hospital for their assistant in this manuscript. This study was supported by a grant from the Vice Chancellor for Research of the Mashhad

University of Medical Sciences for the research project as a medical student thesis with approval number of 910392.

Conflict of Interest

The authors declare no conflict of interest.

References

1. Filler G, Felder S. Trace elements in dialysis. *Pediatr Nephrol.* 2013. [Epub ahead of print]
2. Prasad AS. Discovery of human zinc deficiency: its impact on human health and disease. *Adv Nutr.* 2013;4:176-190.
3. Tomat AL, Costa Mde L, Arranz CT. Zinc restriction during different periods of life: influence in renal and cardiovascular diseases. *Nutrition.* 2011;27:392-398.
4. Nazari H, Nourmohammadi A. Status of zinc, copper and agnesium in serum and hair of insulin-dependent diabetes. *Koomesh.* 1991;1:37-41.
5. D'Ocon C, Alonso de Armiño V, Frassetto V. Levels of Zn and Cu in the serum of a diabetic population. *Rev EspFisiol.* 1978;43:335-338.
6. Jamshidi F, Tasalli E, Heshmat R, et al. Evaluation of Mg, Zn, and Cu serum levels in Diabetic and nondiabetic persons and their

- relationship with anemic situation in food regimes. *Diabetes and Lipid J.* 2010;9:249-259.
7. Metnitz GH, Fischer M, Bartens C, et al. Impact of acute renal failure on antioxidant status in multiple organ failure. *Acta Anaesthesiol Scand.* 2000;44:236-240.
 8. Bowry SK, Gatti E. Impact of hemodialysis therapy on anemia of chronic kidney disease: the potential mechanisms. *Blood Purif.* 2011;32:210-219.
 9. Zaher MM, Gaber M, Alrefaey A, et al. Assessment of some trace elements: copper, zinc and magnesium and their impact on CD3 and CD4 levels in children on chronic hemodialysis. *Life Sci J.* 2013;10:222- 230
 10. El Tayeb AA, Abd El-Mottaleb NA, Abdel Aziz EA. Relationship between serum parathyroid hormone and trace elements (serum zinc and magnesium) in hemodialyzed chronic renal failure children. *Biol Trace Elem Res.* 2009;128:128-134.
 11. Esfahani ST, Hamidian MR, Madani A, et al. Serum zinc and copper levels in children with chronic renal failure. *Pediatr Nephrol.* 2006;21:1153–1156.
 12. Yaghmaee B, Bezarafashani MR, Abiri M. Serum level of aluminum and zinc in hemodialysis patients` in kerman. *Journal of Kerman University of Medical Sciences.* 1994;1:166-170.
 13. MarjaniA, Moujerlou M, Mansourian AR. Serum zincandcopper level before and after haemodialysis. *Journal of Gorgan University of Medical Sciences.* 2003;5:10-14.
 14. Mishra OP, Pooniya V, Ali Z, et al. Antioxidant status of children with acute renal failure. *Pediatr Nephrol.* 2008;23:2047-2051.
 15. Ari E, Kaya Y, Demir H, et al. The correlation of serum trace elements and heavy metals with carotid artery atherosclerosis in maintenance hemodialysis patients. *Biol Trace Elem Res.* 2011;144:351-359.
 16. Apostolidis NS, Panoussopoulos DG, Stamou KM, et al. Selenium metabolism in patients on continuous ambulatory peritoneal dialysis. *Perit Dial Int.* 2002;22:400-404.
 17. Guo CH, Chen PC, Hsu GS, et al. Zinc supplementation alters plasma aluminum and selenium status of patients undergoing dialysis: a pilot study. *Nutrients.* 2013;5:1456-1470.
 18. Palaneeswari M S, Rajan PM, Silambanan S, Jothimalar. Blood arsenic and cadmium concentrations in end-stage renal disease patients who were on maintenance haemodialysis. *J Clin Diagn Res.* 2013;7:809-813.
 19. Elshamaa MF, Sabry S, Mokhtar I, et al. Aluminium and lead abnormalities in children on haemodialysis: relationship with some medications. *Arch Med Sci.* 2010 30;6:420-429.
 20. Tonelli M, Wiebe N, Hemmelgarn B, et al. Trace elements in hemodialysis patients: a systematic review and meta-analysis. *BMC Med.* 2009;7:25.
 21. Vanholder R, Cornelis R, Dhondt A, et al. The role of trace elements in uraemic toxicity. *Nephrol Dial Tranplant.*2002;2:2-8.