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Preliminary Communication



IMPORTANCE OF ZINC CONCENTRATION IN SEMINAL FLUID OF MEN DIAGNOSED WITH INFERTILITY

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SUMMARY - The prevalence and importance of male infertility make it a serious worldwide medical and social problem. The aim of this study was to determine if there were any significant differences in zinc concentrations in seminal fluid in men diagnosed with infertility compared to control group, and if there were, to determine how it affected the number and percentage of sperm cells with normal motility and morphology. A case-control study was conducted in the Osijek-Baranja County from January 2014 to June of 2015. The study included 276 subjects referred to biochemistry laboratory of the Osijek University Hospital Centre, 100 (36.2%) of them with low sperm count confirmed by spermiogram after 3 months. Semen samples were processed according to the World Health Organization criteria, and zinc concentrations were determined using spectrophotometry and direct colorimetry without deproteinization on a chemistry analyzer (Olympus AU 680, Beckman Coulter, Tokyo, Japan). Study results showed the men with low sperm count to be significantly older (Mann Whitney U test: p=0.013) and to have lower zinc levels (χ²-test: p<0.001). Further analysis included 100 (36.2%) patients with low sperm count. In the group of infertile men, those with low sperm count had higher zinc levels compared to men with azoospermia (Mann Whitney U test: p=0.036), suggesting a connection between lower zinc level in seminal fluid and male fertility. Zinc as a biological marker and an antioxidant affects sperm count, motility and morphology. Zinc supplemental therapy could improve seminal parameters in patients diagnosed with low sperm count. Dietary habits of the subjects differ regionally, and future research could make a solid foundation for complementary approach to treatment of male infertility.

Key words: Croatia; Infertility, male; Zinc; Case-control studies; Oligospermia; Semen

Introduction

According to the World Health Organization (WHO), infertility is defined as the inability to spontaneously conceive during one year of normal sexual activity without the usage of contraception, and demo-

(48.5 million couples), with male infertility as the cause identified in 60% and female infertility in 40% of cases. The increase of infertility is an important factor in demographic transition and imbalance, thus it significantly contributes to depopulation and is becoming one of the leading public health problems faced by more and more couples². Experts agree that

the etiology of infertility is multifactorial, with enor-

mous influence of stress, smoking, alcoholism, obesi-

graphic data show increase in the infertility rate in the most developed countries¹. In western countries, 15%

of couples are not able to conceive during one year

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ty^{3,4}, too much physical activity, exposure to toxins, radiation and high temperatures⁵. Alongside these factors, mechanical and reversible damages such as varicocele cause 25%-40% of cases of infertility. In 20%-30% of the cases, it is not possible to determine the etiology, and this kind of infertility is referred to as idiopathic⁶. When male infertility is suspected, one of the first procedures to be used is seminal fluid analysis. This procedure is simple and noninvasive while providing the necessary information for establishing the initial diagnosis. Semen analysis reveals complete absence of sperm cells (azoospermia), decreased number of sperm cells (oligozoospermia), decreased sperm motility (asthenozoospermia), or abnormal morphology (teratozoospermia). Motility, structure, sperm concentration, and the components of semen are extremely important for normal sperm function¹. By determining the biochemical components of semen (fructose, phosphatase, citric acid, zinc), using microbiological analysis, determining sperm antibodies and biological markers we get better view into reproductive health and fertility of the male partner⁷. Microelements are a group of important factors in maintaining the normal function, structure, motility and sperm count8. A few studies have shown how zinc has an antioxidative and anti-inflammatory effect9, and participates in producing antioxidative enzymes important for cell protection¹⁰. The amount of zinc in the human body is relatively constant; it makes up to 24% of all trace elements. Most of it is found in the prostate (100) times more than in blood) and is evenly distributed in the kidneys, liver, muscles and heart, but in some body fluids its concentration is variable¹¹. Zinc level in seminal fluid is related to sperm concentration and motility12. The lack of zinc leads to many different symptoms, so low zinc concentration affects sperm quality and male fertility¹³. Sperm cells are extremely sensitive to damages caused by oxidative stress because of the large amounts of unsaturated fatty acids in their membranes that are essential for sperm capacitation, acrosome reaction and sperm/oocyte interaction. During lipid peroxidation, sperms lose their membrane integrity, which leads to inactivation of inner cell enzymes, structural DNA damages and apoptosis. Lipid peroxidation of sperm membranes is considered to be the main mechanism of reactive oxygen species (ROS)induced sperm damage that leads to infertility¹⁴. The number and motility of sperm decreases due to changes in sperm DNA, thus increasing the number of morphologically abnormal sperm cells, which leads to a decreased ability of penetrating the oocyte and lower chances of insemination^{15,16}. The role of oxidative stress in male infertility has been significantly scientifically supported and evaluation of zinc deficiency could play a key role in diagnosing the cause of sperm damage.

The aim of this study was to determine if there were any significant differences in zinc concentrations in seminal fluid in men diagnosed with infertility as compared to control group, and if there were, to determine how it affected the number and percentage of sperm cells of normal motility and morphology.

Subjects and methods

Study design

A case-control study was conducted in the Osijek-Baranja County from January 2014 to June of 2015. Study subjects were referred to the Osijek University Hospital Centre by their general practitioner. Semen samples were analyzed by routine procedure at the Andrology and Biochemistry Laboratory, Osijek University Hospital Centre. The study protocol was approved by the Ethics Committee of the Osijek Faculty of Medicine (code: 2158-61-07-15-104).

Study participants

The study included 276 men, 100 (36.2%) of them diagnosed with low sperm count and 176 (63.8%) with normal spermiogram according to the 2010 WHO guidelines. Out of 100 subjects diagnosed with low sperm count, 26 had azoospermia. The including criteria were as follows: routine examinaton of the male partner of the infertile couple, inability to concieve during one year of intercourse without protection, age over 18, and at least two semen analyses. The excluding criteria were age <18 years, varicocele, prostatitis, and malignant disease. The subjects were informed on the way of preparing and obtaining the sample. The samples were obtained by masturbating into a small container, and the sampling was done after minimally 2-5 days of sexual abstinence. For this research, demographic data (age, place of residence) were used and the values of seminal parameters (zinc concentration, motility, morphology and sperm count) were established.

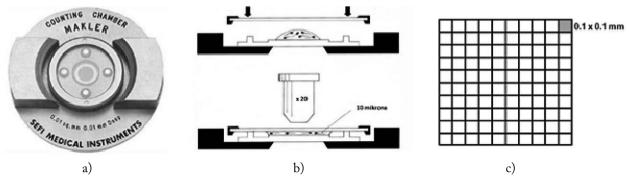


Fig. 1. (a) Makler chamber; (b) scheme of the chamber; (c) net for sperm counting. (source: http://www.sefimedical.com/images/makler_chamber_shema.gif; accessed on July 11, 2013).

Methods

Determining zinc concentrations

Zinc concentrations were determined by atomic absorption spectroscopy (AAS) at the Department of Clinical Laboratory Diagnosis, Osijek University Hospital Centre. The sperm obtained by centrifugation was processed for 10 minutes at 2000 rpm. The supernatant was diluted by adding distilled water (1:21) and zinc concentration was determined on an Olympus AU 680 autoanalyzer (Beckman Coulter, Tokyo, Japan). Zinc levels were measured in mmol/L (reference interval: 1.2-3.5 mmol/L).

Determining sperm count, motility and morphology

The concentration of sperm cells was determined in a Makler chamber. If the sperm count was lower than 15x10⁶/mL (oligozoospermia), or if there was complete absence of sperm cells (azoospermia), it indicated infertility. A small drop of the sample was stirred up with a glass rod, put in the center of the chamber and covered. The cells were counted under a magnification of X20 in 10 squares, and the sum corresponded to the concentration measured in mmol/L (Fig. 1).

Sperm motility in a sample was determined under magnification of X400 on a Nikon YS 2-t (168159) phase contrast microscope (Nikon, Japan). Sperm cell morphology was determined by the Pap test. The whole sperm cell was observed, its shape and size of its head, neck and tail, and the relationship between them. The method allows coloring the acrosomal and postacrosomal parts of the head and tail, as well as distinguishing the basophilic and acidophilic components, thus enabling good look at nuclear chromatin. The

share of morphologically normal sperm cells cannot be less than 4%. If there are no sperm cells found in the sample, the whole sample is centrifuged and the sediment is examined to determine cryptozoospermia or azoospermia.

Statistics

Categorical variables were expressed as absolute and relative frequencies. Numerical data were described using arithmetic mean and standard deviation for normal data distribution, and median and interquartile range for the remaining cases. Differences in categorical variables were tested using the χ^2 -test and the Fisher exact test if necessary. The Kolmogorov-Smirnov test was used to test normal data distribution. Differences in normally distributed numerical variables between two independent groups were tested using Student's t-test, and by Mann-Whitney U test in case of deviation from normal distribution. All p values were two-tailed. The significance of differences determined by statistical testing was expressed at the level p<0.05. The statistical software package Statistica for Windows 2010 (version 10.0, StatSoft Inc., Tulsa, OK, USA) was used for data analysis¹⁷.

Results

The research was conducted on 276 men, out of which 176 (63.8%) had normal spermiogram, and 100 (36.2%) had been diagnosed with low sperm count. The subjects with low sperm count subfertility were significantly older (median 35, interquartile range 31-38.25) compared to those with normal spermiogram (Mann Whitney U test; p=0.013). In the former, me-

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Table 1. Median age, sperm count,	mobility, morpholog	y and zinc level i	in seminal fluid
of study subjects			

	Median [Q1, Q3]		
	Control group	Low sperm count n=100	p*
Age (years)	32 (30-37.25)	35 (31-38.25)	0.013
Sperm count >15x10 ⁶ /mL	117 (59.75-173)	9 (0-19.25)	<0.001
Motility >40%	63 (58-67)	30 (0-40)	<0.001
Morphology >4%	39 (34-46)	26.5 (0-32)	<0.001
Zinc 1.2-3.5 mmol/L	1.8 (1.3-2.53)	1.55 (0.8-2.1)	0.010

Median [Q1, Q3] interquartile range; *Mann Whitney U test

Table 2. Zinc levels in study subjects

Zinc level	Number of participants (%)			
1.2-3.5 mmol/L	Control group	Low sperm count	Total	p*
1.2 3.5 IIIIIO/ L	n=176	n=100		
Lower than reference interval <1.2	32 (18.2)	38 (38.0)	70 (25.4)	
Reference interval 1.2-3.5	132 (75.0)	59 (59.0)	191 (69.2)	<0.001
Higher than reference interval	12 (6.8)	3 (3.0)	15 (5.4)	

*χ²-test

Table 3. Median age and zinc level in the group of men diagnosed with low sperm count

	Mediar		
	Azoospermia	Low sperm count	p*
	n=26	n=74	
Age	34 (28.75-39.5)	35 (31.25-38)	0.880
Zinc	1.1 (0.65-1.6)	1.75 (0.9-2.2)	0.036

Median [Q1, Q3] interquartile range; *Mann Whitney U test

dian sperm count was 9 (interquartile range 0-19.25), motility 30% (interquartile range 0-40), morphology 26.5% (interquartile range 0-32), all significantly lower compared to men with normal spermiogram analysis (Mann Whitney U test; p<0.001). Zinc values in the sperm samples of infertile men had a significantly lower median of 1.55 (interquartile range 0.8-2.1) (Mann Whitney U test; p=0.010) (Table 1).

After examining the results, the participants were put into 3 groups: lower and higher than the reference interval and the ones in the reference interval. The zinc values lower than the reference were found in 70 (25.4%) of the participants, in the reference interval 191 (69.2%), and higher than the reference in 15 (5.4%). Significant-

ly more men, 42 (35.9%), had zinc values lower than the reference ones (χ^2 test, p<0.001) (Table 2).

In the group of infertile men, those with low sperm count had significantly higher median zinc level of 1.75 (interquartile rage 0.9-2.2) compared to those with azoospermia (Mann Whitney U test; p=0.036) (Table 3).

Discussion

This research confirmed the correlation between zinc level and infertility in men diagnosed with infertility in the sample population.

Zinc is an essential oligoelement necessary for a great number of metabolic processes of over 300 metalloenzymes and more than 500 regulatory protein genes, and is involved in many enzyme reactions. Zinc concentration is higher in the male reproductive organs and semen compared with other bodily fluids and tissues, thus it affects sperm quality¹¹. Zinc value in the sperm of the low sperm count men was 1.55 mmol/L, which was much lower compared to the values recorded in the control group (1.8 mmol/L). Other studies have reported similar results that point to lower zinc

levels in the sperm of infertile men^{13,15,16,18}. When looking at the study results, it should be taken in consideration that the observed population did not include patients with chronic illnesses, smokers, alcoholics, and those on any medication or vitamin/mineral therapy because of the possible effects on sperm quality. The environmental factors, diet, lifestyle, exposure to heavy metals, radiation and high temperature were not considered either, since several other studies had already associated them to male infertility⁵. In the present study, the low sperm count men were older than 35, compared with the men with normal spermiogram analysis, who were aged 32 years. Little is still known about the effect of age on infertility; however, a recent study in France done on 8515 couples showed that men over 35 had 50% less chances of parenthood compared to 25-year-olds¹⁹. The reason for this is the longer period of exposure to risk factors that may lead to mutations in the spermatocyte DNA, such as oxidative stress to which men were exposed during their lifetime²⁰. Accumulation of toxic heavy metals (lead, mercury, cadmium) in testicular tissue can also affect male fertility as does the low zinc level in sperm cells²¹. Researchers from the University of Newcastle discovered three times more DNA damages in the spermatocytes of men aged 36-57 than in those younger than 35²². Lifestyle also affects male reproductive health, as well as the global decline in human sperm quality over the past decades. Factors associated with male infertility are smoking, alcohol, drug use, stress, eating habits, as well as excessive drinking of coffee⁵ and obesity that has an epidemic character worldwide, with a 28.8% to 36.9% increase between 1980 and 20133. In addition, other factors such as exposure of testes to high heat, intense exercise, lack of sleep and exposure to electromagnetic radiation have also been related to infertility⁵. The median sperm count found in the infertile men was 9x106, motility 30% and morphology 26.5%, yielding a statistically enormous difference compared to the control group. Zhao and Xiong proved that there was a connection between low zinc levels in the sperm of infertile men and all the parameters listed above²³.

According to their zinc levels, study subjects were divided into 3 groups, as follows: lower and higher than the reference interval, and those within the reference interval. Seventy (25.4%) subjects had zinc levels lower than the reference interval, 191 (69.2%) were

within the reference interval, and 15 (5.4%) had zinc levels higher than the reference interval. More than 38 (38%) subjects diagnosed with low sperm count had zinc levels lower than the reference interval, which was consistent with the above-mentioned research. Moreover, 59 (59%) subjects had zinc levels within the reference interval, whereas only three (3%) subjects had zinc levels higher than the reference interval. The lower zinc levels in the control group may point to inflammation and prostatic changes²⁴, but this may have been overlooked because health status of the patients was not examined during the research²⁵. A few researches have reported that oral zinc supplements improve sperm cell count, mobility and morphology in men with idiopathic infertility, and to some extent also in fertile men diagnosed with asthenozoospermia and/or oligozoospermia^{25,26}. In the group of men diagnosed with low sperm count, those who had azoospermia had significantly lower zinc levels (1.1 mmol/L) compared with the rest of the group (1.75 mmol/L). These results were consistent with other recent researches²⁷, but in contrast with older researches stating that there was no difference in zinc levels between azoospermia patients and those with asthenozoospermia or oligozoospermia²⁸. Azoospermia is found in 1% of the general population and in 10%-15% of infertile men²⁹. In the Osijek-Baranja County, its prevalence is 9.4%, which is consistent with the data from the above-mentioned literature.

Conclusion

The results of the present research point to the importance of zinc in diagnosing male infertility and may help in explaining the etiopathogenesis of the reproductive function of men, thus assisting in therapy and prevention of the disorder in our population. Dietary habits of the subjects differ regionally and future research could provide solid foundation for an additional approach in treating male infertility. Given the fact that this research included men from the Osijek-Baranja County, it may assist in future researches on this subject.

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Sažetak

ZNAČENJE KONCENTRACIJE CINKA U SJEMENOJ TEKUĆINI KOD MUŠKARACA S DIJAGNOZOM NEPLODNOSTI

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Muška neplodnost s obzirom na njenu učestalost i važnost predstavlja ozbiljan globalni medicinski i socijalni problem. Cilj ovoga istraživanja bio je utvrditi postoje li značajne razlike koncentracije cinka u sjemenoj tekućini muškaraca s dijagnozom smanjene plodnosti u odnosu na kontrolnu skupinu te ako postoje, utvrditi kako razina cinka djeluje na broj, postotak pokretnih i broj morfološki normalnih spermija. Studija slučajeva i kontrola provedena je na području Osječko-baranjske županije u razdoblju od siječnja 2014. godine do lipnja 2015. godine. U studiju je bilo uključeno 276 ispitanika koji su bili upućeni u biokemijski laboratorij KBC-a Osijek, od kojih je kod 100 (36,2%) ispitanika potvrđena dijagnoza smanjene plodnosti ponovljenim spermogramom nakon 3 mjeseca. Uzorci sjemene tekućine obrađeni su u skladu s kriterijima Svjetske zdravstvene organizacije, a koncentracija cinka određena je spektrofotometrijskom, direktnom kolorimetrijskom metodom bez deproteinizacije na kemijskom analizatoru tipa Olympus AU 680 (Beckman Coulter, Tokyo, Japan). Istraživanje je pokazalo kako su muškarci s dijagnozom smanjene plodnosti značajno stariji (Mann Whitney U test: p=0,013) i imaju značajno niže vrijednosti cinka (γ^2 -test: p<0,001). Vrijednosti cinka bile su značajno više u skupini ispitanika s dijagnozom smanjene plodnosti kod kojih su nađeni spermiji u odnosu na ispitanike kojima je utvrđena azoospermija (Mann Whitneyev U test: p=0,036). Koncentracija cinka kao značajnog biološkog biljega i antioksidansa utječe na broj, pokretljivost i morfologiju spermija. Nadomjesna terapija cinkom mogla bi poboljšati sjemene parametre kod bolesnika s dijagnozom smanjene plodnosti. Prehrambene navike ispitanika regionalno se razlikuju, a buduća istraživanja mogla bi postaviti dobre temelje za komplementarno liječenje muške neplodnosti.

Ključne riječi: Hrvatska; Neplodnost muškarca; Cink; Studija slučajeva i kontrola; Oligospermija; Sperma