

## Original Research Article

# Role of Micronutrients in Male Infertile Patient having Abnormal Semen Parameter

### ABSTRACT

**Background:** Abnormal semen parameter is the most common cause of infertility in men. Approximately 15% of human couples are infertile and male sub fertility accounts for 50% of cases. About 90% of male factor infertility is idiopathic with no identifiable cause. Affected males has a definite detectable abnormality related to infertility, such as endocrine 1-3% disease, antisperm antibodies 3-13%, varicocele 25.4%, genetic cause 10-15%. A large number of recent studies have focused on the ability of nutraceuticals, to improve the hormonal status and sperm parameters by different mechanisms. Considering the positive effects of micronutrients on sperm motility and count a mixture of micronutrients applied to reverse the sperm parameters.

**Objectives:** To assess the efficacy of micronutrients regarding improvement of sperm count and motility with abnormal semen parameter.

**Methods:** It was a prospective observational study. This study was performed in Reproductive Endocrinology and Infertility Unit of Dhaka Medical College Hospital between 06 October, 2019 to 05 October, 2020 (01 year). Study population consist of all the diagnosed case of infertile males having abnormal semen parameter between 25 to 50 years of age at DMCH. A total of 100 infertile males with abnormal semen parameter were selected for this study but finally 70 infertile male were included. A full assessment includes demographic information and baseline semen analysis. Then anti-oxidant was given one tablet in the morning and another at evening for three months and follow up semen analysis was done at prefixed schedule after three months to analysis the changes that was achieved. Then pre-treatment and post treatment semen parameters, including sperm count and sperm motility were assessed. Statistical analyses were carried out by using the statistical package for social sciences version 22.0 for windows (SPSS Ins, Chicago, Illinois, USA).

**Main outcome measure(s):** Changes in sperm count and motility.

**Results:** The mean age was  $40.5 \pm 5.9$  years with ranged from 25 to 50 years. The Mean sperm count was found  $12.34 \pm 1.84$  mill/mL pre-treatment and  $15.50 \pm 4.69$  mill/mL post-treatment which was statistically significant ( $p < 0.05$ ). The mean rapid progression was found  $28.84 \pm 20.77$  pre-treatment and  $47.44 \pm 25.02$  post-treatment which was statistically significant ( $p < 0.05$ ). The mean non-progressive sperm was found  $40.34 \pm 21.12$  percent before treatment and  $28.32 \pm 15.47$  percent after treatment which was significantly decreased after micronutrient therapy ( $p < 0.05$ ). The improvement of semen parameters after treatment was 72.86%.

**Conclusion:** Majority of the male patients suffer from infertility due to Oligozoospermia and asthenozoospermia. Increasing age, smoking and industrial workers were more common. Sperm count, sperm motility and rapid progression had significantly improved after three months of treatment. Therefore, this study suggests that micronutrients can be helpful to improve the sperm count and motility of infertile male having abnormal semen parameter.

**Keywords:** Oligozoospermia, asthenozoospermia, Infertility, micronutrients, antioxidant.

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### INTRODUCTION

The concept of male subfertility has evolved rapidly since 2000 [1]. About 90% of male factor infertility is idiopathic with no identifiable cause. Only a minority of affected males have a definite detectable abnormality related to infertility. Associated factors are endocrine 1-3% disease, anti-sperm antibodies 3-13%, varicocele 25.4%, and genetic cause 10-15% [2]. Approximately 15% of human couples are infertile and it is generally recognized that male sub-fertility plays a contributory role in up to 50% of cases [3, 4]. Male subfertility falls into two categories either, productive or obstructive. Oligozoospermia is a productive cause of male infertility [3]. The World Health Organization (WHO) classified oligozoospermia, as "sperm count below 15 million per millilitre (ml) of semen as average" and asthenozoospermia as "sperm motility rapid progression  $< 32$  or rapid progression +non-progressive  $< 40$ ". Reproductive organs are highly susceptible to free radicals or oxidative damage from environmental toxins like pesticides, insecticides and heavy metals [2]. A large number of studies have focused on the ability of many

substances, generally termed nutraceuticals, to improve the hormonal status and sperm parameters by different mechanisms. These studies found that micronutrients play an important role in improving abnormal semen parameters [5]. Deterioration of semen parameters may be due to exposure to the environmental toxicant which has detrimental effects on reproductive hormones, spermatogenesis or sperm function. The most widely studied evidence of environmental factors is associated with occupational pollutants, changes in lifestyle, exposure to toxic agents and changes in dietary habits [6,7]. All India Institute of Medical Sciences reported that the majority of men, who were exposed to high temperatures at their workplaces such as welders, dyers, blast furnace workers and those employed in cement and steel factories were more prone to infertility. This is due to excessive environmental heat which increases the temperature of the scrotum, causing a negative effect on sperm production [8]. Lifestyle plays an important role in poor semen quality [5]. Cigarette smoke contains well-known somatic cell mutagens and potent carcinogens. The exact mechanism by which sperm damage occurs is not clear but may be related to an increase in the vulnerability of spermatozoa to oxidative stress given these cells [9,10]. Ageing is an important factor responsible for the decline in semen quality, as sperm concentration decreases with increased age [11]. A balance nutritional supplement with anti-oxidant content can help reverse some of the oxidative damage from environmental toxins and natural ageing. Different micronutrients like vitamin C, vitamin B12, vitamin E, arginine, carnitine, zinc and selenium have specific roles in increasing sperm count and improving function [12-15]. Results of supplementation with folic acid and zinc are particularly relevant because they appear to work synergistically and results in a >70% increase in sperm concentration when taken under study conditions [16]. Selenium supplementation has been shown to yield a significant dose-dependent increase in total sperm count after 26 weeks, although toxicity was reported at excessive doses [17]. When vitamin C intake increases its concentration in seminal plasma prohibits DNA damage [18]. Vitamin E is a fat-soluble antioxidant that neutralizes free radicals and protects cellular membranes against O<sub>2</sub> free radicals. It also prevents lipid peroxidation and therefore improves the functions of other antioxidants [19]. Vitamin E also inhibits the production of ROS in infertile males [20]. L-carnitine (LC) or 3-aminobutyric acid is a naturally occurring compound and also a semi-essential vitamin-like substance required for human metabolism. Findings show a positive relationship between initial sperm movement and increased LC in epididymis and L-acetyl in sperm [21, 22]. CoQ10 also known as ubiquinone is an antioxidant. After 6 months of therapy, CoQ10 increased in the semen of patients who received CoQ10, and sperm motility was improved in these individuals [23]. Zinc supplementation normally protects the spermatozoa against bacteria and also prevents damage to chromosomes [24]. Zinc plays an important role in testicular development and sperm maturation [25]. On the basis of published scientific literature, individual micronutrient has a positive role in male infertility with abnormal semen parameter. Considering the individual benefit of different micronutrients, a blend of micronutrients was used in this study to reverse sperm motility and count in case of abnormal semen parameters. The aim of the study was to assess the efficacy of micronutrients regarding improvement of sperm count and motility with abnormal semen parameter.

## **METHODOLOGY & MATERIALS**

This is a prospective observational study. A total of 70 patients were enrolled and analyzed in this study. The study was conducted at the "Reproductive Endocrinology and Infertility Unit", Dhaka Medical College Hospital (DMCH). The study period was 1 year from 06 October 2019 to 05 October 2020. Male Infertile patients having abnormal semen parameters (oligozoospermia and asthenozoospermia) attending the "Reproductive Endocrinology and Infertility Unit" of DMCH Purposive sampling. The sample was selected by fulfilling the inclusion and exclusion criteria.

### **Inclusion criteria:**

- 25-50 years of age oligozoospermic infertile man whose sperm count (is between 10 to 15 million/ml).
- 25-50 years of age asthenozoospermic infertile man whose sperm motility:
- Rapid progression <32.
- Rapid progression and non-progressive <40.
- Infertile male with abnormal semen parameters with a normal endocrine profile.

### **Exclusion criteria:**

- Medical co-morbidity DM, chronic hypertension, epilepsy, chronic depressive disorder.
- History of any endocrinological disorders that affect semen parameters such as hypothyroidism and hyperprolactinemia.

- Present or past history of drug addiction, smoking and alcoholism.
- History of pelvic surgery–varicocelectomy.
- Present and past history of genital tuberculosis and mumps orchitis.
- Present and past history of taking drugs affecting the semen parameter.
- Present and past history of tumour and trauma.
- Testicular failure with high Follicular Stimulating Hormone.

Infertile males with abnormal semen parameters attending the study centre, DMCH during the study period were included as the study population. Data were collected chronologically by interview, physical examinations and laboratory investigations using a structured questionnaire containing all the variables of interest according to the inclusion and exclusion criteria. A full assessment includes demographic information, clinical presentation, smoking habits, and medical, surgical, and drug histories with a physical examination (height, weight, blood pressure) done. Semen analysis reports (two times at three months intervals), oral glucose tolerance test (OGTT), serum Follicular stimulating hormone, serum Luteinizing hormone, serum Prolactin and serum testosterone were done. All hormonal tests were done at the Institute of Nuclear Medicine, Dhaka medical college hospital. Hormonal tests were done to exclude endocrinological factors related to infertility. For semen analysis, each man provided a semen sample by masturbation into a wide-mounted sterile plastic container labelled with name and ID, in a room close to the laboratory the period of abstinence (3-5 days) was recorded and the semen sample was analyzed by Neubauer counting chamber according to World Health Organization guideline (2010). All semen analysis was done under a controlled condition in one lab by the same Embryologist. The analysis was performed before treatment and three months after starting the first dose.

Statistical analyses were carried out by using the Statistical Package for Social Sciences version 23.0 for Windows (SPSS Inc., Chicago, Illinois, USA). The mean values were calculated for continuous variables. The quantitative observations were indicated by frequencies and percentages. Paired t-test was used for continuous variables. P values <0.05 was considered statistically significant.

## RESULT

This is a prospective observational study which was carried out in the Reproductive Endocrinology and Infertility Unit, Department of Gynecology and Obstetrics, Dhaka Medical College Hospital, Dhaka, between 06 October 2019 to 05 October 2020. A total of 70 infertile males with abnormal semen parameter patients were included in this study maintaining inclusion criteria. Table 1 shows that the majority (40.0%) of patients belonged to age 41-45 years. The mean age was found  $40.5 \pm 5.9$  years with a range from 25 to 50 years. More than one-third (34.3%) patients were industrial workers, and 23(45.7%) patients had monthly family income <10000 BDT. Mean sperm count was found  $12.34 \pm 1.84$  mill/mL before treatment and  $15.50 \pm 4.69$  mill/mL after treatment. Sperm counts were significantly increased after micronutrient therapy ( $p < 0.05$ ) (Table 2). The mean rapid progression was found  $28.84 \pm 20.77$  before treatment and  $47.44 \pm 25.02$  after treatment. Rapid progression was significantly increased after micronutrient therapy ( $p < 0.05$ ) (Table 3). The mean non-progressive sperm was found  $40.34 \pm 21.12$  before treatment and  $28.32 \pm 15.47$  after treatment. Non-progressive sperm was significantly decreased after micronutrient therapy ( $p < 0.05$ ) (Table 4). The mean Immotile was found  $17.64 \pm 16.18\%$  before treatment and  $16.67 \pm 16.22\%$  after treatment. Immotile were decreased after micronutrient therapy but were not statistically significant ( $p > 0.05$ ) (Table 5). Figure 1 shows the improvement of semen parameters after treatment, there was 72.86% improvement.

**Table 1:** Distribution of the study patients by socio-demographic characteristics (N=70).

Socio-demographic characteristics	Frequency	Percentage
Age groups (years)		
25-30	6	8.6
31-35	8	11.4
36-40	13	18.6
41-45	28	40
46-50	15	21.4
Mean $\pm$ SD	$40.5 \pm 5.9$	
Occupational status		
Industrial worker	24	34.3

Private service	17	24.3
Farmer	11	15.7
Gov. service	8	11.4
Abroad	6	8.6
Other	4	5.7
Family income (Taka)		
<10000	32	45.7
10000-30000	24	34.3
>30000	14	20

**Table 2:** Mean sperm count difference between pre-treatment and post-treatment.

Variables	Micronutrients are taken		p-value
	Pre Treatment (n=70)	Post Treatment (n=70)	
	Mean±SD	Mean±SD	
Sperm count (mill/mL)	12.34±1.84	15.50±4.69	0.000 <sup>s</sup>
Range	10.0-15.0	5.0-25.0	

**Table 3:** Mean rapid progression difference between pre-treatment and post-treatment.

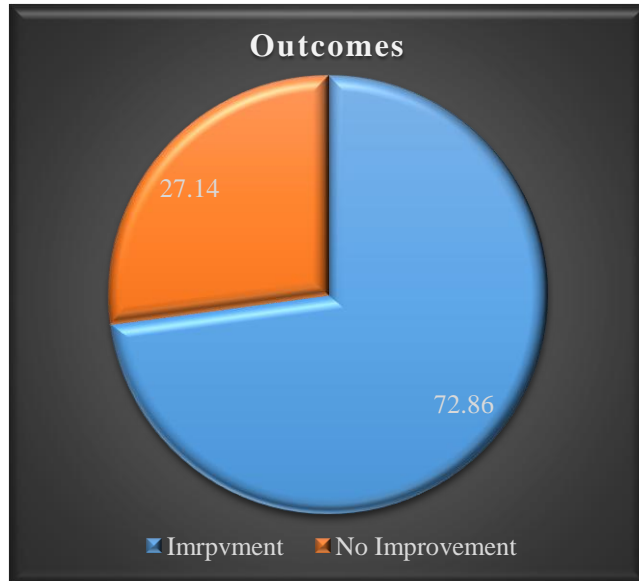
Variables	Micronutrients are taken		p-value
	Pre Treatment (n=70)	Post Treatment (n=70)	
	Mean±SD	Mean±SD	
Rapid progression	28.84±20.77	47.44±25.02	0.000 <sup>s</sup>
Range	(2-90)	(2-90)	

**Table 4:** Mean non-progressive sperm difference between pre-treatment and post-treatment.

Variables	Micronutrients are taken		p-value
	Pre Treatment (n=70)	Post Treatment (n=70)	
	Mean±SD	Mean±SD	
Non-progressive sperm	40.34±21.12	28.32±15.47	0.03 <sup>s</sup>
Range	(8-85)	(8-62)	

**Table 5:** Mean non-progressive sperm difference between pre-treatment and post-treatment.

Variables	Micronutrients are taken		p-value
	Pre Treatment (n=70)	Post Treatment (n=70)	
	Mean±SD	Mean±SD	
Immotile	17.64±16.18	16.67±16.22	0.619 <sup>ns</sup>
Range	(2-80)		



**Figure 1:** Outcome of the treatment of the study respondents (N=70).

## DISCUSSION

This study observed that the majority of the patients (40.0%) belonged to age 41-45 years. The mean age was found  $40.5 \pm 5.9$  years with a range from 25 to 50 years. More than one-third (34.3%) of patients were industrial workers, and 23 (45.7%) patients had monthly family income <10000 BDT. Begum et al also reported that the mean age was found  $34.96 \pm 3.6$  years. Imhof et al. also reported the mean age of men taking the active compound was 34 years (min/max: 18-43 years) [26]. This study showed that the mean sperm count was found  $12.34 \pm 1.84$  million/mL before treatment and  $15.50 \pm 4.69$  million/mL after treatment. The mean rapid progression was found  $28.84 \pm 20.77$  before treatment and  $47.44 \pm 25.02$  after treatment. The mean non-progressive sperm was found  $40.34 \pm 21.12$  before treatment and  $28.32 \pm 15.47$  after treatment. Sperm count and rapid progression were significantly increased after micronutrient therapy ( $p < 0.05$ ) but non-progressive sperm was significantly decreased after micronutrient therapy ( $p < 0.05$ ). Begum et al reported mean count was found  $15.67 \pm 15.15$  million/mL in pre-treatment and  $32.63 \pm 21.18$  million/mL in post-treatment. Mean total motility was found  $39.65 \pm 22.34$  in pre-treatment and  $60.30 \pm 11.29$  in post-treatment. After treatment, there was a significant improvement both in count and motility [2]. Lenzi et al evaluated the effects of a therapeutic formulation, Proxeed Plus, on sperm parameters in oligo asthenozoospermia men. This prospective, randomized, double-blind, placebo-controlled clinical trial involved 175 males (19-44 years) with idiopathic oligoasthenozoospermia. Males received Proxeed Plus or a placebo for 3 and 6 months. Sperm volume, progressive motility and vitality significantly ( $p < 0.001$ ) improved after 6 months compared to baseline [27]. Imhof et al conducted a comparative pilot study at the fertility center IMI, Vienna, Austria. A total of 132 sub-fertile males (active treatment group) took two capsules daily of the active component for a three months period between the first and follow-up semen analysis. Sub-fertile men received no active treatment and served as control. A main outcome measure was the standardized semen analysis [26]. All parameters evaluated by semen analysis significantly increased after three months of treatment with the active compounds. Median ejaculatory volume, sperm cell density, sperm motility (progressive and total) and normal morphology increased by 33.3%, 215.5%, 83.1%, 36.4% and 23.0% respectively. These increments were significantly higher than those observed among controls. A total of 34 pregnancies were reported after six months of follow-up whereas eleven were reported control group [26]. PM Gopinath et al conducted a Placebo-controlled, Double-blind, randomized, Parallel three-arm, Multicentric trial. Compared to placebo, a statistically significant improvement was seen in sperm count ( $14.8$ - $26.35$  in arm 1 and  $14.37$ - $24.8$  million/ml in arm 2,  $p < 0.0001$ ), and sperm total motility ( $39.2$ - $51.6\%$  in arm 1 and  $38.4$ - $50.1\%$  in arm 2,  $p < 0.0001$ ), at 90 days, and treatment further improved these parameters at day 180. No intergroup difference was seen between arm 1 and arm 2. 36 Mahammad K. Moslemi et al included 690 infertile men with idiopathic asthenoteratospermia who received supplemental daily Se ( $200 \mu\text{g}$ ) in combination with vitamin E (400 units) for at least 100 days. They observed 52.6% (362 cases) total improvement in sperm motility, morphology, or both, and 10.8% (75 cases) spontaneous pregnancy in comparison with no treatment (95% confidence interval: 3.08 to 5.52). No response to treatment occurred in 253 cases (36.6%) after 14 weeks of combination therapy. The mean

difference between semen analyses of cases before and after treatment was 4.3% with a standard deviation of 4.29. On the basis of paired t-test results, combination therapy with oral Se and vitamin E was effective for the treatment of oligozoospermia or asthenozoospermia or induction of spontaneous pregnancy ( $P \leq 0.001$ ) [37]. Orv Hetil et al conducted a clinical trial with 100 males with low sperm quality (sperm count 5-20 million/mL, motility 10-40%, and abnormal morphology 30-50) where he found that dietary supplements statistically and clinically significantly improved sperm count and motility. In 74 cases this dietary supplement demonstrated a beneficial effect on sperm quality (more than 10% increase in sperm count, or quality of motility, or shape); in 16 cases the improvement exceeded 30%. No adverse effect could be accounted for by this treatment [28]. A systemic review of randomized studies was conducted to evaluate the effects of oral antioxidants on sperm quality and pregnancy rate in infertile men. Despite the methodological and clinical heterogeneity, 14 of the 17(82%) trials showed an improvement in either sperm quality or pregnancy rate after antioxidant therapy [29]. In 2008, Paradiso Galatioto et al used a multi-drug therapy including several antioxidants (consisting of daily administration of vitamin A 0.06 IU, vitamin C 3 mg, vitamin E 0.2 mg, N-acetyl-cysteine (NAC) 10 mg, zinc 0.01 mg, thiamine 0.4 mg, riboflavin 0.1 mg, pyridoxin 0.2 mg, nicotinamide 1 mg, pantothenate 0.2 mg, biotin 0.04 mg, cyanocobalamin 0.1 mg, calciferol 8 IU, calcium 1 mg, magnesium 0.35 mg, phosphate 0.45 mg, iron 0.2 mg, manganese 0.01 mg, copper 0.02 mg) in 42 oligozoospermic subjects: the treatment group had a 20-fold higher chance of having a normal sperm count than untreated men, and a non-significant increase in the chance of achieving pregnancy [30]. An observational study was conducted to evaluate the effect of multiple micronutrient therapy in sub-fertile males. This study included 103 sub-fertile men. Here 42 patients received 2 tablets of oligocare daily and 61 received 2 tablets with other concomitant therapy. The result of this study suggested that multiple micronutrient monotherapy is equally effective in the management of sub-fertility or idiopathic oligozoospermia in male patients as compared to combination therapy along with multiple micronutrients [31]. The present study showed that for 51(72.86%) patients, there were improvements in semen parameters and for 19(27.14%) there were no improvements in semen parameters. Begum et al. observed 69.55% of patients showed improvement after treatment in terms of count and motility [2]. All studies which examine the effect of multiple antioxidants in a supplementation showed an improvement in semen parameters after therapy [30,32-36]. Above mentioned study's findings are consistent with this study. One study showed a significant improvement in sperm concentration after combination therapy without improvement in motility and morphology [37]. Henkel et al found that the risk of consuming excessive dosages of micronutrient supplements, which may be toxic and results in a phenomenon termed an antioxidant paradox. It may lead to reductive stress, which is reported to be as dangerous to cells as oxidative stress and can be the cause of diseases such as cancer or cardiomyopathy. Therefore, there is a need for more elaborate research to establish the clear benefits and risks involved in antioxidant therapy for male infertility [38].

**Limitations of the study:** The present study had some limitations. The following should be kept in mind while deciding on the implications of the findings of the study: The study population was selected from one selected tertiary hospital in Dhaka city. So, the results of the study may not reflect the exact picture of the entire community. The small sample size was also a limitation of the study. The study was conducted for a short period of time.

### CONCLUSION AND RECOMMENDATIONS

This study was undertaken to evaluate the effect of micronutrients on sperm count and motility in infertile males with abnormal semen parameters. Sperm count and motility had significantly improved after three months of treatment with a micronutrient supplement. Therefore, this study suggested that micronutrients can be helpful to improve sperm count and motility. Improvement of seminal quality may improve the fertility outcome of an infertile couple. Even though a recent study concluded the positive effect of sperm parameters, there is a need for further investigation with randomized control studies to confirm the efficacy and safety of antioxidant supplementation. There is also a need to determine the ideal dose of each compound to improve semen parameters, fertilization rates and pregnancy outcomes. Larger trials with high-quality controls and randomization must be performed to establish clinically relevant guidelines for supplementations.

**Ethical approval:** The study was approved by the Institutional Ethics Committee.

### REFERENCES

1. Salma U, Gill HK, Keith LG, Tilmon S, Jones CA, Sobti A, Patel A. Male subfertility and the role of micronutrient supplementation: clinical and economic issues. *Journal of experimental & clinical assisted reproduction*. 2011;8.
2. Begum MR, Miller D, Salam MA, Quadir E, Begum MS, Khan F, Bhuiyan ZH. Antibiotics and Micronutritional Blend to Enhance Fertility Potential in Male Having Abnormal Semen Parameters. *The Open Clinical Trials Journal*. 2009 Dec 16;1(1).
3. Isidori AM, Pozza C, Gianfrilli D, Isidori A. Medical treatment to improve sperm quality. *Reproductive biomedicine online*. 2006 Jan 1;12(6):704-14.
4. Williams EA, Parker M, Robinson A, Pitt S, Pacey AA. A randomized placebo-controlled trial to investigate the effect of lactolycopene on semen quality in healthy males. *European Journal of Nutrition*. 2020 Mar;59:825-33.
5. Garolla A, Petre GC, Francini-Pesenti F, De Toni L, Vitagliano A, Di Nisio A, Foresta C. Dietary supplements for male infertility: a critical evaluation of their composition. *Nutrients*. 2020 May 19;12(5):1472.
6. Chavarro JE, Toth TL, Sadio SM, Hauser R. Soy food and isoflavone intake in relation to semen quality parameters among men from an infertility clinic. *Human reproduction*. 2008 Nov 1;23(11):2584-90.
7. Tielemans E, Burdorf A, te Velde ER, Weber RF, van Kooij RJ, Veulemans H, Heederik DJ. Occupationally related exposures and reduced semen quality: a case-control study. *Fertility and sterility*. 1999 Apr 1;71(4):690-6.
8. Ajayi VD, Ajayi AB, Ramesh B, Afolabi BM, Biobaku O, Oyetunji I. Comparative Analysis of Hysteroscopic Findings among infertile Women in Nigeria and in India: a Preliminary Investigation. *J Gynecol Women's Health*. 2017;6(2):555681.
9. Sikka SC. Relative impact of oxidative stress on male reproductive function. *Current medicinal chemistry*. 2001 Jun 1;8(7):851-62.
10. ARMSTRONG JS, RAJASEKARAN M, HELLSTROM WJ, SIKKA SC. Antioxidant potential of human serum albumin: role in the recovery of high quality human spermatozoa for assisted reproductive technology. *Journal of Andrology*. 1998 Jul 8;19(4):412-9.
11. Sasano N, Ichijo S. Vascular patterns of the human testis with special reference to its senile changes. *The Tohoku journal of experimental medicine*. 1969;99(3):269-80.
12. Goa KL, Brogden RN. L-Carnitine: a preliminary review of its pharmacokinetics, and its therapeutic use in ischaemic cardiac disease and primary and secondary carnitine deficiencies in relationship to its role in fatty acid metabolism. *Drugs*. 1987 Jul;34:1-24.
13. Prasad AS. Zinc in growth and development and spectrum of human zinc deficiency. *Journal of the American College of Nutrition*. 1988 Oct 1;7(5):377-84.
14. Ursini F, Heim S, Kiess M, Maiorino M, Roveri A, Wissing J, Flohé L. Dual function of the selenoprotein PHGPx during sperm maturation. *Science*. 1999 Aug 27;285(5432):1393-6.
15. Hansen JC, Deguchi Y. Selenium and fertility in animals and man—a review. *Acta Veterinaria Scandinavica*. 1996 Mar;37:19-30.
16. Wong WY, Merkus HM, Thomas CM, Menkveld R, Zielhuis GA, Steegers-Theunissen RP. Effects of folic acid and zinc sulfate on male factor subfertility: a double-blind, randomized, placebo-controlled trial. *Fertility and sterility*. 2002 Mar 1;77(3):491-8.
17. Agarwal A, Prabakaran S, Allamaneni S. What an andrologist/urologist should know about free radicals and why. *Urology*. 2006 Jan 1;67(1):2-8.
18. Colagar AH, Marzony ET. Ascorbic Acid in human seminal plasma: determination and its relationship to sperm quality. *Journal of clinical biochemistry and nutrition*. 2009;45(2):144-9.
19. Brigelius-Flohé R, Traber MG. Vitamin E: function and metabolism. *The FASEB journal*. 1999 Jul;13(10):1145-55.
20. Ross C, Morriss A, Khairy M, Khalaf Y, Braude P, Coomarasamy A, El-Toukhy T. A systematic review of the effect of oral antioxidants on male infertility. *Reproductive biomedicine online*. 2010 Jun 1;20(6):711-23.
21. Johansen L, Bøhmer T. Carnitine-binding related suppressed oxygen uptake by spermatozoa. *Archives of Andrology*. 1978 Jan 1;1(4):321-4.
22. Radigue C, Es-Slami S, Soufir JC. Relationship of carnitine transport across the epididymis to blood carnitine and androgens in rats. *Archives of Andrology*. 1996 Jan 1;37(1):27-31.

23. Balercia G, Buldreghini E, Vignini A, Tiano L, Paggi F, Amoroso S, Ricciardo-Lamonica G, Boscaro M, Lenzi A, Littarru G. Coenzyme Q10 treatment in infertile men with idiopathic asthenozoospermia: a placebo-controlled, double-blind randomized trial. *Fertility and sterility*. 2009 May 1;91(5):1785-92.
24. Hadwan MH, Almashhedy LA, Als Salman AR. Oral zinc supplementation restore high molecular weight seminal zinc binding protein to normal value in Iraqi infertile men. *BMC urology*. 2012 Dec;12(1):1-6.
25. Elgazar V, Razanov V, Stoltenberg M, Hershinkel M, Huleihel M, Nitzan YB, Lunenfeld E, Sekler I, Silverman WF. Zinc-regulating proteins, ZnT-1, and metallothionein I/II are present in different cell populations in the mouse testis. *Journal of Histochemistry & Cytochemistry*. 2005 Jul;53(7):905-12.
26. Imhof M, Lackner J, Lipovac M, Chedraui P, Riedl C. Improvement of sperm quality after micronutrient supplementation. *e-SPEN Journal*. 2012 Feb 1;7(1):e50-3.
27. Lenzi A, Sgro P, Salacone P, Paoli D, Gilio B, Lombardo F, Santulli M, Agarwal A, Gandini L. A placebo-controlled double-blind randomized trial of the use of combined l-carnitine and l-acetyl-carnitine treatment in men with asthenozoospermia. *Fertility and sterility*. 2004 Jun 1;81(6):1578-84.
28. Horváth M, Czeizel E. Effect of new dietary supplement on sperm quality. *Orvosi Hetilap*. 2012 Nov 1;153(45):1787-92.
29. Lombardo F, Sansone A, Romanelli F, Paoli D, Gandini L, Lenzi A. The role of antioxidant therapy in the treatment of male infertility: an overview. *Asian Journal of andrology*. 2011 Sep;13(5):690.
30. Paradiso Galatioto G, Gravina GL, Angelozzi G, Sacchetti A, Innominato PF, Pace G, Ranieri G, Vicentini C. May antioxidant therapy improve sperm parameters of men with persistent oligospermia after retrograde embolization for varicocele?. *World journal of urology*. 2008 Feb;26:97-102.
31. Tayade H P, Rathod O K, Evaluation of efficacy of multiple micronutrients supplements alone or in combination with concomitant therapy in subfertile male. *Journal of pharmaceutical research and clinical practice*. 2012
32. Micic S, Lalic N, Nale DJ, Bojanic N. Effects of L-carnitine on sperm motility and number in infertile men. *FERTILITY AND STERILITY-INTERNATIONAL EDITION*-. 1998;70:O-030.
33. Kessopoulou E, Powers HJ, Sharma KK, Pearson MJ, Russell JM, Cooke ID, Barratt CL. A double-blind randomized placebo cross-over controlled trial using the antioxidant vitamin E to treat reactive oxygen species associated male infertility. *Fertility and sterility*. 1995 Oct 1;64(4):825-31.
34. Vézina D, Mauffette F, Roberts KD, Bleau G. Selenium-vitamin E supplementation in infertile men: effects on semen parameters and micronutrient levels and distribution. *Biological Trace Element Research*. 1996 Jun;53:65-83.
35. Tremellen K, Miari G, Froiland D, Thompson J. A randomised control trial examining the effect of an antioxidant (Menevit) on pregnancy outcome during IVF-ICSI treatment. *Australian and New Zealand Journal of Obstetrics and Gynaecology*. 2007 Jun;47(3):216-21.
36. Matalliotakis I, Koumantaki Y, Evageliou A, Matalliotakis G, Goumenou A, Koumantakis E. L-carnitine levels in the seminal plasma of fertile and infertile men: correlation with sperm quality. *International journal of fertility and women's medicine*. 2000 May 1;45(3):236-40.
37. Safarinejad MR, Safarinejad S. Efficacy of selenium and/or N-acetyl-cysteine for improving semen parameters in infertile men: a double-blind, placebo controlled, randomized study. *The Journal of urology*. 2009 Feb;181(2):741-51.
38. Henkel R, Sandhu IS, Agarwal A. The excessive use of antioxidant therapy: A possible cause of male infertility?. *Andrologia*. 2019 Feb;51(1):e13162.