Investigation of minerals, testosterone, and transaminases in the semen and serum of fertile and infertile men belongs to different age groups

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ABSTRACT
The present study was aimed to assess the potential of infertility to induce adverse effects with reference to testosterone, Triiodothyronine (T3), Thyroxine (T4), Alanine Aminotransferase (ALT), Aspartate Aminotransferase (AST), zinc, copper and iron. All the samples were divided into four groups according to age and disorder (Group 1, 10 infertile men of 25 - 40 years; Group 2, 10 fertile men of 25 - 40 years; Group 3, 10 infertile men of 41 - 60 years and Group 4, 10 fertile men of 41 - 60 years). Semen and blood samples were analyzed by atomic absorption spectrophotometry to determine minerals while, Testosterone, T3 and T4 were determined by enzyme immunoassay kits. ALT and AST were determined using standard kit assay method. The levels of testosterone and T3 and AST in the fertile semen of 41 - 60 years age group were increased significantly (P ≤ 0.001) as compared to that of fertile semen of 25 - 40 years age group. While, the level of T4 in the fertile semen of 41 - 60 years age group was decreased significantly (P ≤ 0.001) as compared to that of fertile semen of 25 - 40 years age group. In case of fertile serum, only the level of testosterone was significantly decreased (P ≤ 0.05) in the 41 - 60 years age group as compared to 25 - 40 years age group. The levels of testosterone and Cu in the infertile serum of 41 - 60 years age group were decreased significantly (P ≤ 0.001). While, the levels of T3, T4, ALT and Fe in the infertile serum of 41 - 60 years age group were increased significantly (P ≤ 0.05) as compared to that of infertile serum of 25 - 40 years age group.

Keywords: Semen; Reproduction; Infertility; Minerals; Testosterone; Transaminases

1. INTRODUCTION
Infertility primarily refers to the biological inability of a person to contribute to conception. According to World Health Organization [1], infertility is a period of two years without conception. After one year of infertility many couples seek a medical opinion because modern artificial reproduction techniques like intra-cytoplasmic sperm injection (ICSI) can help couples to overcome infertility [2]. Sub fertility is the failure to conceive after 1 year of regular, unprotected intercourse with the same partner. Sub fertility is caused by sperm defects or dysfunction [3].

Testosterone is a principal male sex hormone and is primarily secreted in the testes of males. In men, testosterone plays a crucial role in the development of male reproductive tissues such as the testis and prostate as well as promoting secondary sexual characteristics such as increased muscle, bone mass and the growth of body hair [4]. The thyroid hormones, T3, and T4 are tyrosine based hormones formed by the thyroid gland mainly responsible for regulation of metabolism [5]. ALT and AST are enzymes located in liver cells that leak out into the general circulation during the injury of liver cells [6].

Among the metal trace elements Zinc is one of the most interesting nutritional traces in the reproductive system. In man it is a cofactor of more than 200 metallo-enzymes and plays an important role in the normal testicular development, spermatogenesis, and sperm motility [7]. Copper is an essential trace element required in the diet because it is the metal cofactor for a variety of enzymes (amine oxidase, superoxide dismutase, cytochrome oxidase and tyrosinase). However, excess copper can cause problems because of its capacity to oxidize proteins and lipids, bind to nucleic acids and enhancing the production of free radicals. Therefore, it is important to maintain the amount of copper in the body within normal limits. Copper reduces sperm motility by reduc-
ing oxidative processes and glucose consumption [8]. Iron is a vital component of a group of heme proteins that function in oxygen transport or as enzymes in redox systems. A small amount of iron is enclosed in several non heme metalloenzymes. The chief complexes coordinating iron with the cell are heme and heme containing proteins, hemosiderin and ferritin [9,10]. The aim of this study was to assess the potential causes of infertility and old age to induce the adverse effects with reference to testosterone, T3, T4, ALT, AST, zinc, copper and iron.

2. MATERIALS AND METHODS

2.1. Sample Collection

Semen and blood samples were collected from each individual patient of each group. Semen samples were collected by masturbation in a clean specimen container after sexual abstinence for 3 - 5 days followed by liquefaction at 37°C. The semen samples were evaluated according to World Health Organization recommendations (ejaculate volume, pH, time to liquefaction, sperm concentration, motility and morphology). The remaining semen samples were centrifuged at 3000 × g for 10 min to obtain the seminal plasma. The separated seminal plasma was stored at –80°C until further analysis for the detection of testosterone, Zn, Cu, Fe, T3, T4, ALT and AST level. Blood samples were centrifuged at 4000 × g for 15 min after complete coagulation and supernatant serum was collected in Eppendorf tubes and frozen until further analysis for the detection of testosterone, Zn, Cu, Fe, T3, T4, ALT and AST level.

2.2. Measurement of Minerals

Zinc, iron and copper in the samples were measured by atomic absorption spectrophotometer. The samples were aspirated to the instrument and reading was recorded at the Monochromator wavelength 213.9, 285.2 and 342.7 nm for zinc, iron and copper respectively.

2.3. Measurement of Testosterone

Testosterone was determined with the help of DRG Testosterone Enzyme Immunoassay Kit.

2.4. Measurement of T3 & T4

Triiodothyronine (T3) and Thyroxine (T4) were determined by using J D Biotech T3 and J D Biotech T4 Enzyme Immunoassay Kit respectively.

2.5. Measurement of ALT

Human Gasellschaft fur Biochemica and diagnostica mbH (EC 3.1.3.1) kit was used to determine the level of ALT as described by Saher et al. [11].

2.6. Measurement of AST

The AST was evaluated by adopting the methodology as described by Reitman and Frankel [12].

2.7. Statistical Analysis

The results were expressed as mean ± SEM for all continuous variables. Mean ± SEM have been prepared in Microsoft® Excel Windows Version 2007. The data obtained were analyzed by applying student t-test using Minitab 15 Windows Version to assess the significant differences between all selected age groups.

3. RESULTS AND DISCUSSION

The levels of testosterone and T3 and AST in the fertile semen of 41 - 60 years age group were increased significantly (P ≤ 0.001) as compared to that of fertile semen of 25 - 40 years age group. While, the level of T4 in the fertile semen of 41 - 60 years age group was decreased significantly (P ≤ 0.001) as compared to that of fertile semen of 25 - 40 years age group. There was no significant difference among the levels of ALT, Zn, Cu and Fe between the fertile semen of 25 - 40 years age group and 41 - 60 years age group (Figure 1).

In case of fertile serum, only the level of testosterone was significantly decreased (P ≤ 0.05) in the 41 - 60 years age group as compared to 25 - 40 years age group. While, the levels of T3, T4, ALT, AST, Zn, Cu and Fe had no significant difference between the fertile serums of 25 - 40 years age group and 41 - 60 years age group (Figure 2).

In the infertile semen of 41 - 60 years age group, the levels of testosterone, T4, Zn and Cu were decreased significantly (P ≤ 0.001, P ≤ 0.05) as compared to that of infertile semen of 25 - 40 years age group. While, the levels of T3, ALT and AST were increased significantly (P ≤ 0.001, P ≤ 0.05) in the infertile semen of 41 - 60 years age group as compared to that of infertile semen of 25 - 40 years age group (Figure 3).

The levels of testosterone and Cu in the infertile serum of 41 - 60 years age group were decreased significantly (P ≤ 0.001) as compared to that of fertile semen of 25 - 40 years age group. While, the levels of T3, T4, ALT and Fe in the infertile serum of 41 - 60 years age group were increased significantly (P ≤ 0.001, P ≤ 0.05) as compared to that of infertile serum of 25 - 40 years age group (Figure 4).

4. DISCUSSION

The lower level of testosterone in the fertile serum of 41 - 60 years age group as compared to that of fertile serum of 25 - 40 years age group indicates that testosterone level is decreased in older age resulting in the decreased
Figure 1. Biochemical parameters evaluation in fertile semen of age group 2 & 4.

Figure 2. Biochemical parameters evaluation in fertile serum of age group 2 & 4.
Figure 3. Biochemical parameters evaluation in the infertile semen of age group 1 & 3.

Figure 4. Biochemical parameters evaluation in the infertile serum of age group 1 & 3.
sperm production. In consequence, the testosterone concentration is increased in the fertile semen of 41 - 60 years age group as compared to that of fertile semen of 25 - 40 years age group to increase the sperm production. Significantly lower level of testosterone in the infertile serum of 41 - 60 years age group represents that testosterone deficiency is more pronounced in older age. This may be due to atrophy of glandular tissue of pituitary or testes resulting in significantly lower level of testosterone in the infertile semen of 41 - 60 years age group. This may also be due to prostatic inflammation and obstruction or blockage of the male reproductive tract. Hence testis not functioning properly and testosterone production is affected. Zinc deficiency in infertile semen of 41 - 60 years age group also causes lower testosterone production as zinc is involved in steroidogenesis, testicular development and testosterone synthesis [3]. The similar results were found by Khan et al. [13]. They observed the decrease in testosterone concentration in the infertile individuals. Omrani et al. [14] and Weedin et al. [15] also found the decreased testosterone level in the infertile men.

Significantly lower level of zinc in the infertile semen of 41 - 60 years age group in the present study might be due to excessive ROS production. After ejaculation the abnormal spermatozoa are sources of oxidants which bind with the zinc and reduce its concentration in seminal plasma. Hence, zinc is considered to be an important antioxidant protecting the spermatozoa from oxidative stress. Zinc deficiency decreases testicular weight and causes shrinkage of the seminiferous tubules. Zinc deficiency also induces atrophy of the seminiferous tubules and causes failure of the spermatogenesis. Low seminal zinc levels have been correlated with decreased fertility potential [16]. Ali et al. [17] observed the low zinc level in the infertile subjects and found that low zinc level affected the semen parameters. Colagar et al. [18] found that poor Zn nutrition may be an important risk factor for low quality of sperm and idiopathic male infertility.

Significantly lower level of Copper in the infertile semen and serum of 41 - 60 years age group may be due to liver toxicity as it is absorbed by the liver, bound to albumin and excreted into the bile. It is also involved in protein synthesis during certain enzymatic reactions acting as a cofactor. Nutritional deficiency may also lower its level in infertile men. Abdul-Rasheed [19] reported a significant decrease in seminal plasma copper levels in azoospermic patients. In the present research work, significant increase of iron in the infertile serum might be due to oxidative stress in infertile men. Aydemir et al. [20] observed the increased iron level in the infertile men. Eghbali et al. [10] reported that seminal plasma iron content is related with the motility and viability of the spermatozoa after ejaculation. Presence of iron in the seminal plasma will assist spermatozoa to sustain their functions as long as it is needed for semen fertility.

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