Prevalence of Azoospermia in Infertile Males in Thi-Qar Governorate (Iraq)

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ABSTRACT

Objective: Guy infertility is the inability of a sexually mature male to become pregnant by a fertile female. It causes 40% of infertility in humans. Semen quality is utilized as a proxy indicator of male fecundity since defects in the semen are frequently the cause of male infertility. This study's goal was to find out how common azoospermia was among men in infertile couples in the southern Iraqi city of Thi-Qar.

Methods: The study was carried out at infertility unit in Thi Qar city, province of Iraq from January 2018 to January 2021. This study involved an infertile couple, SFA was done to evaluate male factors. Males with Azoospermia were involved. Regarding male, SFA was examined according to WHO 2010. Information such as age, job and duration of infertility were taken.

Results: Number of couples is 2000, patients with abnormal SFA are 600 (30%). From those patients, 30 with azoospermia (5%). Largest job group of Azoospermic Patients is self-employer, which is significantly different from other groups (P<0.01). Non obstructive type is Significantly different (P<0.01) from obstructive type. Half of the couples only undergone IVF program (in vitro fertilization).

Conclusion: The prevalence of azoospermia in our province (5%). Half of the patient only undergone IVF program (in vitro fertilization) because there is no IVF center in our province in addition to high cost of private centers.

Keywords: Azoosperia, SFA (seminal fluid analysis), Infertility

INTRODUCTION

The secretions from the prostate, seminal vesicles, epidydimis, urethral glands, Cowper's glands, and vasa deferentia make up the seminal fluid, often known as semen, which is a clear or gray liquid¹. Millions of spermatozoa (sperm) are present in every milliliter of semen, however the majority of the volume is made up of glandular secretions from the male reproductive organs². Azoospermia affects about 1% of males overall, while ten% to fifteen % of infertile men have the condition. The absence of any seminal fluid emission after ejaculation distinguishes aspermia from azoospermia^{3,4}. The three primary kinds of azoospermia causes are pretesticular, testicular, and post testicular. Endocrine disorders that negatively affect spermatogenesis (secondary testicular failure) are one of the more uncommon pretesticular causes of azoospermia. One of the main reasons of azoospermia is testicular abnormalities of spermatogenesis that occur internally to the testes (primary testicular failure). A ductal obstruction or ejaculatory malfunction that prevents sperm from reaching the urethral meatus is one of the causes of azoospermia in about 40 percent of affected men⁵. Azoospermia is characterized by the total lack of sperm in the ejaculate. A semen sample must be centrifuged for 15 minutes at room temperature in order to confirm this diagnosis. The pellet must also be examined under a microscope at a high magnification, and the centrifugation speed must be at least 3,000 g. It is advised to adhere to the 2010 World Health Organization criteria for semen analysis and to analyze at least two samples of semen taken more than two weeks apart⁶.

METHODOLOGY

The study was carried out at infertility unit in Thi Qar city, province of Iraq from January 2018 to January 2021. This study involved an infertile couple, 2000 couples. SFA was done to evaluate male factors. 600 hundred with abnormal parameters only 30 males with

Azoospermia. Regarding male, SFA was examined according to WHO 2010. Information such as age, job and duration of infertility were taken and analysed for Azoospermic patients⁷⁻¹³.

Seminal Fluid Collection and Examination

3-5 days of sexual abstinence were followed by masturbation to acquire seminal fluid. Within an hour of ejaculation, the samples were taken in sterile, wide-mouthed, non-toxic containers and processed in the lab. The seminal fluid was assessed using macroscopical (appearance, volume, liquefaction, pH, and viscosity) and microscopical (spermatozoa concentration, morphology, and motility) criteria in line with the WHO (2010) manual of seminal fluid examination⁵.

RESULTS

Figure 1 study the patients, from it we notice that number of patients is 2000. patients with abnormal SFA are 600 (30%). Figure 2 describe the percentage of Azoospermic patients (5%). Figure 3 shows Comparative of age groups of Azoospermic patients, there is nonsignificant differences (P>0.05) between them, but age 22-30 year is the largest age group. Figure 4 explains the Comparative of job groups of Azoospermic patients, there is Significant differences (P<0.01) for self-employer. Figure 5 describes Comparative of Azoospermia family history groups of Azoospermic patients. There are non-significant differences (P>0.05) between them. Comparative of IVF training groups of Azoospermic patients shown by Figure 6. There are nonsignificant differences between them. Figure 7 shows Comparative of get pregnant from IVF groups of Azoospermia patients, from it there is significant differences (P<0.05). Comparative of Chromosome test groups of Azoospermic patients explained by Figure 8, Significant differences (P<0.01) between 2 groups. Figure 9 describes Comparative of fertility of female abnormality groups of Azoospermic patients.

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non-significant differences (P>0.05) is noticed. Infertility periods groups are explained by Figure 10, there is non-significant differences (P>0.05). Last figure shows Comparative of obstructive type groups of Azoospermia patients. Significant differences (P<0.01) between 2 groups as non-obstructive type is the largest.



Figure 1: Percentage of patients with abnormal SFA



Figure 2: Percentage of patients with Azoospermia

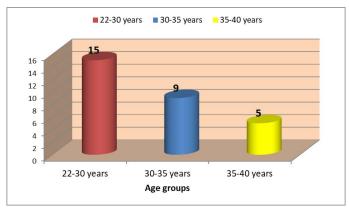


Figure 3: Comparative of age groups of Azoospermic patients Chi²=5.241, Sig=0.07 non-significant differences (P>0.05)

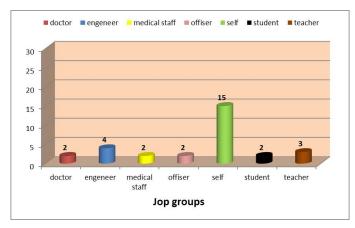


Figure 4: Comparative of job groups of Azoospermic patients Chi²= 32.067, Sig=0.002** Significant differences (P<0.01)

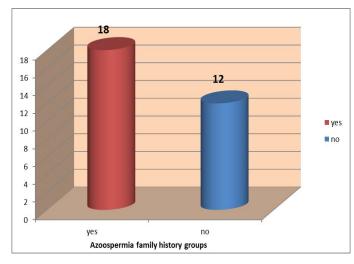


Figure 5: Comparative of family history groups of Azoospermic patients

Chi²= 1.200, Sig= 0.273 non-significant differences (P>0.05)

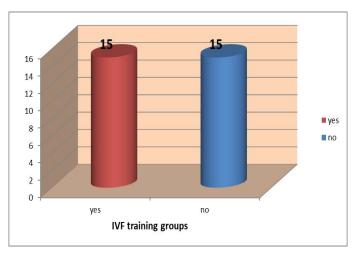


Figure 6: Comparative of IVF training groups of Azoospermic patients Chi²= 0.0, Sig= 1.00 non-significant differences (P>0.05)

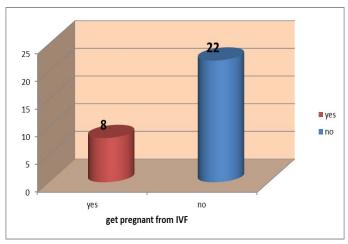


Figure 7: Comparative of get pregnant from IVF groups of Azoospermic patients

Chi²= 6.533, Sig= 0.011* significant differences (P<0.05)

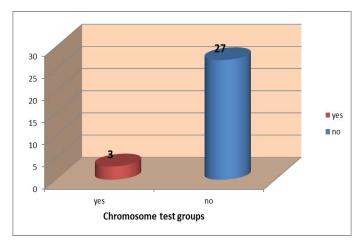


Figure 8: Comparative of chromosome test groups of Azoospermia patients

Chi²= 19.200, Sig=0.0002** Significant differences (P<0.01)

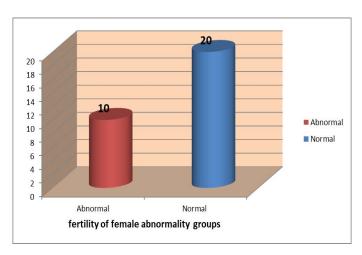


Figure 9: Comparative of fertility of female abnormality groups of Azoospermic patients

Chi²= 3.333, Sig=0.07 non-significant differences (P>0.05)

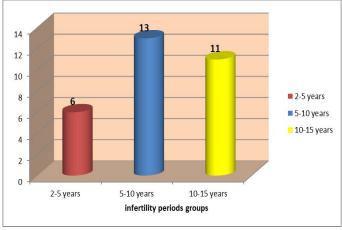


Figure 10: Comparative of infertility periods groups of Azoospermic patients

Chi²= 2.600, Sig=0.27 non-significant differences (P>0.05)

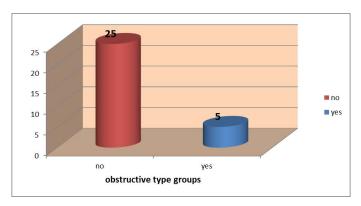


Figure 11: Comparative of obstructive type groups of Azoospermic patients

Chi²= 13.333, Sig=0.007** Significant differences (P<0.01)

DISCUSSION

We observed that 600 males with abnormal SFA out of 2000 (30%) and 30 males with azoospermia from those 600 males (5%). In Iraq study done at 2012, total of 116 males with primary infertility/: 32 azoospermic (AZO)14. In another study conducted in Iraq, 250 infertile couples' male members were assessed, and a 10% prevalence of azoospermia was found¹⁵. In their investigation, Rima D and his colleagues found that 17% of participants had azoospermia. This result is not unexpected given that it has previously been established that around 25% of infertility is attributable to males¹⁶. Previous workers in Pakistan revealed that azoospermia was present in 17.5% and obstructive azoospermia in 8,5% of the population¹⁷. The prevalence of azoospermia varies by area in India, from 38% in Karnool to 37,4% in Jodhpur, while it is 51% in Karnool for oligozoospermia¹⁸. Patients with infertility in Sweden's andrology unit reported oligospermia in 27.6% of cases and azoospermia in 572.4% of cases¹⁹. The American Society for Reproductive Medicine published a study in 2006 that showed 3,017 people had screening semen analysis performed on them. 15% (443) and 3% (95) of those satisfied the criteria for oligozoospermia, respectively. 55 (58%) of the 95 males with azoospermia showed up for additional testing. In 93% (51/55) of individuals, non-obstructive azoospermia was shown to exist. The remaining 7% (4/55) individuals had obstructive azoospermia²⁰, since non-obstructive azoospermia is more common than obstructive azoospermia in our study. According to our research, the majority of self-employed males are from lower socioeconomic backgrounds, and they frequently choose dangerous professions like farming, construction, and industry. Previous research has shown links between each of these professions and reduced male fertility²¹⁻²⁷. Additionally, it has been demonstrated that exposure to chemicals and pesticides, such as those experienced by farm laborers, has a detrimental effect on sperm parameters²⁸⁻³⁰. These elements could all have a role in the rising incidence of non-obstructive azoospermia in our community^{31,32}. Biomarkers for various male reproductive system problems, such as male infertility, may be found in seminal plasma. By identifying and analyzing several molecules expressed in males with normal and defective spermatogenesis' seminal blood, future study may offer insight on the molecular origins of male infertility and possibly azoospermia³³⁻³⁴. Despite a resurgence of interest in genetic research to identify the root causes of idiopathic infertility, the overall pattern has been to disregard advancements in diagnostics in favor of ART, which is more expensive but more successful³⁵⁻³⁶.

CONCLUSION

The prevalence of azoospermia in our province (5%). Half of the patient

only undergone IVF program (in vitro fertilization) because there is no IVF center in our province in addition to high cost of private centers.

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Competing Interest: None

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