Impact of Air Pollution on Semen Quality: The Specific Situation of Terni (Central Italy)

Ilaria Bocci1#, Giulia Poli2#*, Consuelo Fabi3, Francesco Bronzini4, Giulia Ceccotti5 and Stefano Brancorsini2

1Department of Laboratory Medicine, Policlinico Tor Vergata of Rome, Italy
2Department of Medicine and Surgery, University of Perugia, Terni, Italy
3Andrology and Urogynecology Clinic, Santa Maria Terni Hospital, University of Perugia, Terni, Italy
4Santa Maria Terni Hospital, University of Perugia, Italy
5Santa Maria Terni Hospital, University of Perugia, Italy
#These authors contributed equally to this work
*Corresponding author: Giulia Poli, University of Perugia, Department of Medicine and Surgery, Via Mazzieri 3, 05100 Terni, Italy

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ABSTRACT

To evaluate the impact of environmental pollution on male fertility, seminal biofluid parameters of men living in the contaminated urban area of Terni-Papigno, were compared with those who live in rural areas with low risk of pollution. Patients referred to the Seminology Laboratory of the Division of Andrology and Urology Department were enrolled for infertility evaluation. Semen samples were evaluated for seminal volume, pH, total sperm count, progressive motility, and morphology and leukocyte concentration. Data were analyzed with GraphPad Prism 6.0 and results were reported as mean ± standard deviation (SD). We report an association between ambient air pollution and sperm quality. In patients residing in areas with high environmental exposure, sperm concentration and count decrease significantly. Sperm motility, morphology and vitality did not demonstrate statistical differences. Identifying risk factors to improve the management of human wellness and health throughout standardized analysis, which correlates the toxic bioaccumulation of the seminal fluid with semen parameters, might be the main objective to be considered in public prevention policies.

Abbreviations: SD: Standard Deviation; WHO: The World Health Organization; CO: Carbon Monoxide; NO₂: Nitrous Dioxide; SO₂: Sulfur Dioxide; ROS: Reactive Oxidative Species; PM₁₀: Particulate Matter With Diameter Less Than 10 Microns; ARPA: Regional Agency for Environmental Protection

Introduction

Infertility is a prevalent condition affecting an estimated 72.4 million people globally that is well recognized by The World Health Organization (WHO). Although prevalence data are lacking, 9% of couples struggle with fertility issues and male factor contributes to 50% of the issues. Many genetic and lifestyle factors have been implicated in male infertility; however, about 30% of cases are still thought to be idiopathic [1]. The mechanism by which medical conditions affect fertility includes effects on hormonal levels, impairment of sexual function (including ejaculatory function), or impairment of testicular function/spermatogenesis. In the last 70 years, a decrease in sperm fertility and quality has been observed, including sperm count, ejaculate volume, alterations in sperm concentration and morphology [2]. Recent studies suggest that men with abnormal semen parameters have a higher risk of
testicular malignancy [3]. Nowadays, environmental and lifestyle factors could be possible contributors to infertility conditions, such as use of smoke cigarettes, increasing of both parents age conception, abuse of alcohol and drugs, physical inactivity, obesity, social stress, exposure to environmental contaminants (polycyclic aromatic hydrocarbons-PAHs, or heavy metals, for examples) and air pollution [4,5]. In particular, epidemiological and experimental studies explained the link between air pollution and alterations of sperm parameters as the main risk factors for male infertility.

Human activities such as transport, industrial and agricultural emission are considered the main causes of air pollution (solid particles, liquid droplets or gases), and people that living near these area, are more exposed to enhanced emission source of carbon monoxide (CO), nitrous dioxide (NO₂), sulfur dioxide (SO₂), ozone and lead [6]. Ambient air pollution is associated with systemic increases in oxidative stress, to which sperm are particularly sensitive. In this context, reactive oxidative species (ROS) have been related with a broad array of spermatogenesis effects, including the decrease of progressive motile sperm count, viability, abnormal sperm morphology, and fertilization rate and spermatic cell numbers [7]. In Italy, 12,482 areas with a high risk of environmental pollution, mainly due to industrial emission, were been identified. Some central provinces, such as Città di Castello, Foligno and Perugia exceeded the limit set for particulate matter with diameter less than 10 microns (PM₁₀) and O₃ emissions, in 2019. In particular, Terni is one of the most polluted urban and industrial area in Central Italy [8]. In fact, is situated in an intermountain depression, delimited by the Apennine mountain range. This area is characterized by the presence of typical urban PM₁₀ emission sources such as vehicular traffic, domestic heating and industrial emission sources from a power plant for waste treatment. Particular geomorphological and meteorological conditions of Terni basin, limit the dispersion and augment the accumulation of the atmospheric pollutants.

The "Thyssen Krupp AST", a large steel factory founded at the end of 19th century and two more recent chemical industrial areas, are located close to the city center [9]. As a result of the intensive industrial activities and the geographical location, atmospheric pollution is the major local issue with high PM concentrations occurring throughout the year. According to European Commission Law, the daily maximum PM₁₀ concentration allowed in cities is 50 µg·m⁻³ [9]. The threshold has not to exceed more than 35 times per year. In Terni, the atmospheric PM₁₀ concentration exceede daily limit on more than 70 days in 2012, as recorded by Regional Agency for Environmental Protection (ARPA), Umbria. The aim of the present study is to provide an association between ambient air pollution and sperm quality, analyzing seminal biofluid parameters of man living in the urban area of Terni-Papigno, with a high risk of pollution, compared with those who live in rural areas with low risk of pollution.

Materials and Methods

Study Participants

Signed written consent was obtained from all 52 participants (age of 20–40 years) enrolled in this study from January 2018 to December 2019. Patients referred to the Seminology Laboratory of the Division of Andrology and Urology Department for an infertility evaluation. Female partners of the infertile men were subjected to general gynecological evaluation and were reported to have normal reproductive health. Residence in the province of Terni was an inclusion criteria while, systemic and chronic disease, genetic abnormalities, alcohol or drug abuse, hormone treatment, varicocele infection microchidism and cryptorchidism, prostatitis and other factors that could affect semen quality (such as fever, medications, exposure to X rays etc.) were exclusion criteria. Men were divided in group A and B that includes 30 patients from the high pollution environmental risk area of Terni-Papigno and 22 subjects that live in neighboring areas, with a low risk of pollution, respectively (Table 1).

Table 1: Demographic and environmental patient’s classification.

| Groups | Enviromental risk | Population Density | PM₁₀ | NO₂ |
|--------|-------------------|-------------------|------|------|
| A (n=30) | SNI | Urbanized Area | Class 3 (12% of the surface) | Class 4 (11,8%) |
| B (n=22) | - | Rural Area | Prevalence of Class 1 and 2 | Prevalence of Class 1 and 2 |

Note: SNI: Sites of National Interest; PM₁₀, Particulate Matter ≤ 10 µμ; NO₂, Nitrogen dioxide; OCSE Organization for Economic Cooperation and Development, class 1 and 2: rural area with population density <150 inhabitants/ km² and PM₁₀< 10 µg/m³; class 3 and 4: rural area with population density >150 inhabitants/ km² and PM₁₀> 10 µg/m³

Semen Analysis and Preparation of Samples

Semen samples were collected at the Andrology and Urology Laboratory by masturbation into a sterile container after 2–7 days of sexual abstinence and were analyzed immediately after liquefaction, according to the WHO guidelines [10]. Each sample was evaluated for seminal volume, pH, total sperm count, progressive motility, and morphology and leukocyte concentration. Semen volume was measured by graduated pipettes. Calibration strips were used to measure the seminal fluid pH. For the evaluation of the sperm concentration, following semen liquefaction, 10 µL of non-diluted,
well-mixed semen sample was at first loaded in the middle of a clean Burker counting chamber, maintained at the temperature of 37 °C, gently covered with a cover glass, and examined using 200× or 400× magnification. The sample was diluted before proceeding with the sperm count. 1:2 dilution was used, strictly following the WHO 2010 manual recommendations [10]. The final concentration was calculated as: [(number of spermatozoa counted/the number of lines) x dilution factor] and expressed as 106 spermatozoa/mL. To evaluate the sperm motility, immediately after semen liquefaction, 10 μL of undiluted, well-mixed semen sample was loaded in the middle of a clean Neubauer counting chamber, maintained at the temperature of 37 °C, gently covered with a cover glass, and examined using 200× magnification.

Sperm motility was assessed in 200 random spermatozoa and characterized as progressive and non-progressive motility. The total motility was calculated as the sum of progressive and non-progressive motility. Both progressive and total motility were expressed as percentages. Sperm morphology was evaluated in 200 spermatozoa and the value was expressed as percentages. Finally, the vitality was assessed using eosin staining according to the WHO recommendations [10].

**Statistical Analysis**

Data were analyzed with GraphPad Prism 6.0. Results were reported as mean ± standard deviation (SD). Analysis of variance and Kruskal-Wallis tests were performed. The significance threshold was set at 0.05.

**Results**

Patients were classified according to Demographic and Environmental characteristics (Table 1) Total sperm count (Figure 1) and sperm concentration (Figure 2) decreased significantly in seminal biofluid of patients living in urbanized area of Terni-Papigno (group A) when compared to subjects who lived in rural areas (group B). Seminal volume (mL) did not reach significant difference (Figure 3) between the groups considered.

![Figure 1: Total sperm count (n*106) expressed as median with range and interquartile range in group A compared to group B. *P<0.05.](image1)

![Figure 2: Concentration of total sperm count (n/ml) expressed as median with range and interquartile range in group A compared to group B. *P<0.05.](image2)
Discussion

In this study, conventional semen parameters (volume, appearance, pH, motility, and viability, and morphology, presence of aggregations, agglutination and leukocyte concentration) were analyzed for each sample according to the 2010 WHO criteria. Our results showed a significant decrease of total sperm count in group A patients respect to group B subjects (Figure 1). Nowadays, several studies showed how pollution can affect human fertility. Sperm count and sperm motility were reduced in men living in polluted cities. A decrease of sperm count was observed in America and Northern Europe during the last 50 years [2] and the reduction of semen quality and sperm count especially in industrialized countries supports the evidence that adverse environmental factors are key factors for men infertility [11]. Furthermore, different studies reported the enhancement of infertility, urogenital malformation and chronic disease (cancer, diabetes, etc.) in areas with a high environmental pressure [12]. In group A, a significant decrease of sperm concentration was demonstrated, respect to group B patients (Figure 2). In 2019, a longitudinal analysis on 8,945 semen samples, suggested the specific role of O₃ pollution in sperm concentration [13]. The effect of air pollutants and ozone on sperm quality, was strongly elucidated by Sokol, et al. [14]; they reported that pollutants have no effect on sperm quality, however, ozone could make changes in spermatozoa, including sperm DNA fragmentation via oxidative stress, resulting in decreased fertility [14].

The toxicity of O₃ was largely demonstrated to be the major oxidant of photochemical smog and its exposure produces reactive ROS at respiratory level [15,16]. Extra pulmonary toxicity suggests that O₃ or O₂ reaction-products can cross blood-gas barrier and be absorbed into the circulating bloodstream, creating an environment caring to an inflammatory reaction [17,18]. It is unclear how O₃ can negatively affect sperm quality, but O₂-induced oxidative stress could be a possible mechanism, through which testicular and sperm function could be altered [19,20]. Under physiologic conditions, spermatozoa exist in a balanced environment of ROS and antioxidants, where ROS determine the biochemical steps required for normal fertilization (capacitation and the acrosome reaction). However, excessive amounts of ROS produced by leukocytes and immature spermatozoa can damage mature spermatozoa and the integrity of sperm DNA [21,22]. Concerning seminal volume (mL), our results did not reach significant difference between groups considered (Figure 3). Several studies suggested that alterations of semen parameters, including volume, progressive motility, total motility or morphology, may not relate to air pollution [23]. Several works observed a reduction of both sperm motility and morphology, associated with air heavy metals (lead, mercury and cadmium) [24,25], that determine alterations in sperm DNA.

Rubes and colleagues showed that short-term exposure to pollutants caused serious damages in men and women’s reproductive system. Moreover, this study demonstrated that the incidence of sperm DNA alterations due to air pollution is higher in middle-aged men [26]. Also, exposition time of contaminants seem to be determinant, in fact a study of Hammoud showed that 3 months exposure to air pollution decreases motility levels, while removing contamination can restore normal parameters levels [27]. In an extensive cross-sectional study, Xu and co-workers, founded a strong correlation between motility, concentration, and morphology of semen biofluid of men exposed to air pollutants with the use of cigarette and alcohol [28]. On the other hand, Selevan, et al. [29] did not observed any relevant alerations in sperm count, motility and morphology of young men exposed to air pollutants, except for sperm DNA and chromatin [29]. Moreover, several authors specify that the morphological change of spermatozoa after exposure to pollutants is not an indicative diagnostic parameter. Association
between air pollution and alteration of specific semen parameters is not clear and different relevant factors, such as geographic areas and lifestyle, should be considered for male infertility diagnosis.

Conclusion

The current study provides evidence of an association between ambient air pollution and sperm quality. Patient residents in areas with high environment exposure had a significantly decrease in sperm quality especially for sperm concentration and count but had no impact on the other sperm parameters of spermogram (motility, morphology, vitality). The individual role of specific pollutants is difficult to identify, since patient in this study are typically exposed to several pollutants simultaneously. The physiopathology leading to altered fertility is poorly understood. In the literature there are forward four mechanisms to explain the negative impact of air pollution on sperm, as hormonal disturbances, oxidative stress induction (ROS), cell DNA and epigenetic alterations, probably working in combination. Clearly, more research is needed to understand the detrimental effect of the pollutants on sperm and to characterize their action in more detail. Our results suggest the important role of human sperm as an early and sensitive biomarker of environmental pollution as it could represent an ideal tool for investigating and promoting health surveillance especially in environmental risk areas. Environmental contaminants and bad lifestyles can impair reproductive health and overall health, encouraging the development of chronic degenerative diseases affecting the adult and, through the sperm epigenome changes, future generations. Thus, identifying risk factors to improve the management of human wellness and health throughout standardized analysis, which correlates the toxic bioaccumulation of the seminal fluid with the multiple semen parameters, might be the main objective to be considered in public prevention policies.

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Conflicts of Interest

The authors declare that there is no conflict of interest regarding the publication of this paper.

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