Effects of indoor swimming pools on the nasal cytology of pool workers

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Abstract

Objective: We aimed to evaluate the relationship between swimming pool pollutants and allergic rhinitis in swimming pool workers.

Materials and methods: Twenty-seven indoor pool workers (group 1) and 49 control subjects (group 2) were enrolled in the study. A skin prick test was performed and a nasal smear was obtained from each subject to evaluate rhinitis.

Results: When the groups were compared in terms of epithelial cells, group 1 had significantly more epithelial cells than group 2. When the groups were compared with regard to eosinophils, group 1 had significantly more eosinophils than group 2. The skin prick test results for both groups were not significantly different.

Conclusion: Indoor pool workers showed severe symptoms of rhinitis and eosinophilic nasal cytology, likely due to chlorine. Nasal cytology is an easy-to-administer diagnostic test and can be used to follow up rhinitis in indoor pool workers, along with nasal endoscopy, a detailed clinical history and a skin prick test.

Key words: Rhinitis; Cytology; Epithelial Cells; Eosinophils; Swimming Pools; Environmental Impact; Chlorine; Hypochlorite; Pathophysiology

Introduction

Under optimal nasal breathing conditions, air passes over the nasal mucosa resulting in warming, humidification and cleansing. However, these conditions can be influenced by several factors. Indoor air quality may negatively affect the sinonasal mucosa and is associated with occupational and smoking irritants, which can cause chronic or allergic rhinitis.¹ To maintain proper nasal functions, the optimal indoor air temperature should be approximately 23 °C with 50 per cent humidity.² Indoor air is often cleaned to limit the levels of pollutants by modifying temperature and humidity in houses and offices.² However, the improper maintenance, design and operating of air-conditioning systems in non-industrial indoor areas contributes to an increased prevalence of rhinitis.³ In Turkey, smoking is prohibited indoors and in common social areas.

Allergic rhinitis is a symptomatic disorder caused by immunoglobulin E-mediated inflammation of the nasal membranes, which is induced by exposure to an allergen or allergens. Allergic rhinitis frequently causes symptoms that include sneezing, nasal obstruction, itching and discharge, and it may also affect coexisting diseases. Diagnosis is based on a detailed clinical history, an otorhinolaryngological examination and epidermal skin tests. T helper 2 lymphocytes secrete cytokines, such as interleukins 3, 4, 5, 13 and 17 and eotaxin (CCL3 and CCL5), which are released from eosinophils and are associated with the symptoms and severity of allergic rhinitis.^{4–6}

Indoor swimming pools are relatively hot environments (approximately 30 °C) and their humidity ranges between 50 and 60 per cent continuously.⁷ Sodium and calcium hypochlorite, which are commonly used as disinfectants in swimming pools, cause irritation of the nasal mucosa and exacerbate the symptoms of rhinitis.⁸ Rhinitis is common among athletes, especially swimmers.⁹ However, pool workers spend more time in and around swimming pools compared to swimmers and are also exposed to sodium and calcium hypochlorite. A number of studies of swimmers have been performed, but fewer studies have focused on pool workers.^{8–12}

In one study, swimmers had poorer nasal functions compared to runners, but only the post-nasal drip was

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significantly different between the two groups. The authors concluded that the results supported the existence of swimming-induced rhinitis independent of the atopic status of the athlete.¹⁰ In a further study, nasal symptoms increased in competitive swimmers and were not related to seasonal allergen exposure in atopic athletes; thus, these symptoms likely resulted from exposure to chlorine derivatives.¹¹ Evaporation of airborne trichloramine from pool water can cause eye and nasal symptoms among pool workers and trainers, who spend extended periods of time at swimming pools. The most frequent symptoms are red eyes, a runny nose, loss of voice and cold-like symptoms.^{12,13} These symptoms are due to irritation of the nasal mucosa and can be investigated using nasal cytology. Nasal smears can be assessed easily using light microscopy and can reveal relevant information about the predominant cell types infiltrating the nasal cavity. The dominant cell type allows the identification of the type of rhinitis.

In this study, we used nasal cytology and the skin prick test to evaluate the relationship between swimming pool pollutants and rhinitis in pool workers who had high exposure to indoor pollutants compared to a symptom-free control group.

Materials and methods

The study protocol was approved by the Ethics Committee of the Eskişehir Osmangazi Faculty. All subjects gave their written informed consent. The study was performed at the indoor, half-Olympic size swimming pool of Anadolu University, Eskişehir, Turkey. In total, 27 indoor pool workers were enrolled in the study (group 1). These subjects worked 8 hours per day within the pool area, 6 days per week for over one year. The group comprised trainers, lifeguards and cleaners.

The control group (group 2) consisted of 49 office workers with the same age and sex distributions as group 1. The control group worked in a controlled, air-conditioned working environment.

Exclusion criteria included any sign of infection at the time of the study, a previous history of any chronic disease, pregnancy, use of antihistamines, nasal or systemic steroids, or evidence of leukotrienes in the three months preceding the study.

All subjects were evaluated using a detailed clinical history, an otorhinolaryngological examination and a skin prick test. After obtaining a detailed clinical history from each patient, a skin prick test was performed and a nasal smear was obtained to evaluate rhinitis. Nasal smears were taken from the middle third of the inferior turbinate of each nostril using a cotton swab. The smears were fixed in ethyl alcohol and Wright stain solution was applied for examination by light microscopy (BX53 Upright microscope, Olympus America Inc., Center Valley, Philadelphia, USA). The same histology specialist examined each sample in a blinded fashion. The number of goblet

cells, epithelial cells, neutrophils, basophils and eosinophils was counted in each magnified area. Finding one cell in a $10 \times$ magnification field was counted as 1 and finding two cells in that area as 2. Both were considered positive cellular findings. All samples were also checked for dysplasia, metaplasia and atypia.

Pool specifications

The half-Olympic size pool is 25 m long, 16 m wide and 2.25 m deep. It holds 1200 m^3 of water, is filtered approximately every 4 hours and has a standard temperature of 26-28 °C throughout the year. All chemical agents, such as liquid and powdered chlorine, used for cleaning the pool are the same in all of the pools provided by the Ministry of Health in Turkey. Daily analysis of the pool water is performed routinely. Deeper cleaning is performed by a pool robot and by means of deep skimmer nets. The pool is open for 11 months a year, 6 days per week.

Statistical analyses

Statistical analyses were peformed using SPSS version 18.0 for Windows (SPSS, Inc., Chicago, Illinois, USA). The Shapiro–Wilk test was used to test data normality. For normally distributed, continuous data, groups were compared using independent sample *t*-tests; the Mann–Whitney *U* test was used for variables that were not normally distributed. The general linear model for repeated measurements was used for the intra-group comparisons. The chi-square (χ^2) and exact tests were used to compare categorical variables. A *p* value < 0.05 was chosen as indicating statistical significance.

Results

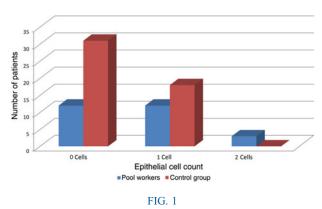
The study included a total of 76 patients: 41 women and 35 men. Both groups 1 and 2 consisted of more women than men. However, there was no significant difference in gender distribution between the two groups ($\chi^2 = 0.064$, p = 0.80, p > 0.05). The groups were also similar with regard to mean age (F = 1.297, p = 0.203, p > 0.05). Group 1 comprised 27 pool workers with an age range of 19–49 years (mean age: 31.33 ± 7.64) and group 2 (control group) comprised 49 subjects with an age range of 25–53 years (mean age: 33.65 ± 6.94). The general characteristics of both groups are summarised in Table I.

Of the 27 subjects in group 1, nasal cytology revealed that 12 (44.4 per cent) had no epithelial cells, 12 (44.4 per cent) had one epithelial cell and three (11.1 per cent) had two epithelial cells. In group 2, nasal cytology revealed that 31 (63.3 per cent) of the 49 subjects had no epithelial cells and only 18 (36.7 per cent) had one epithelial cells. In group 2, no subjects had two epithelial cells. Group 1 had significantly more epithelial cells than group 2 ($\chi^2 = 6.796$, p = 0.033) (Figure 1).

Of the 27 subjects in group 1, nasal cytology revealed that 15 (55.6 per cent) had no eosinophils,

TABLE I			
CHARACTERISTICS OF THE TWO STUDY GROUPS			
	Group 1	Group 2	P value
Subjects	27	49	N/A
Men/women	13/14	22/27	0.80
Age range	19-49	25-53	0.203
Skin prick test positivity	8	9	0.401
Eosinophils	12	12	0.027
Neutrophils	12	16	0.726
Basophils	2	0	0.123
Goblet cells	14	18	0.174
Epithelial cells	15	18	0.033

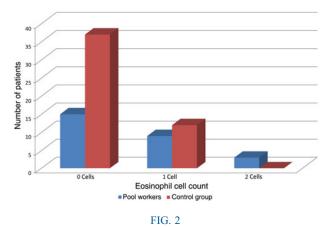
N/A = not applicable



Epithelial cell disturbance in the two study groups.

nine (33.3 per cent) had one eosinophil and three (11.1 per cent) had two eosinophils. In group 2, nasal cytology revealed that 37 (75.5 per cent) of the 49 subjects had no eosinophils and only 12 (24.5 per cent) had one eosinophil. In group 2, no subjects had two eosinophils. Group 1 had significantly more eosinophils than group 2 ($\chi^2 = 6.950$, p = 0.027) (Figure 2).

Dysplasia, metaplasia and atypia were not observed in nasal smears from either group. The number of goblet cells ($\chi^2 = 3.058$, p = 0.174), neutrophils ($\chi^2 =$ 1.427, p = 0.726) and basophils ($\chi^2 = 3.728$, p =0.123) was not significantly different between the two groups.



Eosinophil cell disturbance in the two study groups.

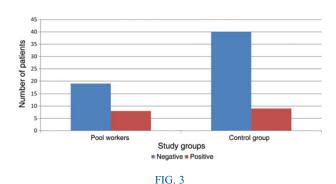
Of the 27 subjects in group 1, 8 (29.6 per cent) had positive skin prick tests, while 9 of the 49 subjects (18.4 per cent) in group 2 had positive skin prick tests to one or more allergens (mostly fungus and mites). The skin prick test results of both groups were not significantly different ($\chi^2 = 0.706$, p = 0.401) (Figure 3). All subjects with a positive skin prick test had eosinophils present in their nasal smears. Of the 27 subjects in group 1 with a negative skin prick test (19 subjects), four had only neutrophils and four had eosinophils and neutrophils.

Discussion

Two main causes of allergic and non-allergic rhinitis affecting indoor swimmers and pool workers have been suggested: namely, fungus and mites in the case of allergic rhinitis, and chlorine and its derivatives in the case of non-allergic rhinitis.^{8,14–17}

Hypersensitivities to fungus and mites are aggravated by the time spent within the pool area, which leads to cytological changes. The concentrations of chlorine by-products in indoor air, such as trihalomethanes and trichloramine, have been found to be correlated with the concentrations found in the pool water.¹⁸ Pool water is more polluted in the winter than in the summer; this is due to the number of swimmers and the low air exchange rate in the pool. Indoor air pollution is also related to the size of the swimming pool hall and the volume of water in the pool. Increasing the flow of fresh air in the pool complex or restricting overtime work and the number of swimmers in the pool could reduce the symptoms of rhinitis.¹⁸

Nasal cytology is easy to perform and can indicate inflammation by identifying the cell types present; it can give accurate and quick results in a short period of time. Eosinophil counts are used for screening allergic rhinitis patients. Therefore, we combined nasal cytology with a skin prick test. Gelardi *et al.* reported predominantly neutrophilic inflammation and allergic rhinitis in symptomatic swimmers and suggested that this could be prevented by avoiding direct contact with chlorinated pool water.⁸ Unlike other reports, in our study, we evaluated the nasal cytology of indoor pool workers who had no direct contact with the swimming pool to assess the indirect effect of the



Skin prick test results for the two study groups.

evaporation of airborne chlorine. In contrast to Gelardi *et al.*⁸ we did not find significant numbers of neutrophils in the nasal smears, and considered that this was likely due to the lack of contact between the nasal mucosa and the pool water.

A recent study from Iran showed that 25.61 per cent of a group of 50 allergic patients had nasal smears positive for eosinophils, with the tests showing a sensitivity, specificity, positive predictive value and negative predictive value of 74, 90, 88 and 77 per cent, respectively.¹⁹ We found significant epithelial and eosinophilic infiltration in the pool worker group (p = 0.033 and p = 0.027, respectively). A nasal smear is the only method for evaluating the predominant cell types within the nasal mucosa. A nasal smear is not a routine method for detecting rhinitis, but we aimed to investigate the correlation between cell count and skin prick test positivity as one of the markers of allergic rhinitis.¹⁹ In our study, all subjects with a positive skin prick test had eosinophils in their nasal smear; thus, allergic rhinitis was more prevalent in indoor pool workers than neutrophilic rhinitis, even though neutrophilic rhinitis has been previously reported in swimmers.⁸ In the skin prick test, 29.6 per cent of subjects in the study group and 18.4 per cent in the control group tested positive. Indoor pool workers had a high prevalence of allergies (as determined by comparing the skin prick test and eosinophil count results with the control group). These results are similar to those reported previously for swimmers and athletes.^{8–10} Swimmers with rhinitis could have concomitant asthma;^{8,12,20} therefore, indoor pool workers should also be checked for allergic disease of the inferior airway.

- Upper and lower respiratory tract disease occurs in swimming pool workers and users
- We wanted to evaluate the relationship between swimming pool pollutants and rhinitis in swimming pool workers
- Pool workers had significantly more epithelial and eosinophilic cells than the control group
- Indoor pool workers, like swimmers, had severe rhinitis symptoms and eosinophilic nasal cytology likely due to chlorine

Temperature, humidity and exposure to high chlorine levels may affect the symptoms of rhinitis and the nasal cytology of indoor pool workers.^{2,8,9,11} This may present as neutrophilic⁸ and eosinophilic⁹ infiltration, as we have reported here. Some common working areas have a stronger association with allergic or non-allergic rhinitis,^{20,21} including indoor swimming pools. Working for many hours in an allergic environment, such as classrooms with polluted air (formaldehyde, acrolein and mould species), can lead to allergic rhinitis or asthma.^{20–22} Indoor swimming

pools should be added to the list of allergy-inducing working areas; fresh air based cooling systems may be the solution to this problem.

Indoor pool workers, like swimmers, have a high prevalence of rhinitis and eosinophilic nasal cytology, likely due to the presence of chlorine. This could be avoided by adopting temperature, humidity and chlorine concentration standards for indoor pools. Nasal cytology is an easy-to-administer diagnostic test and, like nasal endoscopy, a detailed clinical history and a skin prick test can be used to follow up rhinitis in indoor pool workers.

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