Etiological factors of systemic sclerosis in the southeast region of Romania

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Abstract. Systemic sclerosis (SSc) is a relatively rare autoimmune disease with skin and visceral involvement, having a yet unknown etiopathogenesis. Research has shown that professional exposure to various polluting chemicals such as dyes, aliphatic and aromatic organic solvents, inhalable silica dust or certain heavy metals, can be triggering factors for this disease when they overlap a predisposing genetic profile. Smoking is still a debated factor involved in the etiology of SSc, as authors have divergent opinions on this matter. The present study was designed to analyze the etiological factors identified in the group of 37 patients with diffuse and limited SSc from the southeast region of Romania and the results were compared to the literature data. In the group of patients included in this study, occupational exposure and smoking history were not present in all patients, and a hereditary factor was identified only in an isolated case. The majority of patients suffered from a major negative psychological event or from long-term stressful situations and these factors were associated with smoking history or occupational exposure; this suggests that SSc is initiated in a set of cumulative triggering factors.

Introduction

Systemic sclerosis (SSc) is a rare chronic autoimmune disease with high mortality rates (1). The severity of this disease stems from multiple organ morpho-functional damage with multiple complications: Ulcers, gangrene, pulmonary and myocardial fibrosis, pulmonary hypertension, scleroderma renal crisis, esophageal damage and malabsorption syndrome (2-4). Skin induration in SSc requires differential diagnosis with other sclerodermal diseases (5,6). The pathogenesis of this disease is complex and has many yet uncertain variables, very much similar to the pathogenesis of many neoplasms (7-9). Therefore, it is recognized that a number of triggering environmental factors may overlap with predisposing genetic factors, including occupational exposure to various chemicals including aromatic hydrocarbons, silicates, polyvinyl chloride, drugs (amphetamine, bleomycin, pentazocine), hand vibrations and some infections (10-14). In a 2018 review, Abbot et al concluded that exposure to silicates alone is an important risk factor. Various authors express divergent views concerning exposure to organic solvents (15). In another review, Walecka et al highlighted the role of silicates in the etiology of SSc, but also of multiple organic solvents such as ketones, aromatic and aliphatic hydrocarbons, white spirit solvent for oil-based paints, smoke from welding; underlining the need for protective measures for workers with such occupational exposure. The authors pointed out the lack of clear evidence concerning air pollution involvement in the onset of SSc; the same issue concerns asbestos exposure, heavy metals and silicone implant (16). Occupational exposure raises the issue of introducing SSc to the occupational diseases category. The most common occupational exposure is to silica dust, which induces silicosis, rheumatoid arthritis and lung cancer; the association of silicosis with SSc or mixed connective tissue disease, known under the name of Erasmus syndrome, has also been considered for inclusion in the occupational diseases category. Researchers suggest that SSc associated with silica dust exposure should be included in the occupational diseases category, even when silicosis is not present (17,18). Silicon
dioxide nanoparticles (nano-SiO$_2$) in their amorphous state are microparticles with a diameter of less than 100 nm and a very low density, which can easily become airborne and inhaled. It seems that nano-SiO$_2$ primarily affects the skin, and through inhalation induces inflammation and lung and cardiovascular toxicity. Nano-SiO$_2$ toxicity is dependent on the exposure dose, but also on the particle size; the toxicity increases with the decrease of particle size (15-30 nm), being proportionally indirect. SiO$_2$ nanoparticles degrade DNA chains and induce cell apoptosis (19). Among the organic solvents associated with SSc, the most common are aromatic solvents (benzene, toluene, xylene) and trichloroethylene, as well as halogenated solvents. Zhao et al. also revealed the significant exposure to ketones (18). Recent studies show that epoxy resins, which are included in the list of occupational toxins, may be involved in the onset of SSc (17,18,20). Not only occupational exposure to toxins poses a risk, but also constant or accidental pollution through water and air contamination with halogenated compounds and aromatic hydrocarbons, such as benzene, xylene and toluene (21). The study on volatile aromatic hydrocarbon toxicity performed by Niculescu and colleagues on laboratory mice reveals some interesting facts. Experimental exposure of laboratory animals to xylene, toluene and benzene registered no clinical signs for chemical toxicity, only a weight increase starting from day 3 of exposure, becoming significant on day 11. This study opens the opportunity for further research in clarifying the toxic profile of these compounds (22). After in-depth analysis of smoking involvement in the onset of SSc, Walecka et al. concluded that its role is not clearly supported (16). Although there is no solid evidence that smoking could induce SSc, Hissaria et al. revealed that smoking has an adverse effect on disease progression and decreases the survival rate in former and current smoking patients (23).

The present study was designed to analyze the etiological factors identified in the group of 37 patients with diffuse and limited SSc from the southeast region of Romania and the results were compared to the literature data.

**Patients and methods**

*Patients.* We conducted an observational study on a group of 37 patients composed mainly of women (81.1%) from urban areas (64.9%). The average age of the patients was 54.08±11.871 years, with variations between 28 and 76 years. Patients were admitted between February 2019 and March 2020 at the Internal Medicine and Rheumatology Wards of the University Hospital in Bucharest. Patients were diagnosed with SSc using the criteria found in the 2013 American College of Rheumatology (ACR)/European League Against Rheumatism (EULAR) (24,25). Patients with diffuse and limited SSc were included in the study in order to identify the etiological factors involved in the pathogenesis of SSc. Newly diagnosed cases and cases with disease evolution over 10 years were enrolled in the study. The agreement was obtained from the Research Ethics Committee of the Faculty of Medicine in Iasi (24.06.2017) as well as from the Ethics Council of the Clinical Hospital ‘St. Maria’ in Bucharest (5213/04.04.2019). All procedures in this study are in accordance with the Declaration of Helsinki.

**Patient interview.** We analyzed the etiological factors that could be involved in the pathogenesis of the disease. Patients were interviewed regarding smoking status, occupational exposure to various polluting chemicals and major psychological events with significant psychological shock or very stressful situations in the period preceding the onset of SSc. The genetic predisposition for ScS from consanguins (ascendants, kin of blood) was also analyzed. Patients were interviewed regarding the presence of SSc in family members, without the possibility of performing genetic studies, as a limitation of the study. The chemicals recorded in the medical literature for the fibrotic potential were targeted, but other toxic substances from the professional environment declared by the patients were also registered. The duration of the exposures was also noted. Patients' statements about the existence of a major traumatic psychological event, which preceded the onset of the disease and which they incriminated as a trigger, was the starting point for interviewing patients. We analyzed the psychological impact of the stressful situation and the way of reporting the patient to the traumatic episode. The evaluation was performed through a structured interview with 4 questions, through which I appreciated: Easy crying, sadness, insomnia and irascibility.

**Statistical analysis.** The data were obtained by interviewing the patients after they had signed the informed consent and it was introduced into an Excel file. The statistical data processing was carried out using SPSS 24.0 (IBM Corp.) using Microsoft Excel. The quantitative data were characterized through descriptive statistics; the qualitative data were characterized through frequency distributions and contingency tables, and the comparisons between samples were made using the Chi-squared test. All P-values were two-tailed; a P-value of 0.05 was considered significant.

**Results**

The group of 37 patients from the south-east region of Romania were introduced in the study, analyzed, and we concluded that only 24.3% of the investigated patients were exposed to toxic substances and industrial pollutants; no statistically significant differences were observed between the two subtypes of SSC, although the percentage of patients exposed to such toxins was slightly higher in those with limited SSc (28.6%) as compared to those with diffuse SSc (21.7%) (Table I, Figs. 1 and 2).

Most patients (81.1%) also experienced major traumatic psychological events or suffered psychological shocks; from this perspective, no significant statistical differences were observed between disease subtypes, although the patient percentage was higher among those with diffuse SSc (87.0%) as compared to those with limited SSc (71.4%) (Table I, Figs. 3 and 4).

Regarding the predisposing hereditary background, only one female patient was diagnosed with diffuse SSC and reported that the paternal grandmother suffered from identical symptoms, without mentioning whether she was diagnosed or not with SSc; this indicates the likelihood of SSc being genetically transmitted (Table I, Fig. 5).

Among the patients introduced in this study, 48.6% were non-smokers; the percentage of smokers to non-smokers being roughly equal (Table II, Figs. 6 and 7). Of those with a
smoking addiction, little over a half were categorized as heavy smokers. These percentages were found to be almost similarly in the diffuse SSc patient group; small differences were found in those with limited SSc, having a higher percentage of non-smokers (57.1%) with equal percentages of smokers to heavy smokers (21.4% each). An important fact is that only one patient out of the 37 was an active smoker (2.7%) and suffered from diffuse SSc (Table II).
Discussion

This paper is an observational study on a group of patients from the south-east region of Romania, that aimed to identify the etiological factors possibly involved in the pathogenesis of systemic sclerosis (SSc). The number of patients investigated was small due to the low prevalence of this disease. By analyzing occupational exposure to various toxic substances and inhaled pollutants, we found that only a quarter of patients were exposed in their life span for variable time periods of 2 to 5 years or up to 8 consecutive years; the exposure was to organic dyes and solvents such as toluene, xylene and acetonitrile, or to iron oxides and chromium compounds which can be found in synthetic resins used for sound proofing. The patient interviews also identified occupational exposure to hydrogen sulfite, acrolein, aldehydes, benzene, phenols and sulfur dioxide which are released from synthetic rubber, asphalt mixtures, bitumen, diesel and industrial oil. Other exposures identified were to formaldehyde released by pouring plastic into molds, ammonia released from animal manure in the zootechnic field, irritating nitrogen oxide released from pickled feed silos, hydrocyanic acid and isocyanates generated by acrylonitrile and its polymer. All these factors release toxic gases and vapors in the working environment (20). No exposure to respirable crystalline silica powder was identified. Quartz exposure, as a variety of crystalline silica, a well-known factor involved in SSc etiology, was also not found. Among patients exposed to occupational toxic agents, the differences between those with limited and diffuse SSc were not significant. From the remaining patients, a significant percentage of 3/4 (75.7%) did not register exposure to industrial pollutants.

Table II. Group structure by categories of smokers-total and by subsets.

| Smoking status       | SSc limited |          | SSc diffuse |          | Total |          | Chi-square | P-value |
|----------------------|-------------|----------|-------------|----------|-------|----------|------------|---------|
|                      | n | %      | n | %      | n | %      |            |         |
| Smoking history      |   |        |   |        |   |        |            |         |
| No                   | 8 | 57.1   | 10| 43.5   | 18| 48.6   | 0.673      | 0.714 (NS) |
| Yes-smoker           | 3 | 21.4   | 6 | 26.1   | 9 | 24.3   |            |         |
| Yes-heavy smoker     | 3 | 21.4   | 7 | 30.4   | 10| 27.0   |            |         |
| Active smoker        |   |        |   |        |   |        | 0.626      | 0.622 (NS) |
| No                   | 14| 100.0  | 22| 95.7   | 36| 97.3   |            |         |
| Yes                  | 1 | 4.3    | 1 | 2.7    | 1 | 2.7    |            |         |
| Total                | 14| 100.0  | 23| 100.0  | 37| 100.0  | 100.0      |         |

SSc, systemic sclerosis; NS, not significant.
Extremely important are the patients’ statements regarding major psychological events with psychological shock magnitude and highly extremely stressful life situations which they incriminated as factors that preceded the onset of SSC. Most patients strongly claimed that these events (accidents, illnesses and death of a close person or separation from their life partner) triggered the disease. The ratio of patients who had psychological events as a precursor to the disease was higher among those with diffuse SSC.

By analyzing the SSC predisposing hereditary background, a singular case involving a female patient with diffuse SSC was identified in whom hereditary transmission was from the paternal side without the possibility to perform genetic studies, as a limitation of this study. This observation confirms the role of genetic transmission in the predisposition for SSC, being recognized by many authors (10-14).

Concerning tobacco smoking, half of the patients were smokers and heavy smokers during their lifetime and, with one exception, all were in the process of smoking cessation in the course of this study. The ratio of smokers in the diffuse SSC subtype was slightly higher than that found in those suffering from limited SSC. Our observations suggest that tobacco smoking may be involved in the onset of the disease, especially in diffuse forms of SSC, although the findings are divergent (16,23).

The etiological factors identified were not consistently found among all patients from the analyzed group; Most patients investigated had an episode of traumatic psychological in the past, but smaller proportions of patients had a traumatic psychological event associated with history of smoking or with an occupational exposure or even both.

In conclusion, this autoimmune disorder seems to be caused by a combination of several etiological factors, most often in the background of significant psychological events with traumatic potential. Although the current study managed to reveal some potential etiological factors involved in the etiopathogenesis of SSC, it remains uncertain and requires further research in order to be clarified. The identification of the etiological factors of SSC could allow the issuance of a set of recommendations and the early evaluation of target individuals.

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Availability of data and materials

The analyzed data sets generated during the present study are available from the corresponding author on reasonable request.

Authors’ contributions

CB, EN, MC, AIH and LGS were involved in the conception of the study and had major contribution in the writing and revising of the manuscript. IAP, MLD, CO and BS assisted in the acquisition, analysis and interpretation of the data. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

The present study was approved by the Ethics Review Committees of the Medical University of Iaşi (24.06.2017) and of the Ethics Council of the Clinical Hospital ‘St. Maria’ in Bucharest (5213/04.04.2019) and was performed in accordance with the Declaration of Helsinki. All patients provided informed consent and approved the publication of data.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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