Characterization of *Mycobacterium tuberculosis* Complex Isolates from Greek Patients with Sarcoidosis by Spoligotyping

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Spoligotyping was undertaken with 38 *Mycobacterium tuberculosis* isolates from Greek sarcoidosis patients and 31 isolates from patients with tuberculosis. Fifty percent of the isolates from sarcoidosis patients and 16.13% of the isolates from patients with tuberculosis were represented by a unique pattern, whereas the remaining isolates belonged to seven shared types. Interestingly, half of the isolates from sarcoidosis patients did not resemble the spoligotypes of the isolates from patients with tuberculosis, most of which pertained to shared spoligotypes.

Sarcoidosis is a granulomatous disorder of unknown cause that affects multiple organs. The cells of the granuloma are organized spatially as immune granulomas, the characteristic of sarcoidosis, as a result of an immunological response to an antigenic trigger. It has been suggested that infective agents, including mycobacteria, propionibacteria, parasites such as *Schistosoma*, and fungi such as *Coccidioides*, are likely triggers (but not as infection) in a genetically predisposed individual and that this initial event leads to the sarcoidosis granulomatous response (2). Other agents such as beryllium, zirconium, and aluminum can also trigger the granulomatous response (2). Most studies of a possible causal organism have focused on complex organisms, including *Mycobacterium tuberculosis* complex isolates from sarcoidosis patients (4). Nevertheless, the PCR techniques used did not provide a detailed differentiation among isolates of the *M. tuberculosis* complex. Since the discovery of polymorphic DNA in *M. tuberculosis* complex strains, strain differentiation has become a valuable tool in the study of the epidemiology of tuberculosis and perhaps of sarcoidosis as well. Spoligotyping is a typing method that is based on DNA polymorphism in the *M. tuberculosis* complex. The chromosomal region consisting of identical 36-bp direct repeats alternating with 35- to 41-bp unique spacer sequences is the target of the spoligotyping technique (5). Since spoligotyping is PCR-based, it can be performed directly with *M. tuberculosis* complex organisms, including those that are nonviable or found in tissues in paraffin-embedded blocks or in archeological samples (13).

The objective of this study was to determine the molecular epidemiology of sarcoidosis in Greece by spoligotyping. We wanted to determine whether *M. tuberculosis* complex isolates from patients with sarcoidosis have spoligotypes that are common in the world and similar to those isolated from Greek patients with tuberculosis or if they have specific spoligotypes.

A total of 33 formalin-fixed paraffin-embedded tissues (lung and lymph node tissues) and 7 bronchoalveolar lavage samples, previously determined to be *IS6110* positive (4), from Greek patients suffering from sarcoidosis (S1 to S40) were subjected to molecular analysis. In addition, formalin-fixed paraffin-embedded material (lung tissue) from 32 Greek patients with tuberculosis (T1 to T32) was analyzed in parallel as controls. DNA extraction was performed as previously described (4). To confirm the integrity of the DNAs, a 430-bp sequence of the human glyceraldehyde-3-phosphate dehydrogenase gene was amplified.

The DNAs were subjected to direct repeat-specific PCR and hybridization with membrane-bound spacer oligonucleotides and a spoligotyping kit (ISOGEN Bioscience, Maarsen, The Netherlands) according to the manufacturer’s instructions. The spoligotyping results were entered in a binary format as Excel spreadsheets and were compared to the World Spoligotype Database spoIDB3.0 (11). At the time of the matching analysis, spoIDB3.0 contained 13,008 patterns distributed into 813 shared types and 1,300 orphan patterns from >90 countries (11). An isolate was assigned a shared type if the same spoligotype was found for isolates obtained from two or more patients in the world. If no matching spoligotype was identified in the database, the isolate was defined as unique. All of the identified spoligotypes have been submitted to the World Spo-
ligotype Database (http://www.pasteur-guadeloupe.fr/tb/spoldb3). A dendrogram was generated using Bionumerics V4.0 (Applied Maths, Sint-Martin-Latem, Belgium) with the UPGMA Dice comparison setting, an optimization of 2%, and a position tolerance of 1%.

Thirty-eight samples from patients with sarcoidosis and 31 samples from patients with tuberculosis had intact DNA for spoligotyping. A total of 24 distinct spoligotypes were obtained from the 38 samples from sarcoidosis patients, and 9 were obtained from the 31 patients with tuberculosis (Fig. 1). Nineteen (50%) isolates from patients with sarcoidosis were represented by unique patterns, whereas 19 (50%) isolates belonged to six shared types already defined in spoIDB3.0. Five (16.13%) isolates from patients with tuberculosis were represented by unique patterns, whereas 26 (83.87%) isolates were clustered in six shared types. The spoligotypes specific to Greece and the shared type designations of the isolates that matched with worldwide strains, together with their geographic distribution, are indicated in Table 1. All of the patients were epidemiologically unrelated and were born and live in Greece.

As indicated in Fig. 1, 18 isolates from patients with sarcoidosis exhibited similar patterns to those of certain strains from patients with tuberculosis. Only one major spoligotype was observed (type 53), which accounted for 28.95% of the total
isolates from sarcoidosis patients and for 35.48% of the total isolates from patients with tuberculosis. Additionally, it is noteworthy that no association between specific spoligotypes and particular tissue origins was observed.

This is the first report describing the molecular characterization of M. tuberculosis complex isolates from patients with sarcoidosis and tuberculosis in Greece. Almost 34.78% of the spoligotypes were found to be specific to Greece, and thus 50% of the sarcoidosis and tuberculosis in Greece. Almost 34.78% of the spoligotypes localized to Australia, 12). Although during the last few years Greece has accepted a number of communities throughout the world (1, 6, 7, 9, 10, 11). This complex isolates from patients with tuberculosis. Additionally, it is noteworthy that no association between specific spoligotypes and particular tissue origins was observed.

Collectively, our results point to M. tuberculosis complex isolate spoligotypes among Greek patients with sarcoidosis and tuberculosis. Unfortunately, no information is available from Greece, and there is limited information from neighboring countries, regarding the molecular epidemiology of tuberculosis and especially sarcoidosis to aid in explaining and classifying the spread of these spoligotypes. Since a significant number of the samples from patients with sarcoidosis are distributed in different and unique spoligotypes, the analysis of a greater number of samples could reveal spoligotypes specifically associated with sarcoidosis. Through that effort, the hypothesis of the involvement of some types of mycobacteria in the etiopathology of the disease could be tested more robustly. Similarly, the description of the genetic diversity of M. tuberculosis complex isolates from patients with sarcoidosis could be improved.

### References


