

RESEARCH ARTICLE

IDENTIFICATION OF ACTINOMYCETES IN THE BIODEGRADED WALL MATERIAL BY PCR AMPLIFIED GENE FRAGMENT CODING FOR 16S rRNA

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ABSTRACT

Generally, old buildings and monuments are often affected by biodecay. An inventory of the existing microbial consortia which associated with the damage part of the buildings wall is not yet integral part of the restoration process. This stems from the lack of effective means for such stocktaking. Although, Actinomycetes, fungi and bacteria cause severe damage through mechanical process from the growth into wall and its grounding and through their metabolism. Physiological and molecular approach based techniques were used in the detection and identification of DNA sequences encoding rRNA to actinomycetes present on building wall. DNA from isolate was extracted and 194 base pair fragments of the 16S rDNA were amplified with cultured primers. The 16S rDNA fragments of uniform length obtained from the different species were separated according to their sequences difference by PCR. By the sequencing excised and reamplified individual DNA bands, characterized the phylogenetic tree of corresponding actinomycetes. Using this approach, we identified members of the genera of *Streptomyces*, *Nocardia* and *Micromonospora*.

Key words: Actinomycetes, *Nocardia*, *Micromonospora*, *Streptomyces*

INTRODUCTION

It is well recognized that the microorganisms can be responsible for the destruction of building material and cultural heritages including the decay of concrete, marble and sandstone^{1,4}. Thus, it is not surprising that the building material can also support the growth of microorganisms, which are responsible for the biodegradation of building material³. Actinomycetes which, inter alia, grow on the surface of the building wall might discolour the cemented or painted surface but also by excreting metabolic products. Thus, the colonization of external surfaces of buildings by microorganisms cause the aesthetically unacceptable appearance of wall surfaces by biogenic pigment² and the production of extracellular polymeric substances that cause mechanical stress to the mineral structure due to shrinking and swelling cycles of these colloidal biogenic slimes inside the pore system. However, mycelia of the actinomycetes and fungi can penetrate into the paint layer and its grounding, resulting in the mechanical destruction of the cultural heritage and building.

DNA based techniques of identification of microorganisms have revealed that the conventional microbiological efforts to isolate actinomycetes present in building wall material. To obtain a better understanding of the microbial community and potential role of actinomycetes in the process of degradation of building material; we applied the molecular approach of PCR amplified gene fragments coding for 16S rRNA. The 16S rDNA fragments of DNA of actinomycetes, which could be isolate from building wall material by conventional microbiological techniques were sequenced and analysed phylogenetically. Our results demonstrate that the molecular biological approach is a potential tool to identify the actinomycetes in biodegradation process.

MATERIAL AND METHODS

Sample collection: The samples were collected from different sites of building wall material at Hardwar region. The samples were taken from an area which showed a rosy and dirty swelling cover of various intensities by scrapping off surface material and wall plaster to a depth of 1-5 mm. The total sample was split into two parts. One part was used for isolation of aerobic and heterotrophic actinomycetes by standard cultivation techniques and the second part used for scanning electron microscopy.

Electron microscopy: Particles of the samples were mounted with aluminium stubs and sputter coated to a 3 nm depth with gold platinum (80:20), using a magnetic sputter coater. Electron micrography was performed with a Hitachi S-4100 scanning electron microscope.

Isolation of actinomycetes: The sample material from biodegraded wall was suspended into saline solution and shaken for 15 minutes to remove the intensities from the particles⁵ and transferred to agar plates in dilution from 10⁻² to 10⁻⁶. The following media were used: actinomycetes agar, nutrient agar, nutrient broth, glycerol asparagine agar, streptomycetes agar. The plates were incubated at 28°C for 3-7 days. The number of CFU was determined every 3 days. Isolation of well defined single colonies resulted in the isolation of twenty nine isolates as pure culture and three isolates out of them further selected for identification by 16S rDNA sequencing.

DNA extraction: Genomic DNA from the isolated isolates of actinomycetes was obtained by chloroform-phenol extraction⁸. Isolates were dissolved in TE buffer and subjected to three freeze-thaw cycle to lyse the isolates⁵. After centrifugation, 5µl of the supernatant was used as DNA template in the PCR. The protocol for DNA

extraction including microwave heating, freeze thawing, lysis solution were tested.

Sequencing of 16S rDNA fragments: DNA extracted⁸ from isolates of actinomycetes and DNA eluted from the excised DGGE bands were reamplified with the same forward primer (5'-CCTACGGGAGGCAGCAG-3') including an additional sequence extension (5'-AAAATTAACCCTCACTAAAG-3') at its 5' end the reverse primer (5'-ATTACCGCGGCTGCTGG-3') with the additional sequence (5'-AAATTCACACAGGAAACAG-3') at its 5' end to facilitate DNA sequence analysis. The newly obtained PCR fragments were purified⁷ and sequenced directly with a LI-COR model 4000L automatic sequencing system.

Identification of actinomycetes by comparative sequence analysis: The sequence obtained from the isolated actinomycetes and DGGE bands were sent to the ribosomal database project⁸ and to the EMBL nucleotide sequence database for comparison. The similarity rank tool of the RDP and the FASTA search option for the EMBL database were used to search for close evolutionary relatives. Prealigned sequences of representatives of several groups were obtained and used for the identification of similarities to test sequences.

RESULTS AND DISCUSSION

Electron microscopy of biodegraded wall material

The electron microscopy of a biodegraded wall sample, which was taken from the surface, revealed a dense microbial colonization of the material (Figure 1). The dominant colonizing microorganisms showed filamentous morphologies ranging in length from 4 to 15 μ m. The filaments not only were attached to the material but above grew with in the inter sticks of the material occasionally forming tuberous growths. Figure 2 suggests that the growth of these microorganisms resulted in a loosening of the consistency of the ground material.



Figure 1: Microbial activities on domestic building at Haridwar (Uttarakhand)

Isolation of actinomycetes from building wall: As all the isolation process twenty nine isolates of actinomycetes were isolated. On the basis of morphological, physiological characteristic it was said to be isolates belong to actinomycetes family and the all isolates were classified into three groups. Three isolates (A_{11} , A_{20} , A_{27}) which shows better antimicrobial potential, were chosen for further study.

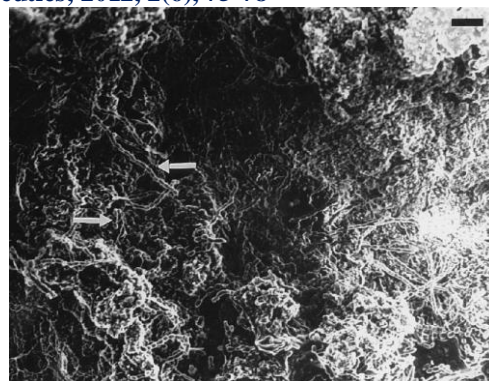


Figure 2: Electron micrograph of surface material of building sample. The Filamentous attached to the material are indicated by arrow.

Phylogentic classification of isolates of actinomycetes:

The alignment of the nucleotide sequence (749 base pair) of actinomycetes isolates were done through matching with the 16S rRNA reported gene sequence in the gene bank. The data base of national centre for biotechnological information (NCBI) basic local alignment search tool (BLAST) available at (www.ncbi.nlm.nih.gov) was used to compare the isolate strains with sequence of the reference species of the isolates A_{11} , A_{20} and A_{27} contained in genomic database bank and produced a ethidium bromide stained gel separation pattern of actinobacteria DNA fragment coding for 16S r RNA (Figure 3). The Figure 3 represents the ethidium bromide-stained DGGE separation pattern of bacterial DNA fragment coding for the 16S rRNA. Lanes 1 to 4 and 7 to 11 show the fragments obtained from actinomycetes isolates A_{11} (lane 2, 3 and 7), A_{20} (lanes 1, 8 and 10), A_{27} (lanes 4, 9 and 11), and (lanes 5 and 6, representing control DNA. A weak band was visible only in the original gel. The position of the band is marked.). Lanes 5

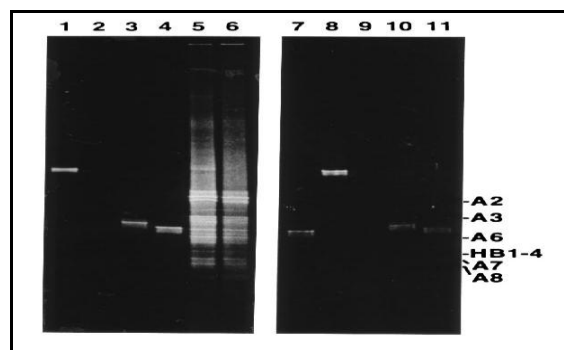


Figure 3: Gel Image of 16S rDNA amplicon

and 6 show the reproduced band patterns of the 16S rRNA genes which were amplified directly from DNA. Bands A_2 , A_3 , A_6 , A_7 , and A_8 were excised and sequenced.

The result exhibited similarity 91% with 10 strains from the uncultured known species of *Streptomyces*, *Nocardia* and *Micromonospora*. The 16S rRNA nucleotide sequence of *Streptomyces*, *Nocardia* and *Micromonospora* consisted of 749 bp each and the G-C content was 70%. These results were in accordance to many authors who mentioned that the G-C content of the actinomycetes DNA is 69-78%⁷. Modern identification system are based on 16S rRNA sequence data, which have provided valuable information about test isolates (A_{11} , A_{20} and A_{27})

systematically and then have been used to identify several new isolates¹⁰. For data analysis, the phylogenetic tree in

Figure 4 was derived from the distance matrices using neighbor joining method¹².

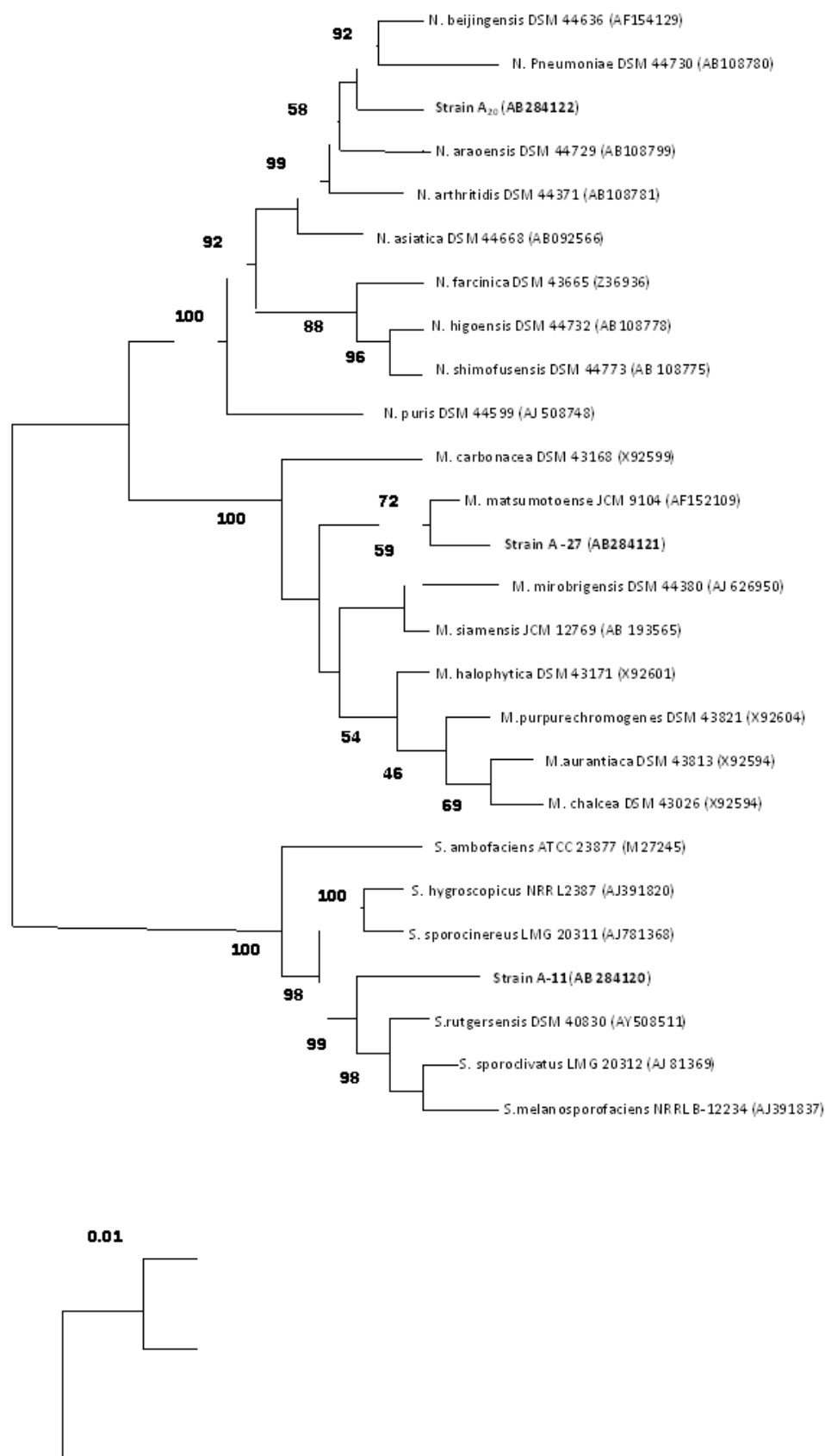


Figure 4: Phylogenetic tree (evolutionary relationships of 26 taxa)

A phylogenetic tree was constructed based on a 1216 bp alignment of the cloned PCR amplicons and known actinomycete sequences (Figure 4). The actinomycete 16S rDNA sequences amplified from DNA isolated from isolates of building walls affiliated with several clusters in the phylogenetic tree. The clones affiliated with three branches, one including *Nocardia beijingensis*, *Nocardia araoensis* and *Nocardia puris* second include the *Micromonospora carbonacea*, *Micromonospora chalcea* and the third including *Streptomyces ambofaciens*, *Streptomyces rutgersensis* and *Streptomyces melanosporefaciens*. Other sequevars were also present; some of them being distantly related to sequences obtained from database. The isolate A₁₁ closely related to *Streptomyces rutgersensis* and *Streptomyces sporocinereus*, isolate A₂₇ closely linked to *Micromonospora matsumotoense*, *Micromonospora siamensis* and *Micromonospora mirobrigensis* and the third isolate linked to the known species of *Nocardia araoensis*, *Nocardia pneumonia* and *Nocardia beijingensis*. Based on maximum identity score first ten sequences were selected and aligned using multiple alignment software program Clustal W. Distance matrix was generated using RDP database and phylogenetic tree was constructed using MEGA4. Therefore, on the basis of sequences producing significant alignment isolates A₁₁, A₂₀ and A₂₇ were suggesting to be new genomic *Streptomyces*, *Nocardia* and *Micromonospora* species and their accession number as under:

Strain A₁₁: accession number (AB284120) strain is *Streptomyces* species.

Strain A₂₀: accession number (AB284122) strain is *Nocardia* species.

Strain A₂₇: accession number (AB284121) strain is *Micromonospora* species.

CONCLUSION

Actinomycetes contribute to the mechanical destruction of wall materials as a result of the production of mycelia, which grow into the building material and its grounding.

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- Sequencing of more bands from the gene sequences pattern to explore the microbial diversity in more detail,
- Attempting to isolate bacteria related to PCR based 16S rDNA sequences such as anaerobic and halotolerant bacteria, and
- Determining the abundance of particular actinobacterial groups which have been found either by microbial isolation or by PCR based analysis¹¹.

It is hoped that an integrated approach of molecular and microbiological techniques will lead to a better understanding of the composition of bacterial species on ancient wall materials and to new ways to restore and conserve this cultural heritage.

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