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Exploration of Deinococcus-Thermus molecular diversity by novel group-specific PCR primers

Nicolas Theodorakopoulos1,2,3,4, Dipankar Bachar5,6, Richard Christen5,6, Karine Alain7 & Virginie Chapon1,2,3

1CEA, DSV, IBEB, SBVME, LIPM, F-13108 Saint-Paul-lez-Durance, France
2CNRS, UMR 7265, F-13108 Saint-Paul-lez-Durance, France
3Université d’Aix-Marseille, F-13108 Saint-Paul-lez-Durance, France
4IRSN, PRP-ENV, SERIS, L2BT, F-13115 Saint Paul-lez-Durance, France
5Université de Nice-Sophia Antipolis, UMR 7138, Systématique, Adaptation, Evolution, Parc Valrose, BP71, F-06108 Nice Cedex 02, France
6CNRS, UMR 7138, Systématique, Adaptation, Evolution, Parc Valrose, BP71, F-06108 Nice Cedex 02, France
7CNRS, IUEM – UMR 6197, Laboratoire de Microbiologie des Environnements Extrêmes (LMEE), Place Nicolas Copernic, F-29280 Plouzanne, France

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Deinococcus-Thermus, group-specific primers, molecular diversity.

Abstract
The deeply branching Deinococcus-Thermus lineage is recognized as one of the most extremophilic phylum of bacteria. In previous studies, the presence of Deinococcus-related bacteria in the hot arid Tunisian desert of Tataouine was demonstrated through combined molecular and culture-based approaches. Similarly, Thermus-related bacteria have been detected in Tunisian geothermal springs. The present work was conducted to explore the molecular diversity within the Deinococcus-Thermus phylum in these extreme environments. A set of specific primers was designed in silico on the basis of 16S rRNA gene sequences, validated for the specific detection of reference strains, and used for the polymerase chain reaction (PCR) amplification of metagenomic DNA retrieved from the Tataouine desert sand and Tunisian hot spring water samples. These analyses have revealed the presence of previously undescribed Deinococcus-Thermus bacterial sequences within these extreme environments. The primers designed in this study thus represent a powerful tool for the rapid detection of Deinococcus-Thermus in environmental samples and could also be applicable to clarify the biogeography of the Deinococcus-Thermus phylum.

Introduction
The phylum Deinococcus-Thermus is currently divided into the orders Deinococcales and Thermales. While the order Thermales encompasses five genera (Thermus, Meiothermus, Marinithermus, Oceanithermus, and Vulcanaithermus), the order Deinococcales is composed of three genera (Deinococcus, Deinobacterium, and Truepera). Deinococcus-Thermus is recognized as one of the most extremophilic phylum of bacteria. Cultured representatives of Thermus are either thermophilic or hyperthermophilic (Brock and Freeze 1969; Bjornsdottir et al. 2009; Zhang et al. 2010; Vajna et al. 2012), while Deinococcus strains exhibit resistance to extreme ionizing and ultraviolet radiations, desiccation, and other DNA damaging conditions (Rainey et al. 1997; Albuquerque et al. 2005; Cox and Battista 2005; Slade and Radman 2011). As these microorganisms or their cellular components are of biotechnological interest with potential applications in bioremediation or molecular biology (e.g., thermostable enzymes), much research has been focused on this particular group of prokaryotes. Deino-
**Material and Methods**

**Bacterial strains, culture conditions, and genomic DNA purification**

The bacterial strains used in this study are listed in Table 1. Biomass for the genomic DNA extraction was prepared by growing *Deinococcus* isolates in TGY medium (0.5% tryptone, 0.1% glucose, 0.3% yeast extract) at 30°C; *Escherichia coli* and *Pseudomonas aeruginosa* were cultivated in Luria-Bertani (LB) medium (1% tryptone, 0.5% yeast extract, 1% NaCl) at 37°C; *Shewanella oneidensis* was grown in LB medium at 30°C; all other isolates were grown in 0.1× Tryptic Soy Broth (DIFCO laboratories, Detroit, MI) at 30°C. Bacteria were harvested by 5 min centrifugation at 10000g, and DNA extractions were performed on the cell pellet with the DNeasy Blood & Tissue kit (Qiagen, Hilden, Germany), according to manufacturer's instructions. *Magnetospirillum magnetica* genomic DNA was provided by J. B. Rioux (IEEB-LBC-CEA Cadarache, France).

**Environmental DNA**

Sample collection and environmental DNA extraction for the sample from the Tataouine desert (located ~100 km west from the Sahara border; see Fig. 1) are reported in Chanal et al. (2006). *Deinococcus* have been detected therein by a combination of molecular and cultural methods. The present study was performed with purified DNA stored at −20°C.

Five water samples from Tunisian geothermal springs, described in Sayeh et al. (2010), were examined in this study: Ain Essalhine (spring 5), Nefta (spring 7), Hammem Bouhlel (spring 9), El Hamma of Tozeur (spring 10), and Ain Atrous (spring 11) (see Fig. 1). Springs 5, 9, and 11 were chosen as control samples as *Deinococcus* taxa in the deserts of Tataouine, Tunisia (Chanal et al. 2006), as well as the presence of *Thermus* taxa in Tunisian geothermal springs (Sayeh et al. 2010). However, the 16S rDNA clone libraries constructed from these studies are insufficient for a thorough exploration of *Deinococcus-Thermus* diversity. Indeed, sequences belonging to these taxa represent less than 2% of the overall community detected in Tataouine, and less than 10% of the overall revealed community in the Tunisian hot springs. An alternative approach to investigate the diversity of a taxonomic group in greater detail is to use group-specific primers specifically targeting a given taxon. This has been a successful strategy for the detection of diverse groups of Bacteria and Archaea at different taxonomic levels, including *Actinobacteria* (Stach et al. 2003); *Bacteroidetes*, *Planctomycetes*, *Firmicutes*, *Cyanobacteria*, α-, β-, and γ-proteobacteria (Mühling et al. 2008); *Korarchaeota* (Auchtung et al. 2011); *Acidobacteria* (Lee and Cho 2011; Gans et al. 2012); *Pseudomonas* (Widmer et al. 1998); and *Francisella* (Duodu et al. 2012). Furthermore, a specific primer for hemi-nested polymerase chain reaction (PCR) that targets the genus *Deinococcus* has recently been described (Chaturvedi and Archana 2012).

Here, we have developed specific PCR primers that target the 16S rRNA gene sequence of the entire *Deinococcus-Thermus* phylum. Following the initial in silico design step, primers specificity was tested with a collection of reference strains. Finally, we constructed 16S rDNA clone libraries to validate the use of these primers with environmental DNA. This approach has enabled us to detect novel representatives of *Deinococcus* and *Thermus* in desert sand samples from Tataouine, as well as in Tunisian geothermal spring water.

**In silico design of *Deinococcus-Thermus* specific primers**

All *Deinococcus-Thermus* sequences were extracted from the SILVA 111 reference sequences and aligned using the Muscle program. From this, a 90% sequence consensus was computed, and encoded using the IUPAC notation.
A 15 nucleotide (nt) sliding window was used to extract each subsequence containing less than three degeneracies. The overlapping extracted 15 nt oligomers were then recombined into longer domains. Each possible primer with a length of 20–30 nts containing less than three degeneracies was then extracted from the domains. A specific program, written in C, was developed to test which sequences would be recognized in the SILVA database, allowing up to three mismatches. Primers were then selected having a high specificity and a wide coverage for sequences of the Deinococcus-Thermus clade. A set of Python programs was used to improve their coverage while remaining specific. At the end of this process, the Deino-f-326-350/Deino-r-758-785 primers displayed good coverage and specificity, and we selected them for in vivo validation.

### PCR amplification and construction of 16S rRNA gene libraries

For the in vitro validation of primers that specifically target Deinococcus-Thermus, an initial set of experiments was conducted.
performed using phylogenetically diverse bacteria (Table 1). Genomic DNA from pure cultures was used as a template for PCR amplification with the primers Deino-f-326-350 (5′-CGGGAGGCAGCAGTTAGGAATCTTC-3′) and Deino-r-758-785 (5′-GTTAGGGYGTGGACTACCGGGGTATCT-3′). Each amplification reaction mixture (50 μL) contained 1× PCR buffer, 2 mmol/L MgCl₂, 0.2 mmol/L of each dNTP, 1 μmol/L of each primer, 1U Go Taq® Hot start polymerase (Promega), and 75 ng of DNA template. Based on the nucleotide content of the primers, the annealing temperature was predicted to be 64.5°C for Deino-f-326-350 and 65°C for Deino-r-758-785. However, when PCR amplification was performed with an annealing temperature of 65°C, all Deinococcus- Thermus strains DNAs were successfully amplified while no amplification was detected for strains belonging to any other phyla, with the exception of Firmicutes. Amplification of Firmicutes was predicted by in silico analysis, and can be accounted for by two primer/template mismatches. To avoid nonspecific amplification of Firmicutes, the PCR protocol was optimized by increasing the annealing temperature to 72°C. After an initial 2 min denaturation step at 94°C, 25 cycles were performed (94°C for 30 sec, 72°C for 1 min 45 sec), followed by a final extension step at 72°C for 5 min.

In a second set of experiments, primer efficiency and specificity for environmental DNA were examined by nested PCR. First, community DNA were used as targets for PCR amplification of the 16S rRNA genes with the universal primers fD1/S17, as described in Chanal et al. (2006). This primer set recognized 39.6, 82, and 88.5% of the 139 eligible sequences of Deinococcus-Thermus by allowing, respectively, 0, 1, or 2 mismatches between primers and sequences. The PCR products were then purified and used as the targets in a second PCR amplification with the primers Deino-f-326-350/Deino-r-758-785. The resulting PCR products were purified, cloned into the pCR2.1-TOPO vector (TOPO TA Cloning kit; Invitrogen, Carlsbad, CA) according to the manufacturer’s instructions, and electro-transformed into E. coli DH5α cells. Single colonies containing inserts were randomly selected and plasmids were extracted using the QIaprep Spin Miniprep Kit (Qiagen), according to manufacturer’s instructions. Plasmids were sent to GATC (Germany) for Sanger sequencing, using the M13F sequencing primer. Sequence quality was ensured manually with the BioEdit Version 7.0.5.3 (Hall 1999).

**Phylogenetic analyses**

To assign taxonomy, the SILVA 111 reference sequence database was downloaded and used to search each clone sequence using a Needleman–Wunsch algorithm, applying a 80% similarity cutoff to retrieve the 20 most similar sequences. Five clone sequences had no close relative in the SILVA database. These five sequences were then submitted to a Blastn query (excluding environmental sequences) using the NCBI nr database. First, we looked for hits with ≥99% similarity, and we calculated a consensus taxonomy. In the event that no hit was found with ≥99% similarity, the threshold was successively lowered in a step-wise fashion to determine at what level a taxonomy could be assigned. This process was repeated until an 80% threshold was reached. A consensus taxonomy corresponded, for example, to a defined genus if all selected hits shared the same genus.

For each of the 142 clone sequence, the two most similar sequences from the SILVA 111 reference database were selected (but with filtering to include at least one cultured bacterial sequence) to create a file of clone sequences and

![Figure 1](image_url). Location map of the study sites (triangles). For the hot springs, the water temperature is indicated in brackets. Genera belonging to the Deinococcus-Thermus phylum retrieved in each site are also indicated.
reference sequences (194 sequences in total). SeaView (Gouy et al. 2010) was used to align these sequences, using the included Muscle program. An initial tree was built from conserved domains, and sequences were reordered as they occurred in this tree using SeaView’s tools. Alignments were checked and manually modified when necessary. This process was repeated until no problem was detected. Trees were built using Neighbor-Joining (with distances corrected using the Kimura 2-parameter method), as implemented in SeaView with 1000 bootstrap replications. Trees were plotted with TreeDyn (Chevenet et al. 2006). To determine OTUs (operational taxonomic units), sequences were pair-wise aligned by a Needleman–Wunsch algorithm, a distance matrix was computed and sequences were clustered by average linkage from 85% to 100% similarity. We used clustering with a similarity of 97% in this study.

Results and Discussion

In silico analysis of primer pair specificity and coverage

The Deino-f-326-350/Deino-r-758-785 primers exhibited high coverage and specificity for the phylum Deinococcus-Thermus (Table 2). By allowing 0, 1, or 2 mismatches between primers and sequences, the primer set, respectively, recognized 89, 95.7, and 97.2% of the 1048 sequences of Deinococcus-Thermus. The coverage values were high for all genera within the phylum except for Marinithermus.

Among the 731,338 sequences in the SILVA database, three (out of 222,804) Proteobacteria sequences and two (out of 890) Aquificae sequences matched exactly with the primers. Allowing one mismatch between primers and sequences increased the number of matches, to 211 sequences from Candidate division TM7 (27.8% coverage value), 89 sequences from Chloroflexi (0.9% coverage value), 30 sequences from BD1–5 (7.6% coverage value), and 10 sequences from Proteobacteria.

The coverage values increased for several phyla, when two mismatches were tolerated between primers and sequences. However, for member-rich phyla (e.g., Proteobacteria), the coverage values remained low (0.1–2.4%) except for Firmicutes (25%). Elevated coverage values were obtained for a number of small phyla, including Candidate Divisions TM7 (39.2%), WS6 (22.2%), and KB1 (12.8%); BD1–5 (69.9%), WCHB1–60 (84.3%), MVP-21 (37.5%), and Kazan-3B-28 (15.4).

In vitro validation of primers with reference strains

Tests for primer specificity and optimization of PCR amplification conditions were performed on a set of 36 bacterial strains comprising representatives from the bacterial phyla Actinobacteria, Bacteroidetes, Deinococcus-Thermus, Firmicutes, and Proteobacteria (Table 1). When
PCR amplification was performed with an annealing temperature of 72°C, Deinococcus and Thermus DNAs were successfully amplified, resulting in an expected amplicon size of 460 bp. No amplification was detected for strains belonging to any other phyla (Fig. S1).

**Contribution of the Deino-f-326-350 and Deino-r-758-785 primers for the efficient detection of Deinococcus-Thermus in environmental DNA**

To validate the use of Deino-f-326-350 and Deino-r-758-785 primers for environmental ecology applications, primer specificity was examined using six environmental community DNAs extracted from two distinct environments: the Tataouine desert sand and Tunisian geothermal springs. A direct PCR using the Deino-f-326-350 and Deino-r-758-785 primers with a 72°C annealing temperature did not yield any product with our six environmental DNA samples. This could be related to the high annealing temperature used here, in combination with low amount of Deinococcus-Thermus derived DNA and large quantity of nonspecific DNA in template DNA sample. Therefore, we performed a nested PCR protocol: the universal bacterial primers F1 and S17 were used in a first round, and the resulting amplicons were targeted in the second round with the Deino-f-326-350 and Deino-r-758-785 primers (with an annealing temperature of 72°C). This protocol resulted in PCR products of the expected size for the six DNA samples tested.

Then, to check primer specificity, six gene libraries were constructed with the PCR products and 142 sequences were used for a phylogenetic analysis (Fig. S2); a simplified tree is shown in Figure 2. The sequences clustered into 33 OTUs at 97% sequence identity. The rarefaction curves plotted at this level did not reach an asymptote indicating the number of clones sequenced (Fig. S3). One hundred twenty-four out of 142 sequences were affiliated with the targeted phylum Deinococcus-Thermus, whereas 18 sequences were affiliated with nontargeted phyla such as Chloroflexi, Firmicutes, and the Candidate Divisions KB1, TM7, and OD1. Among these 18 sequences, 15 displayed low similarity values (<93%) to known 16S rRNA gene sequences and three were related to Firmicutes; the latter suggests either amplification linked to primer mismatches (that could not be avoided even with an annealing temperature of 72°C) or PCR-induced artifacts. In spite of this, these nonspecific reactions occurred with a low frequency (13%) and did not prevent the detection of the targeted sequences within complex environmental DNA samples.

**Molecular diversity of taxa affiliated with Deinococcus-Thermus in the desert of Tataouine**

All twenty-four 16S rRNA gene clone sequences from the desert of Tataouine were identified as belonging to Deinococcales, and could be divided into two groups: the first group affiliates with the genus Deinococcus and comprises 14 sequences clustered into three OTUs; the second group is affiliated with the genus Truepera and comprises 10 sequences clustered into two OTUs (Fig. 2).

Nine sequences from Deinococcales were grouped into OTU 9. Within this OTU, four sequences were closely related to Deinococcus desertii (with similarity values ranging from 98.3–98.7%), a bacterium previously isolated from the Tataouine desert sand (Chanal et al. 2006). By contrast, the other sequences were affiliated to Deinococcus hopiensis and D. navajovicensis (95.5–99.8% similarity), two radioreistant strains recovered from arid soils in the Sonoran hot desert (Rainey et al. 2005). Four sequences grouped into OTU 13 were not closely related to any cultured strains, and their closest neighbor (94.1–95.2% similarity) was a 16S rRNA gene sequence detected in a quartz hypolith from the Acatama desert (Lacap et al. 2011).

Ten sequences, clustered into OTUs 6 and 10, affiliated with Truepera, and exhibited 93–94% 16S rRNA gene sequence similarity with Truepera radiovictrix, the sole type strain of this genus, isolated from hot spring run-offs (Albuquerque et al. 2005). These sequences displayed higher similarity (98.4–100%) to uncultured bacteria sequences from diverse biotopes and notably from extreme environments, including: rock samples from the Black Canyon of the Chihuahuan desert (NM; Northrup et al. 2010); the Mars desert research station (UT; Direito et al. 2011); sunlight-exposed biofilms from Chernobyl (Ukraine; Ragon et al. 2011); and saline biological desert crusts (China; Li et al. 2013).

Thus, application of the Deino-f-326-350 and Deino-r-758-785 primers revealed sequences that were detected in our previous study (such as Deinococcus desertii), as well as several new sequences that have not previously been recovered. In particular, the presence of Truepera in this environment was unsuspected until now.

**Molecular diversity of taxa affiliated with Deinococcus-Thermus in Tunisian geothermal springs**

One hundred sequences belonging to the Deinococcus-Thermus group were recovered from hot springs 5, 7, 9, 10, and 11. These were marked by an uneven distribution among the five springs, as the 26 sequences derived from
Figure 2. Neighbor-Joining tree based on 16S rRNA sequences recovered from Tataouine (labelled with the prefix DTat) and from hot springs 5, 7, 9, 10 and 11 (labelled with the prefixes Dtun-S5, Dtun-S7, Dtun-S9, Dtun-S10 and Dtun-S11, respectively). The tree is simplified to include one sequence per OTU and per collection site, as well as the most similar sequences from public databases. The number of sequences within each OTU is indicated at the leaves (n = x). A complete tree is shown in Figure S2.
springs 5 and 7 all affiliated with *Thermales* while the 74 sequences derived from springs 9, 10, and 11 all affiliated with *Deinococcales* (Fig. 2). This result differs to some extent from that previously reported by Sayeh et al. (2010), in which *Thermales* sequences were identified in springs 5, 9, and 11, and no sequences were found to affiliate with *Deinococcales*. This discrepancy between the two studies could be explained by the fact that, for this study, new DNA extractions were performed from samples stored at −80°C, implicating that the DNA extracts were not exactly the same in both cases. The discrepancies could also be due to differences in DNA extraction procedures and PCR conditions (e.g., primer pair and/or amplification programs).

The 74 *Deinococcales* sequences recovered from springs 9, 10, and 11 all affiliated with the genus *Truepera*. Springs 10 and 11 displayed a very similar molecular diversity, whereas spring 9 had a distinct profile. Springs 10 and 11 were dominated by OTUs 11 and 14, representing, respectively, 93 and 96% of the sequences; these OTUs were also characterized by a lower than 90% similarity to cultured species. Sequences recovered from spring 9 also clustered into these two OTUs, but with much less abundance (three sequences). Sequences from OTU 14 were most closely related (95–99.8%) to sequences recovered from “hot” environments such as alkaline hot springs (JF935173, Papua New Guinea), arid soils of northwestern China (FR849462), and hot Calamita ferromagnetic sand (Perfumo et al. 2011). Sequences from OTU 11 were most closely related (95–96% similarity) to uncultured bacteria derived from an impressive number of saline and hypersaline environments, including: French Guiana coast mud (KCO10001); the northern subtropical Pacific Ocean (Eiler et al. 2011); hypersaline microbial mats (Mexico, JNS01803); north Pacific subtropical gyres (Pham et al. 2008); activated sludge from a seawater-processing wastewater treatment plant (Sánchez et al. 2011); the northeast subarctic Pacific Ocean (HHQ64210); coastal sediments of the Ariake Sea (Japan, AB560052); the Xiao Chaidan salt lake (China, HM128252); tailing material from Chanaral Bay (Acatama desert, HF558617); hypersaline sediments from Lake Kasin (Russia; Emmerich et al. 2012); marine sponge (Florida; Montalvo and Hill 2011); and the Sapelo Island salt marsh (GA, AY711411). Despite their great geographic separation, springs 9 and 10 (located near Tozeur, at the border of the “Chott el-Djerid” endorheic salt lake) and 11 (located near Korbous, on the Mediterranean Sea) each showed elevated salinity (3, 5, and 11 g/L, respectively). Taken together, these data strongly suggest that sequences from OTUs 11 and 14 belong to ubiquitous halophilic *Truepera* found in saline environments worldwide.

Nine OTUs related to *Truepera* (2, 4, 6, 12, 20, 25, 35, 47, and 50), and which represent 30 sequences, were detected exclusively in spring 9. Sequences belonging to the OTUs 2, 4, 12, 25, 47, and 50 were most closely related to uncultured organisms derived from marine environments, such as surfaces of marine macroalgae (GU451513; Lachnit et al. 2011) and marine sandy sediments from the North Sea (AM040122; Musat et al. 2006). Bacteria belonging to these OTUs may correspond to salt-tolerant species. This hypothesis is consistent with the location of spring 9 near the endorheic salt lake “Chott el-Djerid”. The OTU 6 encompassed six sequences that were also detected in spring 9, and which displayed close relationships with sequences detected in several extreme environments such as deserts and radioactive sites (see above). This OTU was singular in that it was additionally represented in the Tataouine desert sample; this phenomenon could be the signature of exchanges between the Tataouine site and spring 9, possibly occurring through the dust and sand storms that frequently occur in this region.

The 26 *Thermales* sequences derived from spring 5 (13 sequences) and spring 7 (13 sequences) all affiliated with the genus *Thermus* and clustered into two OTUs. OTU 1 was the most abundant and accounted for 77% of the sequences from spring 5, and 100% of the sequences from spring 7. Sequences from this OTU have no close cultured neighbor, and exhibit high similarity (96.1–97.0%) with a sequence detected in hot mineral soils (Antarctica; Soo et al. 2009). In addition, three sequences from spring 5 clustered into OTU 5 and were closely related to *Thermus thermophilus* (97.4–99.6% similarity) and *Thermus arcticus* (97.6% similarity), two thermophilic strains isolated from hot springs (Murzina et al. 1988; Zhang et al. 2010). As most *Deinococcus* species are mesophilic or moderately thermophilic, the absence of these bacteria from springs 5 and 7 could be explained by the elevated temperatures recorded at these sites (73°C and 70°C, respectively).

**Conclusions**

We have demonstrated that the primers developed for this study are highly specific and allow the detection of *Deinococcus-Thermus* sequences within environmental samples. They represent a powerful tool to detect novel *Deinococcus-Thermus* sequences through the sequencing of a limited number of clones, which will provide new insight into *Deinococcus-Thermus* molecular diversity in extreme environments. In line with this, our results indicate the presence of previously undescribed salt-tolerant bacteria in three springs.

These primers could be used in ecological studies for a rapid screening of environmental DNA samples, and could also be applicable to clarify the biogeography of the *Deinococcus-Thermus* phylum.
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Conflict of Interest

None declared.

References


Supporting Information

Additional Supporting Information may be found in the online version of this article:

Figure S1. PCR amplification using Deino-f-326-350 and Deino-r-758-785 primers and genomic DNA of diverse Deinococcus, Meiothermus, Thermus, Paenibacillus, Bacillus and Lysinibacillus strains.

Figure S2. Neighbor-Joining tree (1000 replicates) based on 16S rRNA gene sequences recovered from Tataouine (labelled with the prefix DTat) and from hot springs 5, 7, 9, 10 and 11 (labelled with the prefixes Dtun-S5, Dtun-S7, Dtun-S9, Dtun-S10 and Dtun-S11, respectively). The tree includes the most similar sequences from public databases.

Figure S3. Rarefaction curves of the OTUs at 97% sequence similarity.