Unexpected homology between inducible cell wall protein QID74 of filamentous fungi and BR3 salivary protein of the insect Chironomus

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ABSTRACT A gene, qid74, of mycoparasitic filamentous fungus Trichoderma harzianum and its allies encodes a cell wall protein that is induced by replacing glucose in the culture medium with chitin (simulated mycoparasitism conditions). Because no trace of this gene can be detected in related species such as Gibberella fujikuroi and Saccharomyces cerevisiae, the qid74 gene appears to have arisen de novo within the genus Trichoderma. Qid74 protein, 687 residues long, is now seen as highly conserved tandem repeats of the 59-residue-long unit. This unit itself, however, may have arisen as tandem repeats of the shorter 13-residue-long basic unit. Within the genus Trichoderma, the amino acid sequence of Qid74 proteins has been conserved in toto. The most striking is the fact that Qid74 shares 25.3% sequence identity with the carboxyl-terminal half of the 1,572-residue-long BR3 protein of the dipteran insect Chironomus tentans. BR3 protein is secreted by the salivary gland of each aquatic larva of Chironomus to form a tube to house itself. Furthermore, the consensus sequence derived from these 59-residue-long repeating units resembles those of epidermal growth factor-like domains found in divergent invertebrate and vertebrate proteins as to the positions of critical cysteine residues and homology of residues surrounding these cysteines.

When a distinctive body part is shared between certain organisms belonging to different phyla or classes, convergent evolution traditionally has been invoked as an explanation, implying unrelated mutations affecting different sets of gene loci resulted in the manifestation of that distinct body part. An oft-invoked example of convergent evolution has been compound eyes of insects versus singular eyes of vertebrates. This proved to be a wrong example, however, because development of all metazoan eyes recently has been interpreted as consequences of exon shufflings (5, 6). Nevertheless, how, by exon shufflings, the same domain can be propagated to be included in a number of unrelated proteins has never been made clear. If a particular protein donated the domain in question to the other by an exon shuffling, that donor itself had to lose that domain, in return receiving an unrelated domain.

In the present paper, we present Qid74 protein of a mycoparasitic filamentous fungus Trichoderma harzianum and show its sequence conservation within the genus Trichoderma. Then we show a surprising identity shared between this fungal Qid74 and the previously determined sequence of BR3 protein of the dipteran insect Chironomus, thus raising the possibility of convergent evolution at the protein level.

MATERIALS AND METHODS

Fungal Materials and Growth Conditions. Trichoderma harzianum CECT 2413, Trichoderma viride CECT 2423, Gliocladium virens CECT 2460 (reclassified as Trichoderma virens), Trichoderma longibrachiatum CECT 2606, Trichoderma koningii CECT 2412, Trichoderma reesi CECT2414, Gibberella fujikuroi CECT 2152, and Saccharomyces cerevisiae CECT1329 were obtained from the Colección Española de Cultivos Tipo, Burjasot, Valencia, Spain; T. harzianum IMI 206040 was obtained from the Imperial Mycological Institute, Kew, U.K. Hypocrea jecorina was a generous gift from C. Kubicek, University of Technology, Vienna. Glucose/agar/potato medium was used for the maintenance of cultures (7).

cDNA Library and Differential Screening. A cDNA library was constructed from mRNA isolated from 33-h cultures as described (8). The library was constructed in the Agt11 SfI-NotI vector by using the Riboclone kit (both from Promega). The library was screened by using first-strand cDNA synthesized on poly(A)+ mRNA populations as differential probes; they were obtained from control cultures (incubated with 10% glucose) and cultures induced with 1.5% chitin. The cDNA generated was labeled with [α-32P]dCTP (Amersham) by using a random primer-labeling kit from Boehringer Mannheim. The library was plated, transferred to nitrocellulose filters (Millipore), and hybridized as described (9). Plaques hybridizing only with cDNA from the induced culture were isolated and purified, and the cDNA was isolated according to the kit manufacturer’s instructions (Promega). The cDNA inserts were excised by digesting the recombinant phage DNA with EcoRI and NotI. These cDNAs were subcloned into pBlue-script SK+(Stratagene), obtaining the recombinant plasmid pQID74.

PCR. Total DNA from 4-day-old cultures was prepared according to Murray and Thomson (10) and was used as a
template for PCR by using, as a sense primer, the oligonucleotide Q74up, 5'-ATGTTGCTTAAAGCGTTCTGTTG-GCC-3' (nucleotides 1–28 of the ORF) and, as antisense primer, the oligonucleotide Q74ol1, 5'-ATGTTCTTTGCA-CAGCAGCTTGTGTTCCT-3' or Q74ol2, 5'-TCAAGGATGATCTCATTCAAGTGTTCTTCT-3' (2074–2100). To clone the 5' region inverse PCR was carried out with the oligonucleotides IQ74up, 5'-GCAAGAACTGCGC-CAAGCTTTGATGAT-3' (1832–1860) as sense primer and IQ74ol1, 5'-GCAAGAACTGCGC-CAAGCTTTGATGAT-3' as antisense primer. The amplification protocol used was as follows: 95°C, 1 min; 60°C, 30 sec; 72°C, 1–4 min. This standard cycle was repeated 35 times. PCR experiments were always carried out with PFU polymerase from Stratagene.

**DNA Sequencing and Data Analysis.** The recombinant plasmid (pQID74) was subjected to restriction analysis. Using the information obtained, constructions and nested deletions made from the original full cDNA were subjected to double-stranded DNA sequencing reactions by using the dideoxy chain termination method of Sanger et al. (11) and following the Sequenase protocol (United States Biochemical). Both strands were fully sequenced from overlapping constructs. The clones amplified by PCR were subcloned in pBluescript SK(−) (Stratagene), and nucleotide sequence was determined with an ALF automatic DNA sequencer from Pharmacia. Sequence data and predictions of protein structure were analyzed by using programs developed by the University of Wisconsin Genetics Computer Group (12). Other computer analyses were performed by using programs developed by Altschul et al. (13).

**RESULTS**

**Isolation of Genomic and cDNA Clones of Qid74 from T. harzianum.** As already noted, Qid74 protein is a cell wall component of *T. harzianum*, which is induced by replacing glucose in the culture medium with chitin as a sole carbon source (simulated mycoparasitism conditions). This permitted us to use a differential hybridization approach to single out cDNA copies of chitin-induced mRNAs. A Agt11-based cDNA expression library was prepared. Upon screening of approximately 2 × 10⁷ plaques, 20 clones were isolated, all of them showing a strong hybridization signal with the chitin-induced probe and no signal with the noninduced probe. One of the clones that showed the strongest hybridization signal was λQID74, which was further purified. DNA from λQID74 was isolated and sequenced as described in Materials and Methods. From the cDNA sequence, oligonucleotides were designed that allowed the isolation of the genomic DNA corresponding to the ORF region and the promoter region from *T. harzianum* and from other *Trichoderma* species. To isolate these genomic clones, a PCR approach was followed (see Material and Methods). Under the experimental conditions described, a band of ca. 2.1 kbp was initially amplified from *T. harzianum* 2413 chromosomal DNA. This band was then subcloned and sequenced.

**Sequence Analysis of qid74.** The complete nucleotide sequence for the QID74 cDNA and the genomic clone was determined. The most relevant characteristic of the sequence was its internal organization into tandemly repeating units. The genomic sequence of qid74 was identical to that of the QID74 cDNA, indicating a lack of introns in this gene. The full cDNA clone is 2,546 bp in length, with an ORF of 2,115 nucleotides. Such an ORF encoded a protein of 704 aa residues with a predicted molecular mass of 77,872 Da. The amino acid composition of the mature protein coded by qid74 was rather unusual, with a high proportion of lysine (13.5%), cysteine (12.8%), and glycine (9.6%). The codon usage of the *T. harzianum* qid74 gene resembles that of other *Trichoderma* genes that in the induced state demonstrated a very high transcription rate (14). Out of the 61 codons, 6 were not used at all whereas 24 were used 10 or more times. For example, of 76 glycine residues, 47 were encoded by the same triplet. With regard to serine, 40% of the residues also were encoded by the same triplet, in spite of the fact that there are six different serine codons available. The amino acid sequence of Qid74 also contains a potential signal peptide of 17 aa, which has an α-helical structure and is highly hydrophobic (15). This structure has been observed in secreted proteins such as cell wall proteins.

**Internal Organization of qid74.** A dot matrix analysis of the nucleotide or amino acid sequence of the qid74 gene indicated a number of regions that contained internally repetitious sequences. Multiple alignment of randomly generated fragments was carried out to identify the nature of apparent repetitiveness. Such an analysis showed internal repetitiveness in qid74 at two hierarchical levels: (hereafter) major and minor repeating units.

**Major repeating units.** The protein consisted of five incomplete and nine complete copies of the 59-residue-long unit. Thus, almost the entire ORF of the gene (from the residue position 46–680) can be divided into these tandemly repeated copies, as shown in Fig. 1. The pattern of sequence conservation among 59-residue-long units follows a pattern characteristic of other proteins with internal repeats (16). The units from the central region of the protein are more similar to one another as compared with those from the terminal regions. In fact, although the overall sequence identity was 60% among the repeats, units G and H demonstrated 97% identity with G, 75% with E, 61% with C, and only 41% with repeat A (Fig. 2).

The alignment of the sequence repeats indicates that the shorter amino acid sequences of units A, B, C, L, and M may be because of small deletions that had occurred in these copies. The observed tendency of copies in the central position remaining more conserved than those located in the periphery has been observed previously in all repeating DNA sequences, regardless of whether they are coding or noncoding (17). Yet, in the case of our Qid74, degeneracy most often affected the third base of codons, and even missense degeneracies were of the kind that led to conservative amino acid substitutions, e.g., glutamate to aspartate. Thus, degeneracy of repeating units in our case was under the strict surveillance by natural selection.

![Alignment of the deduced Qid74 amino acid sequence of the 14 repetitions that are 59 aa in length. The sequence of the repetitive regions have been aligned to achieve maximal homology by using the PILEUP and BESTFIT programs of the Wisconsin Gene Computer Group package.](image-url)
The qid74 gene apparently was conserved in toto within the genus Trichoderma, no trace of this gene was found in related fungi such as G. fugikuroi and S. cerevisiae (data not shown).

Qid74 and Homologous Proteins Found in Other Organisms. The BLAST program (13) has been used to perform a database search to find a protein or proteins that are reasonably homologous with Qid74. The maximal identity (p = 4.3e-44) with Qid74 was found in BR3 protein precursor of the

Cysteine is a highly reactive residue and has the tendency to form a disulfide bridge with another cysteine, thereby profoundly affecting a secondary structure of a protein. Not surprisingly, cysteine residues were conserved in fixed positions in all repeating units. Furthermore, they appeared in two motifs: CXC tripeptidic motif, where X is Val, Ser, or Ala and CXXC pentapeptidic motifs, which shall be discussed later (Figs. 1 and 2). Motifs related to the above also have been found in other cell wall proteins such as the ice nucleation protein (18).

Minor repeating units. In all repeating sequences, the larger repeating unit of today tends to be made of multiple copies of the smaller and more ancient repeating unit (19). This apparently was the case with Qid74. Observing Fig. 1, we note that the tridecapeptide KSKTCSCPQGNYW (in single-letter code) occupied the carboxyl-terminal position of the K copy. Furthermore, its apparent derivatives were found not only in the carboxyl half but also in the amino-terminal half of the C copy. In fact, two or more derivatives, at times truncated, of the above-noted tridecapeptide were found in nearly every copy of the 59-residue-long unit. Thus it would appear that the ultimate ancestor of the qid74 gene was tandem repeats of the 39-bp-long unit.

The qid74 Gene in Other Trichoderma Species. Southern blot and PCR analyses showed that the qid74 gene of the same ORF length was present in all the Trichoderma species tested. To determine a degree of sequence conservation among qid74 genes of various Trichoderma species, the study on a variation of tandemly repeating sequences was done. This analysis revealed almost the identical pattern shown by all the Trichoderma species tested. To verify the above finding based on PCR analyses by actual sequencing, a 1.1-kb band (Fig. 3C) from T. reesei and T. koningii and two 600-nt-long segments representing 5′ as well as 3′ ends of T. virens were sequenced. In addition, one 1-kb-long fragment corresponding to the upstream 5′ sequence of qid74 was cloned and sequenced on the strain T. harzianum 2413 and a species, T. reesei. To do so, inverse PCR using IQ74up and AQ74lo as sense and antisense primers was carried out.

As expected from PCR analyses, the ORFs of T. reesei, T. virens, and T. koningii indeed were identical with ORF of T. harzianum 2413. As to the 5′ upstream noncoding region, by contrast, 7 base substitutions per 600 bases separated T. harzianum 2413 from T. reesei. The promoter region apparently tolerated more base substitutions than the coding region. In addition to the conserved qid74 noted above, T. reesei was endowed with its redundant copy, which was rendered functionally defunct by an insertion of a 180-nt-long segment at the positions between 677 and 678. This pseudogene maintained only 65% identity with the real qid74 gene.

Although the qid74 gene apparently was conserved in toto within the genus Trichoderma, no trace of this gene was found in related fungi such as G. fugikuroi and S. cerevisiae (data not shown).
dipteran insect *Chironomus tentans* (accession number Q03376) and its homolog of *C. thumii* and *C. pallidivittatus*. BR3 protein is secreted in an enormous amount by salivary glands of an aquatic larvae of the midge *Chironomus* and it is used to weave a tube in which an aquatic larva houses itself. In preparations of *Chironomus* polytene salivary gland chromosomes, a gene encoding BR3 is readily recognized as an enlarged RNA puff (Balbiani Ring) reflecting its enormous transcriptional activity, and BR3 protein too is encoded by a coding sequence that is made of tandem repeats (6, 19). As shown in the alignments of Fig. 4, Qid74 and the carboxy-terminal half of BR3 demonstrated 25.3% sequence identity.

Other proteins with lesser homology included an unknown protein of *Caenorhabditis elegans* (p = 11e-16), fibrillins, Notch, and others, all of which contained repeating epidermal growth factor-like domains. Another reasonably homologous protein was an oocyst wall protein of *Chrysonodidium parvum* (accession number A48456). The homology score between Qid74 and this oocyst wall protein was p = 8.9e-13. All these proteins noted above had a few characteristics in common. As indicated by the presence of a signal peptide at each protein’s N-terminal region, these proteins were meant to be secreted by the cell, they were made of repeating domains rich in cysteine residues, and they were also rich in charged residues presenting similar hydropathic profiles.

**DISCUSSION**

A major problem is encountered whenever an attempt is made to classify various strains of deuteromycetal fungi, because there is an inherent uncertainty in deciding whether they are, in fact, different species belonging to the same genus or there is an inherent uncertainty in deciding whether they are, in fact, different species belonging to the same genus or they are, in fact, different species belonging to the same species. However, because we found no gene among these in *S. cerevisiae*, it would appear that this gene arose *de novo* in the ancestor of the teleomorphic genus *Hypocrea* whence the presently studied anamorphic genus *Trichoderma* sprung (21).

Let us now turn our attention to BR3 protein of the dipteran insect *Chironomus* with which Qid74 protein shared 25.3% sequence identity. Of the insect order Diptera, *Chironomus* together with mosquitoes belong to the suborder Nematocera, whereas the familiar *Drosophila* together with house flies belong to the other suborder Cyclorrhapha. Whereas larvae of the first suborder tend to lead aquatic lives, those of the second suborder are terrestrial. Indeed, no gene homologous to that encoding BR3 was found within the *Drosophila* genome (6, 19). Thus it would appear that BR3 gene also arose *de novo* either within the suborder Nematocera or much later within the genus *Chironomus*.

As indicated by the presence of a signal peptide at each protein’s N-terminal region, these proteins were meant to be secreted by the cell, they were made of repeating domains rich in cysteine residues, and they were also rich in charged residues presenting similar hydropathic profiles.

**FIG. 4.** Alignment between Qid74 (704 residues) and BR3 (1,700 residues). The alignment was carried out using the program sim, which is available in the Molecular Biology Server Expasy.
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