DNA sequence-based identification of *Fusarium*: Current status and future directions

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Received: 14 May 2015 / Accepted: 20 August 2015 / Published online: 26 September 2015
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Introduction

Fusarium ranks as one of the world's most economically destructive and species-rich groups of mycotoxigenic plant pathogens (Aoki *et al.* 2014). These ubiquitous molds produce a plethora of toxic secondary metabolites, such as trichothecenes, zearalenone, fumonisins,

and enniatins, which pose a significant threat to agricultural biosecurity, food safety, and plant, human and animal health (Marasas *et al.* 1984). Fusarial-induced diseases of virtually every economically important plant cost the global agricultural economy multi-billion euro losses annually. Moreover, phylogenetically diverse fusaria, including plant pathogens (Short *et al.* 2011), cause infections in humans, with those involving the cornea and nails being the most common (Chang *et al.* 2006 and references therein). Because fusaria are broadly resistant to the spectrum of antifungals currently available, disseminated infections in patients who are artificially immunosuppressed or immunocompromised and severely neutropenic are typically fatal (Balajee *et al.* 2009). The likely reservoir of nosocomial fusarioses is the plumbing system, which has been shown to harbor the most common human opportunistic fusaria (Kuchar 1996; Short *et al.* 2011). Accurate identification of the etiological and/or toxigenic agent is central to disease management and infection control (Wingfield *et al.* 2012). Thus, the primary focus of this mini-review is to provide a contemporary guide to the following three web-accessible resources for DNA sequence-based identification of *Fusarium*: FUSARIUM-ID (<http://isolate.fusariumdb.org/>; Geiser *et al.* 2004; Park *et al.* 2010), *Fusarium* MLST (<http://www.cbs.knaw.nl/fusarium/>; O'Donnell *et al.* 2010), and NCBI GenBank (<http://www.ncbi.nlm.nih.gov/>). The following brief overview of *Fusarium* phylogenetic diversity is provided as background information for the sections on DNA sequence-based identification.

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Phylogenetic limits, subgeneric clades and species recognition

Fusarium Link (Hypocreales, Nectriaceae) was circumscribed originally in 1809 based on the production of its iconic fusiform multiseptate macroconidia. However, we now know from morphological and molecular phylogenetic studies (Gräfenhan *et al.* 2011) that this character evolved convergently in different lineages of ascomycetes, and that it has been lost at least once within the *F. solani* species complex (e.g., *F. neocosmosporiellum* and related self-fertile species formerly classified in *Neocosmospora*; O'Donnell *et al.* 2013). As a result, *Fusarium* is non-monophyletic in the three mostly widely used morphology-based taxonomic treatments (Nelson *et al.* 1983; Gerlach and Nirenberg 1982; Leslie and Summerell 2006) that recognize 30, 78 and 70 species, respectively. With the exception of the turf grass pathogen '*Fusarium*' *nivale*, which is now recognized as *Microdochium nivale* within the distantly related order Xylariales (Samuels and Hallet 1983), and '*Fusarium*' *tabacinum*, which was reclassified as *Monographella cucumerina* in the Xylariales (Palm *et al.* 1995), the other taxa removed from *Fusarium* and reclassified in four different genera within the Nectriaceae by Gräfenhan *et al.* (2011) are not known to be pathogenic or mycotoxigenic. Fortunately, the molecular phylogenetic circumscription of a monophyletic *Fusarium* that includes at least 20 clades referred to as species complexes and nine monotypic lineages (Fig. 1) has received overwhelming support by the *Fusarium* research community worldwide (Geiser *et al.* 2013). Given the poor correspondence between the robust *RPB1/RPB2* molecular phylogeny and the morphology-based sectional classification adopted by Gerlach and Nirenberg (1982) and Nelson *et al.* (1983), the nonmonophyletic sections were abandoned in favor of strongly supported, monophyletic species complexes (Geiser *et al.* 2013; O'Donnell *et al.* 2013). Following the demise of dual nomenclature 1 January 2013 under the newly named International Code of Nomenclature for Algae, Fungi and Plants (Hawksworth *et al.* 2011), plant pathologists and other applied biologists are encouraged to only use the *Fusarium* name (Geiser *et al.* 2013). *Fusarium* has priority over the teleomorphs (i.e., *Albonectria*, *Cyanonectria*, *Gibberella*, *Haematonectria*, *Nectria* and *Neocosmospora*), and

the anamorphs *Bisfusarium* and *Rectifusarium*, which were recently proposed, respectively, for members of the *F. dimerum* (FDSC) and *F. ventricosum* (FVSC) species complexes (Lombard *et al.* 2015). Comparative phylogenomic analyses of low-coverage genomes of the 93 fusaria included in O'Donnell *et al.* (2013) are in progress to more critically evaluate the circumscription of *Fusarium* (J. Stajich, pers. commun.).

Although *Fusarium* as circumscribed morphologically by Gerlach and Nirenberg (1982) has shrunk in size by approximately 20 species, largely due to the seminal study by Gräfenhan *et al.* (2011), application of phylogenetic species recognition based on genealogical concordance and non-discordance over the past 20 years (GCPSR sensu Taylor *et al.* 2000; Dettman *et al.* 2003) has resulted in its explosive growth. Today at least 300 phylogenetically distinct species have been resolved as genealogically exclusive lineages based on phylogenetic analyses of representative fusaria in the ARS Culture Collection (NRRL), the CBS-KNAW Biodiversity Centre (CBS) and the Fusarium Research Center (FRC). However, the majority of these species are unnamed and many of these are morphologically cryptic (Fig. 1). Four complementary but distinct technological and theoretical advances have been key to the greatly accelerated species discovery within *Fusarium* over the past two decades. These include the marriage of PCR and automated DNA sequencing in the early 1990s, the acceptance of GCPSR-based studies as the gold standard for species recognition within the *Fusarium* community, the wealth of fusaria (~40,000 isolates) accessioned in publically accessible culture collections (e.g., CBS, FRC and NRRL), and a highly collaborative global phytopathological community.

Based on biodiversity studies that estimate the ~100,000 named and accepted fungi only comprise one-tenth (Hawksworth 2001) to one-fiftieth (Blackwell 2011) of the species in nature, coupled with the fact that most of the fusaria studied to date were isolated from agronomically important plants (i.e., <1% of all vascular plants; Simpson and Ogorzaly 1995) cultivated on a fraction of our planet's surface, it is reasonable to suggest that the number of novel fusaria in nature could exceed our current estimate of 300 by an order of magnitude. Looking to the future, typing schemes will need to take advantage of next-generation sequencing (NGS) technology (Boers *et al.* 2012) to rise to the challenge posed by the anticipated seismic shift in species discovery within *Fusarium* and

Fig. 1 Diagrammatic representation of *Fusarium* phylogeny inferred from a combined *RPB1* + *RPB2* dataset (3383 bp) rooted on sequences of *Neonectria* and *Ilyonectria* (modified from Fig. 1 in O'Donnell *et al.* 2013). GCPSR-based analyses indicate that the genus comprises at least 300 phylogenetically distinct species, 20 species complexes (highlighted in gray), and 9 monotypic lineages. The approximate number of phylogenetically distinct species within each species complex is indicated

other agriculturally important plant pathogens. This discovery will be driven in part by metagenomics studies (LeBlanc *et al.* 2014), surveys of endophytes inhabiting endemics in biogeographically interesting areas (Walsh *et al.* 2010), and by the inexorable introduction of novel pathogens into nonindigenous areas by the globalization of world trade (Fisher *et al.* 2012). Much is at stake because an accurate species-level identification is essential for elucidating and communicating all facets of a pathogens' biology (Wingfield *et al.* 2012).

Sequence-based identification of fusaria: Some loci reveal while others conceal

Only three of the marker loci tested to date meet three important criteria for phylogenetic species recognition in that they are: 1) applicable across the phylogenetic breadth of *Fusarium* (Fig. 1), 2) informative at or near the species-level, and 3) orthologous across the genus. These are: translation elongation factor 1- α (*TEF1*), DNA-directed RNA polymerase II largest (*RPB1*) and

second largest subunit (*RPB2*). The latter two marker loci were developed as part of the NSF-funded Deep Hyphae and the Assembling the Fungal Tree of Life (AFTOL) projects (Lutzoni *et al.* 2004; James *et al.* 2006). In contrast to *TEF1*, whose highly variable introns can only be aligned reliably across members of a species complex, or several closely related ones, the portions of *RPB1* and *RPB2* sequenced can easily be aligned across *Fusarium* (O'Donnell *et al.* 2013). While all three genes are included in our ongoing GCPSR-based studies that span the breadth of the genus, sequence data from only one of these loci is needed to obtain a reasonably accurate placement of an unknown within a species complex by conducting a nucleotide BLAST query of FUSARIUM-ID and/or *Fusarium* MLST (Fig. 2A–B), or by phylogenetic analysis (see Bruns *et al.* 1998 for an example). As discussed below, we also conduct BLASTn queries of GenBank, but these require careful scrutiny of the top ‘hits’ because: 1) many sequences in NCBI are misidentified (Bidartondo *et al.* 2008; Kang *et al.* 2010), 2) sequences deposited in NCBI for the majority of newly discovered fusaria employing GCPSR lack binomials, and 3) the taxonomy for many records is out of date (Fig. 3). Because most of the fusaria within the *F. solani* and *F. incarnatum-equiseti* species complexes lack binomials, an ad hoc species/multilocus haplotype nomenclature was adopted to allow for accurate communication of information regarding these pathogens within the scientific community (O'Donnell *et al.* 2008, 2009b).

At the suggestion of Christopher L. Schardl, University of Kentucky (Tsai *et al.* 1994), β -tubulin was the first protein-encoding gene that we used for molecular phylogenetics in *Fusarium* (O'Donnell and Cigelnik 1997; O'Donnell *et al.* 1998a). However, we discovered that its utility is limited due to the presence of divergent paralogs within the *F. solani*, *F. incarnatum-equiseti*, and *F. chlamydosporum* species complexes. After Stephen A. Rehner (ARS-USDA, Beltsville, MD) called our attention to the utility of *TEF1* for species-level studies in insects (Cho *et al.* 1995), we developed degenerate PCR primers that amplify the intron-rich 5' end of this ortholog in all fusaria (O'Donnell *et al.* 1998b), and we have used this sequence data to populate FUSARIUM-ID (Geiser *et al.* 2004; Park *et al.* 2010) and *Fusarium* MLST (O'Donnell *et al.* 2010). Although the internal transcribed spacer (ITS) region of the nuclear ribosomal DNA was selected as the official ‘barcode’ locus for the Fungi (Schoch *et al.* 2012), its utility within

Fusarium, and many other groups of fungi (Bruns 2001; Du *et al.* 2012; Gazis *et al.* 2011), is limited by the fact that it is often uninformative at the species-level, and like *TEF1*, it can only be aligned reliably across members of a species complex or several closely related ones. In recently evolved phytopathologically-important lineages such as the *F. graminearum* (Sarver *et al.* 2011), *F. oxysporum* (O'Donnell *et al.* 2009a) and *F. fujikuroi* species complexes (O'Donnell *et al.* 1998a), many of the species share identical or nearly ITS rDNA alleles. Using the ITS rDNA for species identification and phylogenetic inference in *Fusarium* is further complicated by the presence of highly divergent ITS2 rDNA paralogs or xenologs within every strain tested within six closely related species complexes (i.e., *concolor-to-fujikuroi* in Fig. 1; O'Donnell *et al.* 1998a). In addition, we recently detected this phenomenon within the *F. buharicum* species complex, which suggests that the gene duplication or horizontal gene transfer event took place at least 49 million years ago (O'Donnell *et al.* 2013).

ITS rDNA and domains D1 and D2 at the 5' end of the ribosomal large subunit (LSU rDNA) were used in GCPSR-based MLST schemes for the *F. solani* (O'Donnell *et al.* 2008), *F. dimerum* (Schroers *et al.* 2009), *F. incarnatum-equiseti* (FIESC) and *F. chlamydosporum* species complexes (O'Donnell *et al.* 2009b), but this locus consistently contributed the least number of phylogenetically informative characters among the loci sampled. For example, in the FIESC study, 162/717 (23%) of the aligned nucleotide positions in *TEF1* and 220/1766 (12%) in *RPB2* were parsimony informative; however, only 11/1125 (1%) were synapomorphic within the ITS+LSU rDNA (O'Donnell *et al.* 2009b). In hindsight, inclusion of the fungal ‘barcode’ locus in this typing scheme was a mistake. However, we should note that in the numerous instances where ITS+LSU rDNA sequence data did not resolve at the species level, it does have utility for placing unidentified fusaria within a species complex (Balajee *et al.* 2009).

Currently we are generating low-coverage genome sequences for the 93 fusaria included in our published molecular phylogeny of *Fusarium* (O'Donnell *et al.* 2013) to mine them for additional phylogenetically informative loci for species-level studies (López-Giráldez and Townsend 2011) and for comparative phylogenomics (Stajich, unpubl.). This and other whole genome sequence data will be critical for developing

NGS typing schemes necessary to characterize the deluge of novel species that will be discovered by phytopathological, biogeographical and metagenomic studies in the future. The newly developed marker loci should prove to be invaluable in expanding the ad hoc species-haplotype nomenclature (O'Donnell *et al.* 2008, 2009a, b) to all agriculturally and medically important fusaria so that information concerning the unnamed, morphologically cryptic mycotoxigenic plant and human pathogens can be accurately communicated within the scientific community. This informal naming system is also useful for identifying agriculturally and medically important species that should be formally described with Latin binomials.

Sequence-based identification of fusaria: A primer for conducting BLASTn queries via the Internet

Querying one of the web-accessible databases using partial DNA sequence data from *TEF1*, *RPB1* and/or *RPB2* to identify an unknown ordinarily is the easy part; however, interpreting the results is often challenging. As previously discussed in detail (Geiser *et al.* 2004;

O'Donnell *et al.* 2010), the advantage of conducting nucleotide BLAST queries of FUSARIUM-ID or *Fusarium MLST* first, rather than NCBI GenBank, is that they house broadly sampled, well-characterized phylogenetically informative sequences from isolates that can be obtained from FRC (<http://plantpath.psu.edu/facilities/fusarium-research-center>), the ARS Culture Collection (NRRL, <http://nrri.ncaur.usda.gov/cgi-bin/usda>) or the CBS-KNAW Biodiversity Centre (<http://www.cbs.knaw.nl/Fusarium/>).

When conducting BLASTn queries of the CBS-KNAW's *Fusarium MLST* database, we recommend only searching the reference files for *Fusarium*, which is the default setting, and setting the 'Minimum similarity to keep results' to 50% (Fig. 2B). Most of the sequences housed in FUSARIUM-ID are also present in *Fusarium MLST*, so a query of only one of these databases may be needed. Once the results of a BLASTn query of *Fusarium MLST* are returned, with the top 'hits' displayed in tabular form, click on Expand Alignments to see the alignment of your query sequence with each of the reference sequences retrieved from the database. Results obtained from BLASTn queries of GenBank and FUSARIUM-ID differ slightly in that

Fig. 2 (A) FUSARIUM-ID (<http://isolate.fusariumdb.org>) at Pennsylvania State University (Park *et al.* 2010) and (B) *Fusarium MLST* at the CBS-KNAW Fungal Biodiversity Centre (<http://www.cbs.knaw.nl/Fusarium/>) host two complementary web-accessible databases dedicated to the identification of fusaria via

nucleotide BLAST queries. Conducting the same BLASTn queries of GenBank (<http://blast.ncbi.nlm.nih.gov/Blast.cgi>) can be informative, but also challenging given the large number of misidentified sequences in NCBI (see Fig. 3)

horizontal red lines are used to identify sequence length of the top ‘hits’, which span the length of the region if the sequences are full-length (see % query cover in Fig. 3). GenBank differs from FUSARIUM-ID and *Fusarium* MLST in that all of the top BLASTn hits are displayed together in tabular form immediately below the horizontal red lines, followed by the alignment of your query sequence with each of the reference sequences that were recovered (Fig. 4). All three databases provide a hyperlink to each accession record, a description that includes the taxon name that the accession was deposited under and the locus. Also included are several statistical measures, with the two most important ones being percentage identity and query coverage. Using the loci mentioned herein, the E-value for each ‘hit’ should be zero (Fig. 3), which means the match is not due to chance.

After conducting a query of *Fusarium* MLST or FUSARIUM-ID, we frequently use the same nucleotide sequence to query GenBank, bearing in mind that many

sequences in NCBI are misidentified, others are deposited as *Fusarium* sp. without further annotation, and the taxonomy especially for older records may be out of date (e.g., GenBank accession AF178356.1 was deposited as *F. solani* f. sp. *glycines* in 1999, a year before this soybean pathogen was formally described as *F. virguliforme*). For those who opt to query GenBank, we recommend the use of *TEF1*, *RPB1* and/or *RPB2* sequences, rather than sequences from the ITS or LSU rDNA, and look for sequences obtained from NRRL strains among the top ‘hits’. The majority of the sequences we deposited in these databases were listed as *Fusarium* sp., because they represent unnamed species based on the results of several GCPSR-based studies (O’Donnell *et al.* 2010, 2014 and references therein). However, notes or comments were often included in the accession records to help identify the phylogenetic species and/or multilocus haplotype (Fig. 5; i.e., Haplotype=“FSSC 11-b” from O’Donnell *et al.* 2008). Another tip for correctly interpreting BLASTn results is

Sequences producing significant alignments:

Select: All None Selected:0

Alignments						Download	GenBank	Graphics	Distance tree of results	
	Description	Max score	Total score	Query cover	E value	Ident	Accession			
■	Fusarium equiseti isolate NFCCI 2157 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	878	878	100%	0.0	100%	JN014954.1			
□	Fusarium equiseti strain HP7 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	867	867	100%	0.0	99%	KJ677237.1			
■	Fusarium cf. equiseti MY-2011 isolate AM-28 18S small subunit ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	867	867	100%	0.0	99%	JN038469.1			
□	Fusarium cf. equiseti MY-2011 isolate AM-25 18S small subunit ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	867	867	100%	0.0	99%	JN038466.1			
□	Fusarium cf. equiseti MY-2011 isolate AM-24 18S small subunit ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	867	867	100%	0.0	99%	JN038465.1			
□	Fusarium equiseti genomic DNA containing 18S rRNA gene, ITS1, 5.8S rRNA gene, ITS2, 28S rRNA gene, isolate DBT-97	867	867	100%	0.0	99%	FR872728.1			
□	Fusarium sp. 91D2/F 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	867	867	100%	0.0	99%	GQ407102.1			
□	Fusarium equiseti 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1 and 5.8S ribosomal RNA gene, complete sequence; and internal transcribed spacer 2, partial sequence	863	863	99%	0.0	99%	KM273261.1			
■	Fusarium culmorum strain SX2 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1 and 5.8S ribosomal RNA gene, complete sequence; and internal transcribed spacer 2, partial sequence	863	863	99%	0.0	99%	KC329615.1			
□	Fusarium equiseti strain Fe1 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	863	863	99%	0.0	99%	HQ718414.1			
□	Fusarium equiseti isolate 2F internal transcribed spacer 1, partial sequence; 5.8S ribosomal RNA gene and internal transcribed spacer 2, complete sequence; and 28S ribosomal RNA gene, partial sequence	861	861	100%	0.0	99%	KM580656.1			
■	Fusarium sacchari culture-collection NCCPF-580033 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	861	861	100%	0.0	99%	KM921664.1			
□	Fusarium equiseti isolate USM3-2A 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	861	861	100%	0.0	99%	KM111481.1			
■	Uncultured fungus clone FA2-O9 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	861	861	100%	0.0	99%	JX984757.1			
★	Fusarium sp. NRRL 22244 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	861	861	100%	0.0	99%	GQ505685.1			
■	Hypocreales sp. LM512 18S ribosomal RNA gene, partial sequence	861	861	100%	0.0	99%	EF060807.1			
■	Fusarium chlamydosporum 18S rRNA gene (partial), 5.8S rRNA gene, 28S rRNA gene (partial), ITS1 and ITS2, strain WM 04.494	861	861	100%	0.0	99%	AJ853773.1			
■	Fusarium incarnatum voucher NJM 0177 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	861	861	100%	0.0	99%	AY633745.1			
■	Fusarium tricinctum genomic DNA containing 18S rRNA gene, ITS1, 5.8S rRNA gene, ITS2 and 28S rRNA gene, isolate DBT-43Amravati	861	861	100%	0.0	99%	FR851237.1			
■	Septogloeum mori strain EUF10 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	859	859	100%	0.0	99%	HQ339998.1			
■	Fusarium oxysporum isolate A2S3-D1 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S rDNA	856	856	100%	0.0	99%	KJ767070.1			

Fig. 3 Twenty-one of the 100 BLASTn ‘hits’ that were recovered from GenBank using accession JN014954.1 as the query. Black boxes were added to the GenBank query results to identify sequences deposited under 11 different names. The black star inserted

by the record for *Fusarium* sp. NRRL 22244 indicates the accession was identified in the notes/comments as phylogenetic species FIESC 25 (O’Donnell *et al.* 2009b)

Fig. 4 One of the 100 BLASTn nucleotide alignments that followed the tabular display shown in Fig. 3. Alignment of ITS rDNA GenBank accession JN014954.1 used as the query and GQ505685.1 NRRL 22244 FIESC 25-a strongly suggests that the

query sequence contains three sequencing errors (indicated by arrows). Alignments such as this are ideal for identifying nucleotide positions within chromatograms that should be checked for errors before sequences are analyzed and deposited in NCBI

to check the taxon names. Identical or nearly identical sequences deposited under multiple names, as illustrated in Figs. 3 and 6, should raise suspicions that some sequences are misidentified. In addition, sorting through the display of BLASTn ‘hits’ in some searches is complicated by the legacy of dual nomenclature (e.g., *Gibberella zeae* and *F. graminearum* for the same species).

We don’t recommend using sequences from the nuclear ITS rDNA and/or domains D1+D2 of the LSU rDNA from an unknown to query GenBank because 50% or more of the *Fusarium* sequences from this locus are misidentified in NCBI. Besides, as previously mentioned, ITS+LSU rDNA sequences are too conserved to resolve species limits of most fusaria. The 10 steps outlined below are recommended to increase the likelihood of obtaining an accurate species- or species complex-level identification of an unknown *Fusarium* (also see Fig. 7). Here, we used a study by Suthar and Bhatt (2011) to illustrate common mistakes/errors associated with species identification based on BLASTn queries and how to avoid them. They deposited their ITS rDNA sequence of a putative cumin wilt pathogen (NFCCI 2157) in GenBank incorrectly as *F. equiseti* (accession JN014954.1). Of the 100 ITS rDNA sequences that were recovered from GenBank, using their

accession JN014954.1 as the BLASTn query, we determined that all 48 sequences deposited with binomials were incorrectly identified to the species level. This finding highlights the importance of carefully inspecting the name attached to each of the records (Fig. 3) and consulting the primary literature (O’Donnell *et al.* 2009b). This simple but critical step would have revealed to these authors that their queries of GenBank and FUSARIUM-ID yielded contradictory results. A cursory inspection of the 48 named accessions retrieved by the BLASTn query of GenBank revealed that they were incorrectly deposited under seven different species names (Fig. 3), representing fusaria that are nested within six phylogenetically divergent species complexes (Fig. 1; i.e., *incarnatum-equiseti*, *chlamydosporum*, *sambucinum*, *tricinctum*, *fujikuroi* and *oxysporum*), and as an unrelated fungus, *Septogloeum mori*.

Sequence-based identification of fusaria: 10 simple steps to increase your odds of obtaining an accurate species-level identification

1) Carefully check sequence chromatograms for errors before conducting a BLASTn query (also check

[Display Settings:](#) GenBank

[Send to:](#)

Fusarium sp. NRRL 22244 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S ribosomal RNA gene, partial sequence

GenBank: GQ505685.1

[FASTA](#) [Graphics](#) [PopSet](#)

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LOCUS       GQ505685                1132 bp    DNA     linear     PLN 04-DEC-2009
DEFINITION  Fusarium sp. NRRL 22244 18S ribosomal RNA gene, partial sequence;
              internal transcribed spacer 1, 5.8S ribosomal RNA gene, and
              internal transcribed spacer 2, complete sequence; and 28S ribosomal
              RNA gene, partial sequence.
ACCESSION   GQ505685
VERSION     GQ505685.1   GI:262476544
KEYWORDS    .
SOURCE      Fusarium sp. NRRL 22244
  ORGANISM  Fusarium sp. NRRL 22244
            Eukaryota; Fungi; Dikarya; Ascomycota; Pezizomycotina;
            Sordariomycetes; Hypocreomycetidae; Hypocreales; Nectriaceae;
            Fusarium; Fusarium incarnatum-equiseti species complex.
REFERENCE   1  (bases 1 to 1132)
  AUTHORS   O'Donnell,K., Sutton,D.A., Rinaldi,M.G., Gueidan,C., Crous,P.W. and
            Geiser,D.M.
  TITLE     Novel multilocus sequence typing scheme reveals high genetic
            diversity of human pathogenic members of the Fusarium incarnatum-F.
            equiseti and F. chlamydosporum species complexes within the United
            States
  JOURNAL   J. Clin. Microbiol. 47 (12), 3851-3861 (2009)
  PUBMED    19828752
REFERENCE   2  (bases 1 to 1132)
  AUTHORS   O'Donnell,K., Sutton,D.A., Rinaldi,M.G., Gueidan,C., Crous,P.W. and
            Geiser,D.M.
  TITLE     Direct Submission
  JOURNAL   Submitted (21-AUG-2009) Microbial Genomics, USDA, ARS, NCAUR, 1815
            N. University St., Peoria, IL 61604, USA
FEATURES             Location/Qualifiers
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                     /organism="Fusarium sp. NRRL 22244"
                     /mol_type="genomic DNA"
                     /strain="NRRL 22244"
                     /db_xref="taxon:679408"
                     /note="Fusarium incarnatum-equiseti species complex
                     multilocus haplotype 25_a"
     rRNA             1..42
                     /product="18S ribosomal RNA"
     misc_RNA         43..183
                     /product="internal transcribed spacer 1"
     rRNA             184..340
                     /product="5.8S ribosomal RNA"
     misc_RNA         341..489
                     /product="internal transcribed spacer 2"
     rRNA             490..>1132
                     /product="28S ribosomal RNA"

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Fig. 5 GenBank files for NRRL strains included in several GCPSR-based studies were deposited with notes or comments that identify the phylogenetic species and/or multilocus haplotype (indicated by arrow; O'Donnell *et al.* 2008, 2009a, 2009b, 2010, 2012, 2013, 2014)

problematic nucleotide positions at step 4 if necessary). Based on a detailed GCPSR-based study of the *F. incarnatum-equiseti* species complex (O'Donnell *et al.* 2009b), it seems likely that GenBank accession JN014954.1 contains three sequencing errors (Fig. 4); errors are most commonly found at either end of the sequence and within homopolymers.

2) Avoid using ITS+LSU rDNA sequences to identify unknown fusaria because, compared with *TEF1*, *RPB1* and *RPB2*, they are frequently uninformative at the species-level. However, we do plan to deposit ITS+LSU rDNA sequences of the 93 fusaria included in a robust phylogeny of the genus (O'Donnell *et al.* 2013) in FUSARIUM-ID, *Fusarium MLST* and GenBank to

BLASTn query of GenBank reveals putatively novel fusaria.

Sequences producing significant alignments:

Select: All None Selected:0

Alignments							Download	GenBank	Graphics	Distance tree of results	
	Description	Max score	Total score	Query cover	E value	Ident	Accession				
☐	Fusarium venenatum isolate 64 translation elongation factor 1 alpha gene, partial sequence	1219	1219	100%	0.0	100%	FJ939721.1				
☐	Fusarium venenatum isolate 03 translation elongation factor 1 alpha gene, partial sequence	1094	1094	100%	0.0	97%	FJ939720.1				
☐	Fusarium equiseti isolate DAOM167768 Tomato Albera translation elongation factor-1 alpha (EF-1) gene, partial sequence	1088	1088	89%	0.0	100%	DQ842081.1				
☐	Fusarium venenatum strain FRC R-09186 translation elongation factor-1 alpha (TEF1) gene, partial cds	1086	1086	99%	0.0	97%	GQ915515.1				
☐	Fusarium venenatum isolate 162 translation elongation factor 1 alpha gene, partial sequence	1083	1083	100%	0.0	96%	FJ939722.1				
☐	Fusarium venenatum partial tef-1alpha gene for translation elongation factor 1 alpha, exons 1-4, V101174	1048	1048	96%	0.0	96%	AJ543632.1				
☐	Fusarium venenatum partial tef-1alpha gene for translation elongation factor 1 alpha, exons 1-4, V101176	1046	1046	96%	0.0	96%	AJ543634.1				

Fig. 6 Seven of the 100 BLASTn ‘hits’ from a search of NCBI using a partial *TEF1* sequence of GenBank accession FJ939721.1 (deposited as *F. venenatum*) as the query. However, the relatively low 97% similarity to accession GQ915515.1, which was obtained from an authentic isolate of *F. venenatum* (FRC R-9186), likely indicates the query sequence is from a novel species within the *F. sambucinum* species complex (see Fig. 1). The latter appears to

be conspecific with DAOM 167768 accession DQ842081.1, which was incorrectly deposited in GenBank as *F. equiseti*. This finding suggests that if an *emerencia*-like system (Nilsson *et al.* 2005) was constructed to mine *Fusarium TEF1*, *RPB1* and *RPB2* sequences in GenBank, it could prove to be an invaluable tool for discovering many novel phylogenetically distinct species

facilitate identification to species complex, and in some instances, to several phylogenetically divergent species (Fig. 1; Balajee *et al.* 2009).

3) Avoid querying GenBank with ITS+LSU rDNA sequences because the majority of named sequences are misidentified. When GenBank ITS rDNA accession JN014954.1 was used to query FUSARIUM-ID, it showed 99.78% identity to sequences of four different species within the *F. incarnatum-equiseti* species complex (i.e., FIESC 15, 17, 23 and 25). Instead of depositing the sequence as *Fusarium* sp., which would have been the only correct identification based on the ITS

rDNA sequence data, it was deposited under the name attached to the majority of the named sequences retrieved in their BLASTn query of GenBank (i.e., *F. equiseti*). Our search of the NCBI GenBank database, using accession JN014954.1 as the query, found that the 48 accessions with binomials were all misidentified, including 28 deposited as *F. equiseti*. The real *F. equiseti*, however, corresponds to phylogenetic species FIESC 14 (O'Donnell *et al.* 2009b).

4) Nucleotide polymorphisms or gaps in the alignments displayed after a BLASTn query should **always** be confirmed by rechecking the chromatograms. The

Fig. 7 Flowchart outlining 10 steps that are recommended to increase the likelihood of obtaining an accurate DNA sequence-based species- or species complex-level identification of an unknown *Fusarium*

Sequence-based Identification of *Fusarium* in 10 Simple Steps

- 1) Correct sequence chromatograms completely and carefully
- 2) Use *TEF-1*, *RPB1* and/or *RPB2* and avoid ITS+LSU rDNA
- 3) Query *Fusarium MLST* or FUSARIUM-ID before GenBank
- 4) Recheck chromatograms at positions where gaps are in the BLASTn alignments
- 5) Carefully check species names associated with top BLASTn hits
- 6) Collect DNA sequence data from two independent loci where possible
- 7) Published GCPSR studies may help assess when % similarity indicates conspecificity
- 8) Become familiar with ad hoc species-haplotype nomenclature used in some clades
- 9) Contact culture collections (CBS-KNAW, FRC and NRRL) to inquire about strains
- 10) Misidentifications are more likely to be caught and corrected at better journals

available evidence suggests that the three mismatches in the ITS rDNA alignment of GenBank accessions JN014954.1 and GQ505685.1 NRRL 22244 FIESC 25 (Fig. 4) are due to sequencing errors in the former.

5) Carefully check the taxon names associated with the top hits (i.e., % identity and sequence coverage), who deposited the sequence, and look for notes/comments included in the accession record (Fig. 5). Be prepared to scrutinize multiple names among the top ‘hits’ in BLASTn queries of GenBank. The example provided by GenBank accession JN014954.1 might seem extreme (i.e., all 48 ITS rDNA sequences with binomials were misidentified); however, it serves to highlight an intractable problem created by open deposit of uncurated sequences without allowing third party annotation (Bidartondo *et al.* 2008). By contrast, sequences deposited in FUSARIUM-ID and *Fusarium MLST* were generated primarily as part of GCPSR-based studies of *Fusarium*. When conducting queries of GenBank, it is good practice to look for NRRL strains among the top hits, and if they were deposited as *Fusarium* sp., then look for notes/comments in the accession records.

6) It is prudent to compare BLASTn results from multiple loci where possible; the identification tools built into *Fusarium MLST* allow for queries using single or multiple sequences. Generating a partial *TEF1* sequence is an excellent place to start because both FUSARIUM-ID and *Fusarium MLST* are well-

populated with data from this locus and because it frequently resolves at the species level. Where possible, it is highly desirable to try to identify unknowns by conducting molecular phylogenetic analyses of published MLST datasets to which the unknowns have been added; MLST datasets can be downloaded from FUSARIUM-ID or *Fusarium MLST*. Bootstrap analyses of the individual and combined dataset often yield the most reliable identification of unknowns, when the criteria of genealogical exclusivity and non-discordance under GCPSR are employed (Taylor *et al.* 2000; Dettman *et al.* 2003).

7) The results of prior GCPSR-based studies provide invaluable guidance in interpreting when a given % similarity equates with conspecificity. For the following interpretations to hold true, we assume that carefully edited *TEF1*, *RPB1* and/or *RPB2* sequences are being used as a query against FUSARIUM-ID and/or *Fusarium MLST*. As discussed previously (Geiser *et al.* 2004; O'Donnell *et al.* 2012), an identical match with 99–100% sequence coverage (Fig. 3), in most but not all cases can be interpreted as a definitive species identification. However, it is important to note that our GCPSR-based studies have shown that sequences from these genes sometime fail to distinguish recently evolved sister species (Kasson *et al.* 2013; Sarver *et al.* 2011; Schroers *et al.* 2009). Queries that show similarity at or below 99.4% (i.e., $\geq 4/680$ nucleotide differences in

Ambrosium Fusarium Clade species are distinguished by AF and number between 1–12 (ex., AF-6).

Sequences producing significant alignments:

Select: All None Selected: 0

Alignments Download GenBank Graphics Distance tree of results						
	Description	Max score	Total score	Query cover	E value	Ident Accession
☐	Fusarium ambrosium isolate 31-14 translation elongation factor 1-alpha (EF1-alpha) gene, partial cds	1262	1262	100%	0.0	100% JX677563.1
☐	Fusarium solani strain NRRL 23244 translation elongation factor 1-alpha (EF1-alpha) gene, partial cds	1262	1262	100%	0.0	100% DQ247568.1
☐	Fusarium ambrosium strain NRRL 62605 elongation factor 1-alpha (EF1-a) gene, partial cds	1254	1254	99%	0.0	100% KC691531.1
☐	Fusarium ambrosium strain NRRL 36510 elongation factor 1-alpha (EF1-a) gene, partial cds	1254	1254	99%	0.0	100% KC691530.1
☐	Fusarium ambrosium strain NRRL 20438 translation elongation factor 1 alpha gene, partial cds	1251	1251	100%	0.0	99% AF178332.1
☐	Fusarium sp. AF-6 strain NRRL 62590 elongation factor 1-alpha (EF1-a) gene, partial cds	1210	1210	99%	0.0	99% KC691546.1

Fig. 8 Because binomials for 11/12 species farmed by *Euwallacea* ambrosia beetles were lacking, an informal nomenclature was adopted for the 12 phylogenetically distinct species within the Ambrosia *Fusarium* Clade (AFC). The latter is nested within clade 3 of the *F. solani* species complex (Kasson *et al.* 2013; O'Donnell *et al.* 2014). AFC species are distinguished by AF followed by a

unique number between 1 and 12 (see AF-6 below). Although a partial *TEF1* sequence can be used to identify most of the species within the AFC, multilocus sequence data is recommended for a definitive identification because several species share an identical *TEF1* allele, and because hybrid introgression involving this locus was detected

TEF1) should be subjected to a GCPSR analysis, given the likelihood the unknown represents a novel phylogenetic species (Fig. 6). To increase confidence in *TEF1* queries that show 1–3 base pair differences from the top ‘hit’ in FUSARIUM-ID and *Fusarium MLST*, it is advisable to conduct similar queries using *RPB1* and/or *RPB2* together with bootstrapped phylogenetic analyses as outlined in step 6. To increase the taxonomic representation of fusaria in agronomically important species complexes that haven’t been subjected to GCPSR, and to promote phylogenetic species recognition, we plan to update FUSARIUM-ID and *Fusarium MLST* by depositing several hundred *TEF1*, *RPB1* and *RPB2* unpublished sequences from the *F. sambucinum*, *F. tricinctum* and *F. lateritium* species complexes (Fig. 1).

8) The primary literature should be consulted to take advantage of the ad hoc species and/or species-haplotype nomenclature developed for the *F. incarnatum-equiseti* and *F. chlamydosporum* (O’Donnell *et al.* 2009b), *F. solani* (O’Donnell *et al.* 2008), *F. oxysporum* (O’Donnell *et al.* 2009a) species complexes and Ambrosia *Fusarium* Clade (AFC; Kasson *et al.* 2013; O’Donnell *et al.* 2014). The AFC represents a monophyletic lineage within clade 3 of the *F. solani* species complex that comprises at least 12 phylogenetically distinct mutualistic symbionts of the fungus-farming ambrosia beetle *Euwallacea* (Fig. 8).

9) When questions concerning the metadata or availability of strains arise, contact Kerry O’Donnell at kerry.odonnell@ars.usda.gov and the curators of FUSARIUM-ID (David M. Geiser, dmgl7@psu.edu) or *Fusarium MLST* (Vincent Robert, v.robert@cbs.knaw.nl) for clarification.

10) Publish in journals where submissions are more likely to be reviewed by those knowledgeable in the pitfalls associated with sequence-based identification. Here, it is worth reiterating the point made in Wingfield *et al.* (2012) that an accurate species identification is crucial for communicating findings to the scientific community.

Although *Fusarium* molecular phylogenetics and systematics has been characterized as a model (Kang *et al.* 2010), species limits and evolutionary relationships within several of the most important plant pathogenic lineages (i.e., *F. fujikuroi*, *F. oxysporum*, *F. solani* and *F. sambucinum* species complexes) are not fully resolved, especially within the most recently evolved clades (O’Donnell *et al.* 2013). As mentioned earlier,

one of the primary objectives for generating genome-scale data across the breadth of *Fusarium* is to mine it for additional phylogenetically informative loci for NGS-based species-level studies. In the not too distant future, our goal is to identify additional genes that are as informative as *TEF1*, *RPB1* and *RPB2* to further *Fusarium* identification, molecular diagnostics and robust phylogenetic inference.

Acknowledgments Special thanks are due to Stacy Sink for assistance with the DNA sequence data, Nathane Orwig for running DNA sequences in the NCAUR DNA core facility, and Bongsoo Park for maintaining FUSARIUM-ID. The mention of firm names or trade products does not imply that they are endorsed or recommended by the US Department of Agriculture over other firms or similar products not mentioned. The USDA is an equal opportunity provider and employer.

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