

The Myrtle rust pathogen, *Puccinia psidii*, discovered in Africa

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Abstract: *Puccinia psidii*, the cause of a disease today commonly referred to as Myrtle rust, is considered a high priority quarantine threat globally. It has a wide host range in the *Myrtaceae* and it is feared that it may result in significant damage to native ecosystems where these plants occur. The fungus is also of considerable concern to plantation forestry industries that propagate Australian *Eucalyptus* species. In May 2013, symptoms of a rust disease resembling those of *P. psidii* were observed on an ornamental Myrtaceous shrub in a garden in South Africa. The fungus was identified based on DNA sequence data of the ITS and 5.8S nrRNA gene regions and here we report, for the first time, the presence of *P. psidii* in Africa.

Key words:

Guava rust
Eucalyptus rust
Myrtus communis
Pucciniaceae
Uredo rangellii

Article info: Submitted: 13 June 2013; Accepted: 14 June 2013; Published: 24 June 2013.

INTRODUCTION

Puccinia psidii (*Uredinales*, *Pucciniaceae*) has been considered as an important quarantine threat to many countries (Glen *et al.* 2007). It was first described from native guava (*Psidium guajava*) in Brazil in 1884 and gained notoriety when it was found to infect various other members of the *Myrtaceae*, an unusual feature for most rust fungi (Coutinho *et al.* 1998, Glen *et al.* 2007, Carnegie *et al.* 2010a, Morin *et al.* 2012). *Puccinia psidii* became particularly prominent in the literature when it was found causing disease on non-native *Eucalyptus* species in Brazil (Jollify 1944) and it was rapidly considered as a significant threat to the commercial production of *Eucalyptus* species globally (Coutinho *et al.* 1998). It was also feared to threaten the survival of native *Myrtales* in countries such as Australia where this order represents a mega-diverse group of plants (Glen *et al.* 2007, Morin *et al.* 2012).

Since the first report of *P. psidii* in Brazil, the rust has been recorded in several countries of South and Central America, including the Caribbean (Coutinho *et al.* 1998, Glen *et al.* 2007, Graça *et al.* 2011, Morin *et al.* 2012). It has moved, increasingly rapidly, to new environments including California, Florida and Hawaii in the USA (Marlatt & Kimbrough 1979, Rayachhertry *et al.* 1997, Uchida *et al.* 2006), Japan (Kawanishi *et al.* 2009), Australia (Carnegie *et al.* 2010), and China (Zhuang & Wei 2011). Rusts in the guava/eucalyptus rust complex are native to South and Central America (Alfenas *et al.* 2005) and are feared because of their wide host range in *Myrtaceae*, including over 125 species (Morin *et al.* 2012). Thus, the recent appearance of *P. psidii* in Australia has resulted in many studies to consider its likely long-term impact (Carnegie & Cooper 2011, Morin *et al.* 2012, Kriticos *et al.* 2013).

Several common names have been used for the disease caused by *P. psidii*. That it was first found on guava led to it being known as Guava rust, but its appearance on more commercially important *Eucalyptus* species led to it commonly being referred to as Eucalyptus rust. When it first appeared in Australia, there was debate regarding its taxonomy and whether the fungus infecting a wide range of trees in *Myrtaceae* might not be the rust that had been described as *Uredo rangellii* (Carnegie *et al.* 2010, Carnegie & Cooper 2011). While there are clearly taxonomic issues relating to this fungus that remain to be resolved, such as that it is not phylogenetically related to other members of the genus *Puccinia* (M. Wingfield & W. Maier, unpubl. data), the disease caused by the fungus currently treated as *P. psidii* is best referred to as Myrtle rust. This captures the occurrence of the pathogen on a very wide host range including, numerous genera and species of *Myrtales*.

In May 2013, an ornamental *Myrtus communis* plant growing in a residential garden in the KwaZulu-Natal province of South Africa was discovered showing typical symptoms of infection by *P. psidii*. The aim of this study was to identify the fungus using DNA sequence data and to determine whether one of the globally most important invasive alien plant pathogens might have entered South Africa, which would represent the first confirmed record of this pathogen on the African continent.

MATERIALS AND METHODS

Leaves and shoots were taken from the infected *Myrtus communis* plant, collected in brown paper bags, and transported to the laboratory for study. The plant showed typical symptoms of infection by *Puccinia psidii*. These

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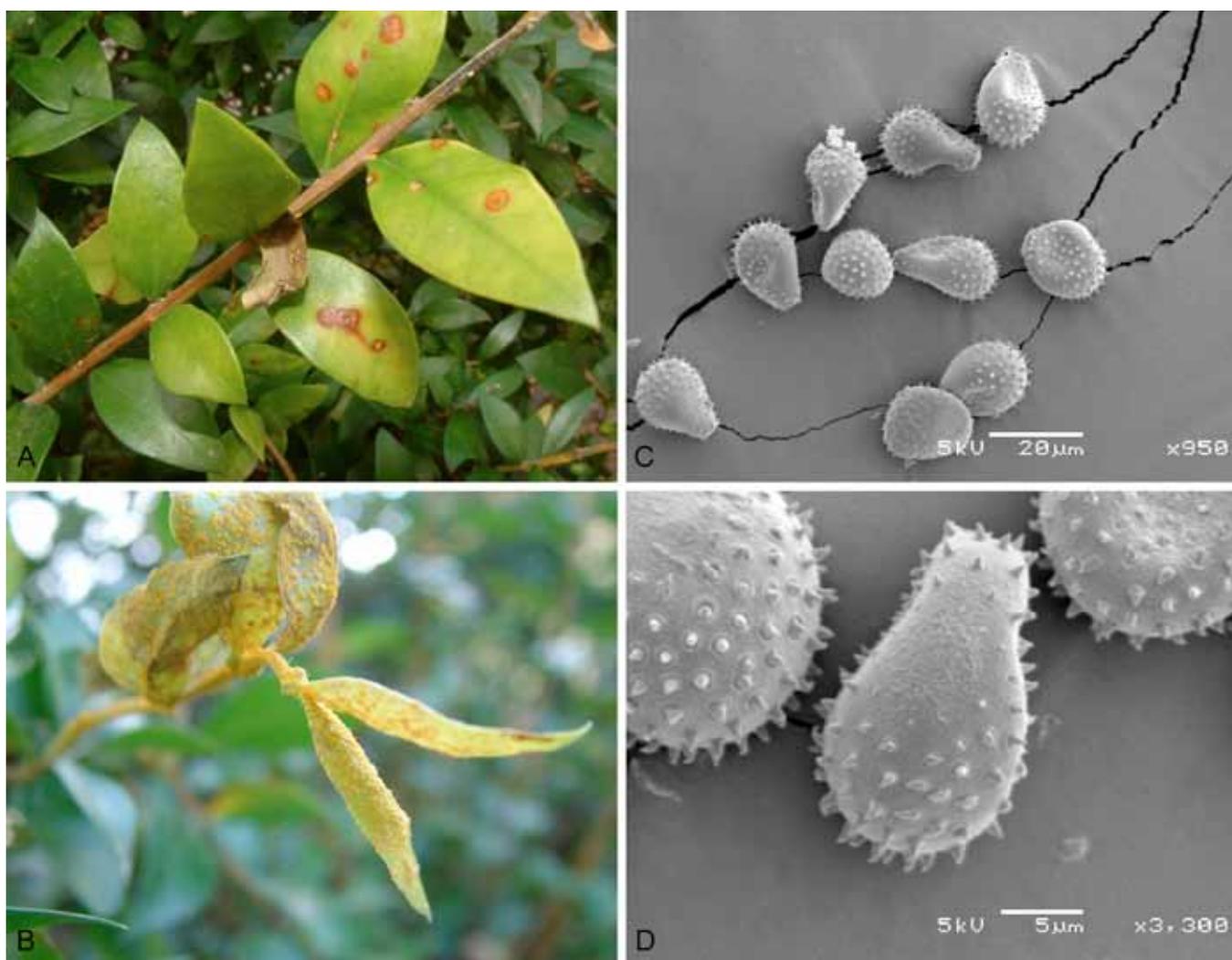


Fig. 1. *Puccinia psidii* on *Myrtus communis* in South Africa. **A.** Leaf spot on *Myrtus communis*. **B.** Yellow masses of urediniospores covering a dying *M. communis* shoot tip. **C–D.** Urediniospores with echinulations and smooth pathes (tonsures).

included leaf spots and cankers on young shoots/petioles (Fig. 1). Infected tissues were covered with yellow urediniospores. Spores were collected directly from infected material and used for morphological and DNA sequence studies (Table 1).

For DNA sequence studies, urediniospores were scraped from the surface of infected material into 18 μ L sterile SABAX water. The SABAX water containing urediniospores were incubated at 94 °C for 10 minutes and used as template in subsequent reactions for amplification of the ITS1, 5.8S and ITS2 gene regions of the Internally Transcribed Spacer regions of the nuclear rDNA. Amplification reactions were performed in a final reaction volume of 25 μ L containing: 5 μ L 5 \times MyTaq™ Reaction Buffer (Bioline, London), 0.2 mM of each of the universal primers ITS 1-F (Gardes & Bruns 1993) and ITS 4 (White *et al.* 1990) and 1 U of MyTaq™ DNA polymerase. The PCR conditions were as follows: Initial denaturation at 94 °C for 3 min followed by 30 cycles of denaturation at 94 °C for 1 min, annealing at 53 °C for 1 min, and elongation at 72 °C for 1 min. A final elongation step at 72 °C for 10 min followed. Products were separated using gel electrophoresis and visualised using GelRed™ (Biotium, CA).

PCR Amplification products were purified using the Zymo research DNA Clean & Concentration™ - 5 kit (CA). Fragments were sequenced, using forward and reverse primers as described above, using the ABI Prism® Big Dye™ Terminator 3.0 Ready Reaction Cycle sequencing Kit (Applied Biosystems, Foster City, CA). Sequences were determined with an ABI PRISM™ 3100 Genetic Analyzer (Applied Biosystems). DNA sequences of opposite strands were edited and consensus sequences obtained using CLC Main workbench v. 6.1 (CLC Bio, www.clcbio.com) and MEGA v. 5 (Tamura *et al.* 2011). Sequences obtained were submitted to NCBI's GenBank (<http://www.ncbi.nlm.nih.gov/Genbank/index.html>) with accession numbers KF220289 – KF220293.

Sequences obtained for the rust fungus from South Africa were subjected to a Blastn search on the NCBI database (<http://www.ncbi.nlm.nih.gov>) and thereafter incorporated into a dataset of closely related sequences for phylogenetic analyses. After online alignment using MAFFT v. 7 (<http://mafft.cbrc.jp/alignment/server/>), the programme MEGA v. 5.1 (Tamura *et al.* 2011) was used to check the alignments and conduct a Maximum Likelihood analysis of the data set.

Table 1. List of *Puccinia* isolates used in DNA sequence analyses.

Species	Origin	GenBank Accession nr.	Host
<i>Puccinia psidii</i>	Australia	HM448900	<i>Agonis flexuosa</i>
	Brazil	AJ536601	<i>Psidium guajava</i>
	Brazil	AJ421801	<i>Eugenia uniflora</i>
	Brazil	AJ421802	<i>Melaleuca quinquenervia</i>
	Colombia	EU711423	<i>Syzygium jambos</i>
	Hawaii	EF599768	<i>Metrosideros polymorpha</i>
	Hawaii	EU071045	<i>Melaleuca quinquenervia</i>
	Florida	AJ535659	<i>Pimenta dioca</i>
	Japan	AB470483	<i>Metrosideros polymorpha</i>
	South Africa	KF220289	<i>Myrtus communis</i>
	South Africa	KF220290	<i>M. communis</i>
	South Africa	KF220291	<i>M. communis</i>
	South Africa	KF220292	<i>M. communis</i>
	South Africa	KF220293	<i>M. communis</i>
	Uruguay	EU348742	<i>Eucalyptus grandis</i>
	Uruguay	EU348743	<i>E. globulus</i>
<i>P. cygnorum</i>		EF490601	<i>Kunzea ericifolia</i>
<i>P. hordei</i>		AF511086	n/a
<i>P. recondita</i>		AF511082	<i>Triticum turgidum</i>

Puccinia cygnorum (EF490601), *P. hordei* (AF511086), and *P. recondita* (AF511082) were used as outgroup species in the analyses.

Measurements of 20 urediniospores were made using a Zeiss Axioscop compound microscope and photographic images were captured using a Zeiss AxioCam MRc digital camera and the AxioVision v. 4.8 (Carl Zeiss) software. Scanning electron micrographs (SEM) were obtained directly from urediniospores scraped from infected material using a JSM-840 SEM (JEOL, Tokyo) at 5 kV and images captured with Orion v. 6.60.4 (E.L.I. s.p.r.l., Brussels, Belgium).

RESULTS

The infected *Myrtus communis* plant showed symptoms of leaf spot, with red margins, and the presence of abundant yellow spore masses on young shoots and leaves (Fig. 1). Infection resulted in the death of shoots and leaves. Only urediniospores were observed on the plant. These ranged in size from 15–20 (av = 19) × 12–16 (av = 14) µm. SEM of the urediniospores revealed the presence of tonsures (smooth patches) on the surfaces of some spores (Fig. 2).

Amplification reactions of the ITS and 5.8S gene regions resulted in fragments of ~700 bp in length. The Blast search on the NCBI database showed that the rust fungus from South Africa (KF220289 – KF220293) was most closely related to *Puccinia psidii*. Subsequent comparisons using ML in MEGA, of a dataset comprising 17 sequences (Table 1) of 626 bp in length, showed that there were no differences in the ITS sequences between the South African collection and those from other parts of the world. All *P. psidii* isolates grouped

together in a single clade, separately from the other *Puccinia* species included in the analyses.

DISCUSSION

This is the first confirmed report of the Myrtle rust pathogen, *Puccinia psidii*, from South Africa. There have been two previous reports of a rust fungus on *Eucalyptus* in South Africa (Knipscheer & Crous 1990, Maier *et al.* 2010), but both were morphologically different to *P. psidii*. The present study provides robust evidence that *P. psidii* is now present in South Africa. This is an important discovery and it is one that is of considerable concern, especially as the species is unrecorded elsewhere in Africa.

The discovery of *P. psidii* in South Africa is significant to both the commercial plantation forestry industry, as well as the conservation of native plants and associated ecosystems. The impact of *P. psidii* on plantation grown *Eucalyptus* species has been shown in several previous studies (Tommerup *et al.* 2003, Graça *et al.* 2011, Silva *et al.* 2013). More recently, a number of studies have shown the broad host range of the pathogen on other *Myrtaceae* and the fungus has been described as a significant threat to native ecosystems in Australia (Morin *et al.* 2012). Importantly, a study has also shown that native South African *Heteropyxis natalensis* (*Myrtales*, *Heteropyxidaceae*) is highly susceptible to infection by *P. psidii* (Alfenas *et al.* 2005).

The majority of *P. psidii* hosts reside in the subfamily *Myrtoideae* of the *Myrtaceae* (Morin *et al.* 2012). The susceptibility, in greenhouse studies, of South African *H. natalensis* in *Heteropyxidaceae*, clearly shows the potential

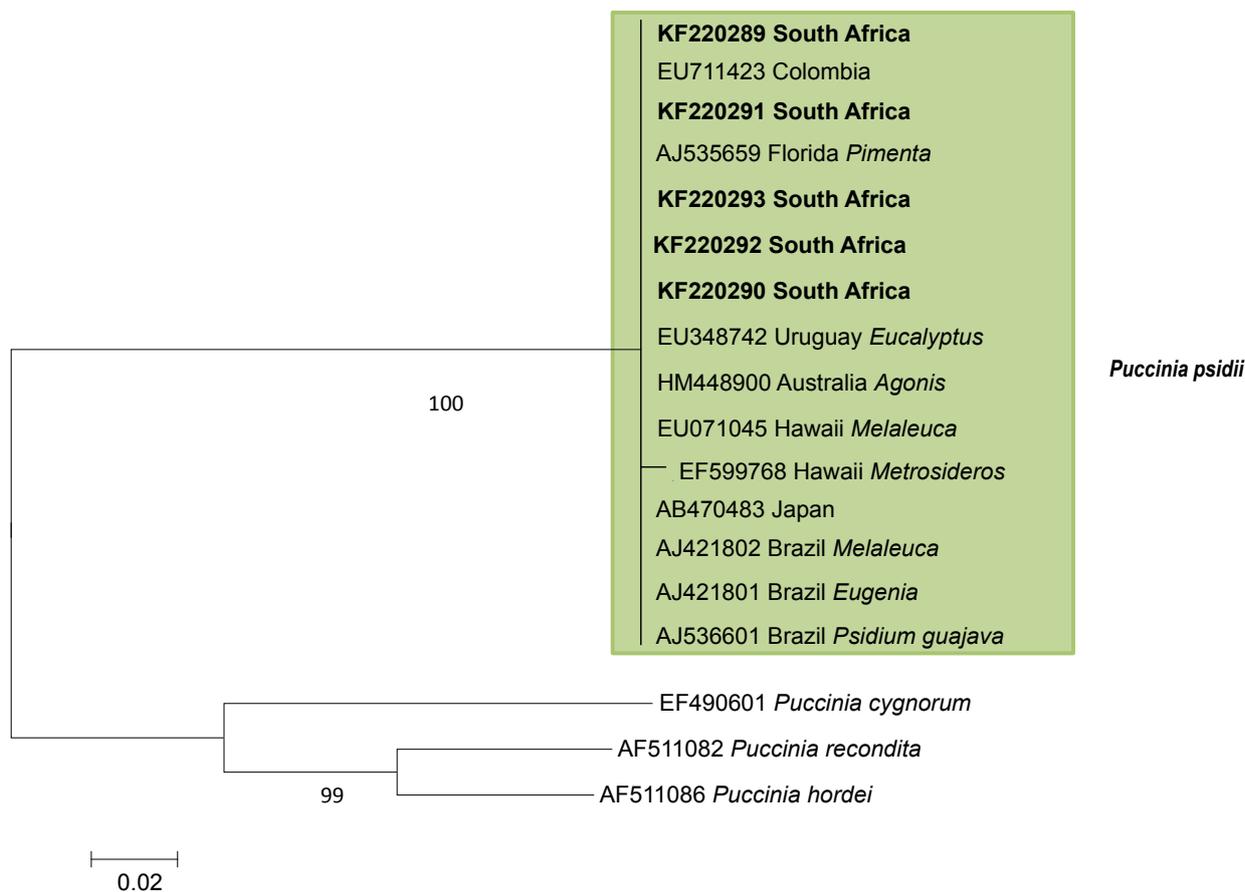


Fig. 2. Phylogenetic tree of ITS sequence data showing identity of South Africa *Puccinia psidii* isolates. Maximum likelihood tree based on 626 bp from 17 taxa. Numbers below branches indicate bootstrap support values.

damage that the pathogen might pose to other families in *Myrtales*. South Africa is home to three families in *Myrtales* (Palgraves & Palgraves 2002), including species that are endemic and thus of significant conservation importance. Of these, five genera (*Eugenia*, *Heteropyxis*, *Memecylon*, *Syzygium*, and *Warneckea*) occur in the KwaZulu-Natal Province where *P. psidii* has now been detected. Clearly, much work has yet to be done to consider the potential impact that *P. psidii* might have in South Africa. It is known that *P. psidii* has considerable race specialisation (Coelho et al. 2001, Aparecido et al. 2003) and it will be necessary to determine the host range of the fungus now present, but known only on a single host species from a single locality in South Africa.

The fungus on *Myrtus communis* in South Africa is morphologically similar to *P. psidii* from elsewhere in the world. Urediniospores of the South African collection were, however, smaller than those reported for *P. psidii* from Uruguay (Perez et al. 2011) and for *Uredo rangelii* (Simpson et al. 2006). SEM showed the presence of typical spines as well as tonsures on the urediniospores. There have been previous arguments that these tonsures are characteristic of *U. rangelii*, and not *P. psidii* (Simpson et al. 2006, Carnegie et al. 2010a). However, they are common on the rust now considered to be *P. psidii* in Australia (Carnegie et al. 2011) and this is also true for collections that infect *Eucalyptus* and native *Myrtaceae* in Uruguay (Carnegie et al. 2010, Perez et

al. 2011). Although *P. psidii* might represent a suite of related cryptic species (Wingfield, personal observation), in-depth studies are needed to elucidate this question.

There are relatively limited options to manage the *P. psidii* infection. Eradication of this new invasion is unlikely to be effective because rust fungi produce abundant air-borne spores, and will be highly dependent on rapid action. Selection of resistant plant material will be the most durable approach. For example, considerable variation has been identified in susceptibility of *Eucalyptus* genotypes to infection by *P. psidii* (Carvalho et al. 1998, Junghans et al. 2003, Silva et al. 2012), which implies that it will be possible to restrict the damage that might occur in commercial plantation situations. The deployment of resistant native *Myrtaceae*, if present, will, however, be more complicated than that of a commercial crop such as *Eucalyptus* in South Africa. Dealing with the disease in native ecosystems will be much more complex. Therefore, great effort should be made to slow the movement of the pathogen into native ecosystems, particularly in areas such as the Western Cape, where only a single, endemic species of *Myrtaceae* (*Metrosideros angustifolia*) occurs. In Australia, *Puccinia psidii* was described as “the pinnacle of pathogens we wanted to keep out of Australia” (Dayton & Higgins 2011). Its appearance in South Africa is likely to have substantial negative long-term consequences for both forestry and plant conservation. South Africa is also likely to now provide the bridge for *P. psidii* to move northwards in Africa.

ACKNOWLEDGMENTS

We thank members of the Tree Protection Co-operative Programme (TPCP), DST/NRF Centre of Excellence in Tree Health Biotechnology (CTHB) and the University of Pretoria for funding and facilities to undertake this work. We also thank Mr. Alan Hall of the microscopy unit of the University of Pretoria for the SEM photos of the urediniospores.

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