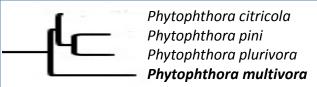
Determining the origin of the emerging pathogen, *Phytophthora multivora*

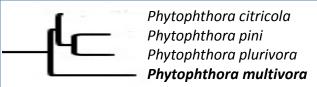
Alex Rea, Giles Hardy, Mike Stukely and Treena Burgess









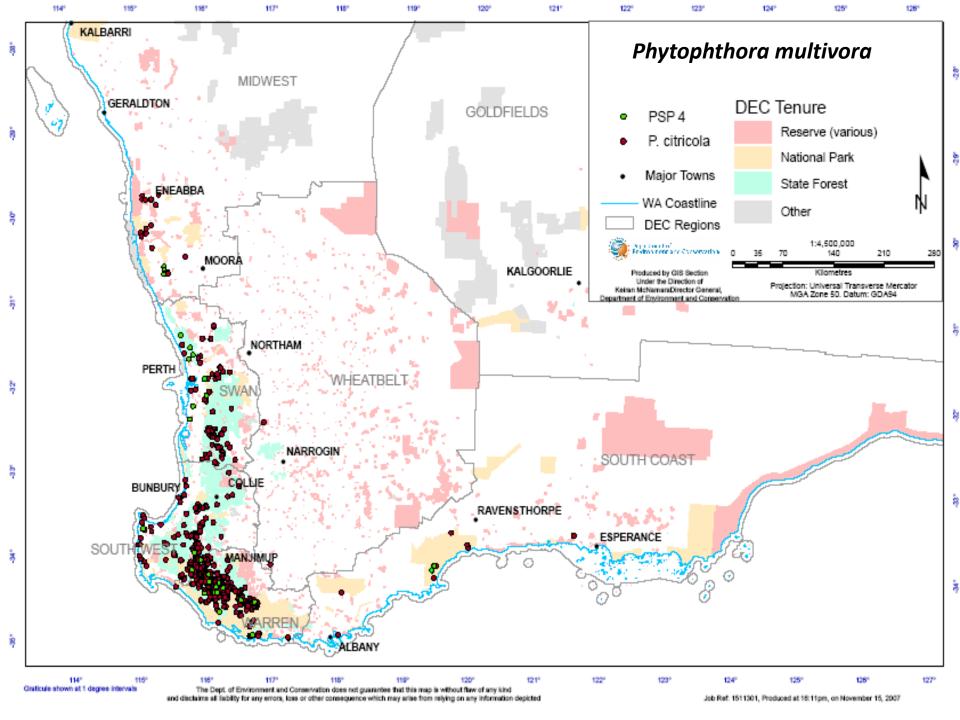


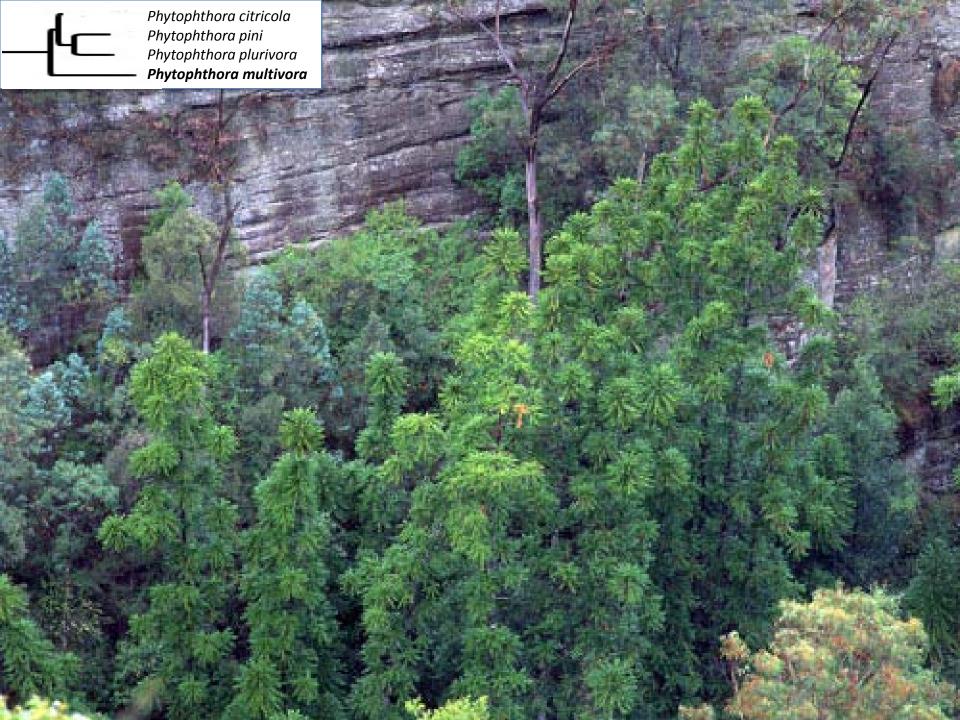












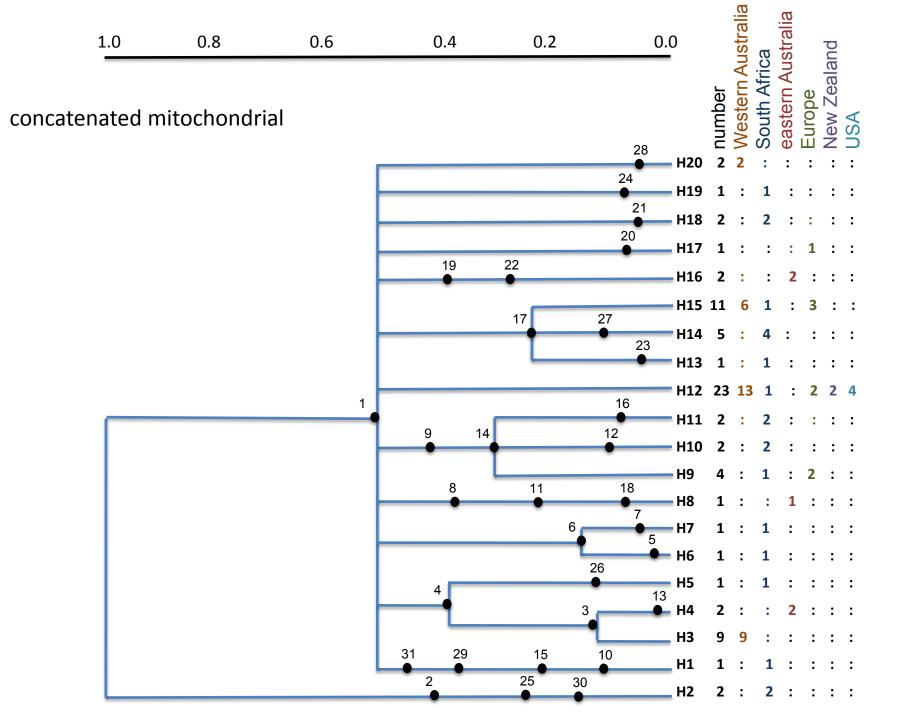


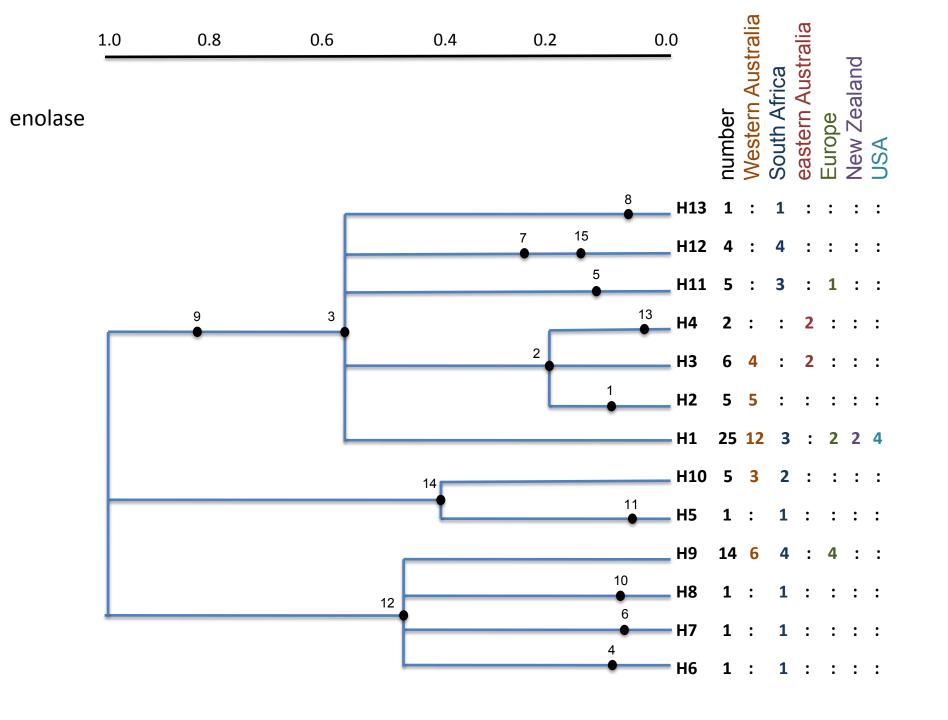


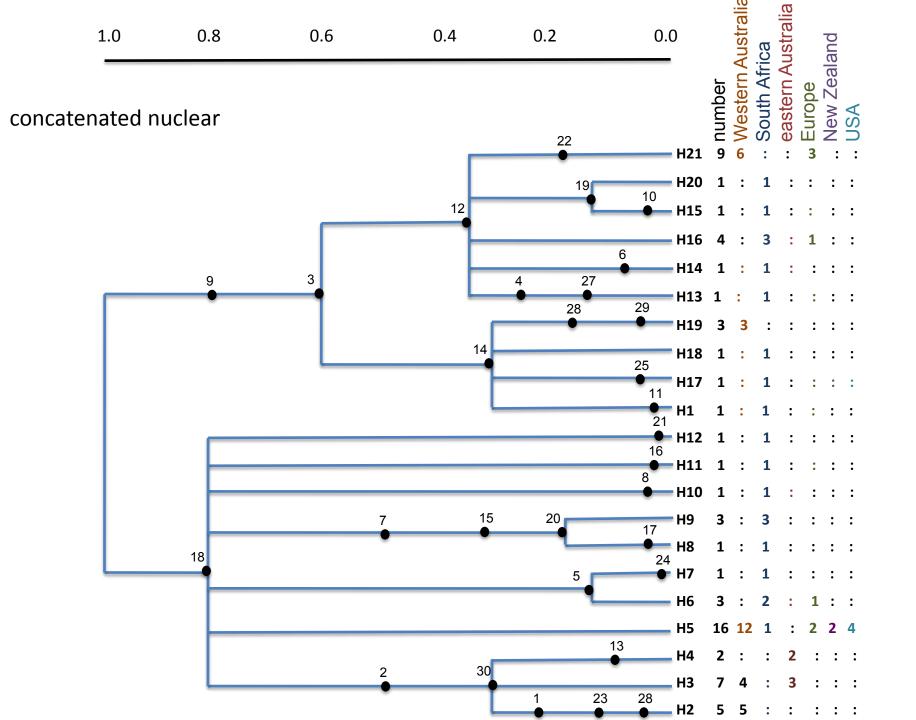


- When P. multivora was described in 2009; there were 7 matching ITS sequences on GenBank from Hungary, Canada, Switzerland, Spain, Korea and Japan
- due to sequence polymorphism and widespread occurrence we postulated and Australian origin for this species
- Since the description *P. multivora* was identified in South Africa, New Zealand, USA and other European countries
- In Australia, *P. multivora* has been isolated once in NSW from Wollemi pine and once in Victoria from a pipeline survey
- In Western Australia approximately 500 sequence verified isolates and it is now being routinely isolated from peri-urban and urban areas
- through collaboration at FABI in South Africa found P. multivora to be the most common species in health natural forest ecosystems and readily isolated in forest streams
- we gathered a set of isolates to test the hypothesis that P. multivora was endemic to Australia
- sequenced 5 nuclear genes and 3 mitochondrial genes and conducted coalescence analysis

Location	no.	Hosts
Western Australia	30	numerous hosts, isolates selected from wide geographic distribution, commonly isolated
Australia (elsewhere)	5	Wollemi pine (NSW), rarely from natural vegetation (VIC)
South Africa	21	from soil beneath asymptomatic natural vegetation, also from water
New Zealand	2	forest survey (highly impacted region)
Europe	8	nurseries and young plantings
USA	4	forest survey (encountered rarely)







	/	n	5	h	k	π ± SE (x10 ⁻³)	θw
Mitochondrial loci	1072	70	32	22	3.769	3.61±0.23	6.648
WA	1070	30	8	5	2.759	2.58±0.19	2.019
R5A	1044	21	21	16	4.608	4.41±0.40	6.008
Nuclear loci	2140	70	37	21	6.505	3.40±0.20	7.963
WA	2140	30	14	5	3.929	1.77±0.36	2.544
RSA	2140	21	27	16	7.170	3.35±0.24	7.725

/= length of sequence;

n = sample size;

s = number of segregating sites;

h = number of haplotypes;

k = mean number of pairwise nucleotide differences;

 π = mean number of base differences per site;

 θ w = mean population mutation rate.

Mitochondrial loci = concatenated coxIGS and nadh1 sequence data;

Nuclear loci = concatenated enolase, hsp90, ras, and asf-like sequence data

- at the time of its description, P. multivora was virtually unknown outside Western Australia
- subsequently it is appearing more and more often in Europe
- also in Western Australia it has become the most common species isolated in the urban and peri-urban environment
- in South Africa, it is widely distributed and readily isolated from natural forest with no disease symptoms
- no unique isolates from Europe, USA and NZ, *P. multivora* appears to be introduced to these regions but to date dataset no large enough
- in coalescence analysis, more variable sites, more haplotypes and highest diversity among isolates from South Africa
- based on coalescence analysis, isolate diversity and biology we propose South Africa is the origin of *P. multivora*