Figure S1. Growth curves of *Knoxdaviesia capensis*, *K. proteae* and *C. albifundus* on all PM1 substrates. The 96 well plate is shown in four separate blocks for clarity: (A) wells A-B, (B) wells C-D, (C) wells E-F, (D) wells G-H.



(C)



Figure S2. Growth curves of *Knoxdaviesia capensis*, *K. proteae* and *C. albifundus* on all PM2 substrates. The 96 well plate is shown in four separate blocks for clarity: (A) wells A-B, (B) wells C-D, (C) wells E-F, (D) wells G-H.





(C)



Figure S3. *Ceratocystis albifundus* growth curves for all PM1 (A-D) and PM2 (E-H) substrates. The thick black curve is the negative control. The names of active and ambiguous substrates are indicated on the plot.



(D)



(E)





se	D03 Sedoheptulosan
al and a second s	D04 L-Sorbose
tose	D05 Stachyose
	D06 D-Tagatose
-D-Glucoside	D07 Turanose
D-Galactoside	D08 Xylitol
nyl-D-Glucose	D09 N-Acetyl-D-Glucosaminito
D-Glucuronic Acid	D10 y Amino-n-Butyric Acid
-D-Mannoside	D11 d Amino Valeric Acid
-D-Xylopyranoside	D12 Butyric Acid
se	

D01 D-Raffinose

D02 D-Salicin

Supplementary File 4: Aylward *et al.* Contrasting carbon metabolism in saprotrophic and pathogenic Microascalean fungi from *Protea* trees





(H)





Figure S4. Carbohydrate utilization pathways in *Knoxdaviesia capensis*, *K. proteae* and *Ceratocystis albifundus*. Substrates tested in the PM assays are circled and coloured according to usage: used by all (green), used by none (white), used by both *Knoxdaviesia* species (purple), used by *K. capensis*

only (red) and used by C. albifundus only (blue). The same colouring scheme applies to the presence of enzymes (boxed). The grey double-lined ovalrepresentsthefungalcell.



Figure S5. Predicted enzymes associated with pectin backbone (homogalacturonan) degradation in *Ceratocystis albifundus* and *Knoxdaviesia*. None of the species are apparently able to complete the pathway.