Supplementary material

Table S1 Values of the mycelial and biofilm growth of *F. circinatum*

Azole	I ₅₀ mg/L	
	Mycelia	Biofilms
Tebuconazole	0.04±0.01 b	0.46±0.07 b
lmazalil	0.26±0.13 b	0.74±0.05 b

^{*}b – Indicates statistically different means between samples using student *t*-test with a *P* value (p<0.01). n = 3



Figure S1. Growth of *Fusarium circinatum* FSP34 isolate at the air/liquid interface in ½ strength Potato Dextrose Broth (PDB). The cells were left to incubated for 7 days at stationary phase, following which a culture with a consistency of slime or a pellicle was observed

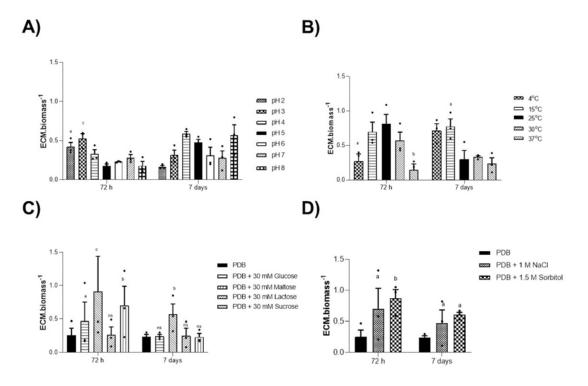


Figure S2. ECM production per unit of biomass under different conditions. A - pH, B – temperature, C - sugars and D - osmotic stress. Means that were statistically different (p<0.01, one-way ANOVA) between the various treatments at a particular time point are indicated with different letters, while those lacking significant differences were indicated with "NS". Error bars indicate standard errors. n = 3, PC - positive control, PDB - potato dextrose broth

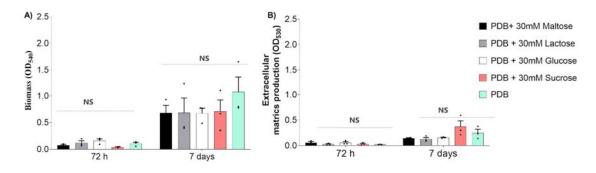


Figure S3. Fusarium circinatum biofilm formation in the presence of sugars. The formation of biofilms was assessed using crystal violet for biomass (A) and basic fuchsin (B) for extracellular matrix (ECM) using 72-h and 7-day-old biofilms. The means that were statistically different are assigned a P value (p<0.005, one-way ANOVA) while those lacking significant differences were indicated with "NS". Error bars indicate standard errors. n = 3, PDB - potato dextrose broth