

Table S1. *C. albicans* strains used in this study.

Strain	Genotype	Source
<i>C. albicans</i>		
SC5314	Wild type	[1]
CAF2-1	<i>ura3Δ::imm434/URA3</i>	[2]
SPCa2	<i>pmt1Δ::hisG/pmt1Δ::hisG ura3Δ::imm434/URA3</i>	[3]
SPCa4	<i>pmt2Δ::hisG/PMT2 ura3Δ::imm434/URA3</i>	[3]
SPCa6	<i>pmt4Δ::hisG/pmt4Δ::hisG ura3Δ::imm434/URA3</i>	[3]
SPCa8	<i>pmt6Δ::hisG/pmt6Δ::hisG ura3Δ::imm434/URA3</i>	[3]
SPCa10	<i>pmt5Δ::hisG/pmt5Δ::hisG ura3Δ::imm434/URA3</i>	[3]
NGY152	<i>ura3Δ::imm434/ura3Δ::imm434, RPS1/rps1Δ::CIP10</i>	[4]
NGY145	<i>ura3Δ::imm434/ura3Δ::imm434, mnt2Δ::hisG/mnt2Δ::hisG, RPS1/rps1Δ::CIP10</i>	[5]
NGY158	<i>ura3Δ::imm434/ura3Δ::imm434, mnt1Δ::hisG/mnt1Δ::hisG, RPS1/rps1Δ::CIP10</i>	[5]
NGY337	<i>ura3Δ::imm434/ura3Δ::imm434, mnt1-mnt2Δ::hisG, RPS1/rps1Δ::CIP10</i>	[5]
NGY516	<i>ura3Δ::imm434/ura3Δ::imm434, mnt4::hisG/mnt4Δ::hisG, mnt5Δ::dp1200/mnt5Δ::dp1200, RPS1/rps1Δ::CIP10</i>	[6]
NGY1227	<i>ura3Δ::imm434/ura3Δ::imm434, mnt3Δ::dp1200/mnt3Δ::dp1200, mnt5Δ::hisG/mnt5Δ::hisG, RPS1/rps1Δ::CIP10</i>	[6]
NGY357	<i>ura3Δ::imm434/ura3Δ::imm434, och1Δ::hisG/och1Δ::hisG, RPS1/rps1Δ::CIP10</i>	[7]
CDH15	<i>ura3Δ::imm434/ura3Δ::imm434, mnn4Δ::hisG/mnn4Δ::hisG, RPS1/rps1Δ::CIP10</i>	[8]
NGY355	<i>ura3Δ::imm434/ura3Δ::imm434, pmr1Δ::hisG/pmr1Δ::hisG, RPS1/rps1Δ::CIP10</i>	[9]
SN250	Wild type	[10]
och1	From Noble's mutant library	[10]

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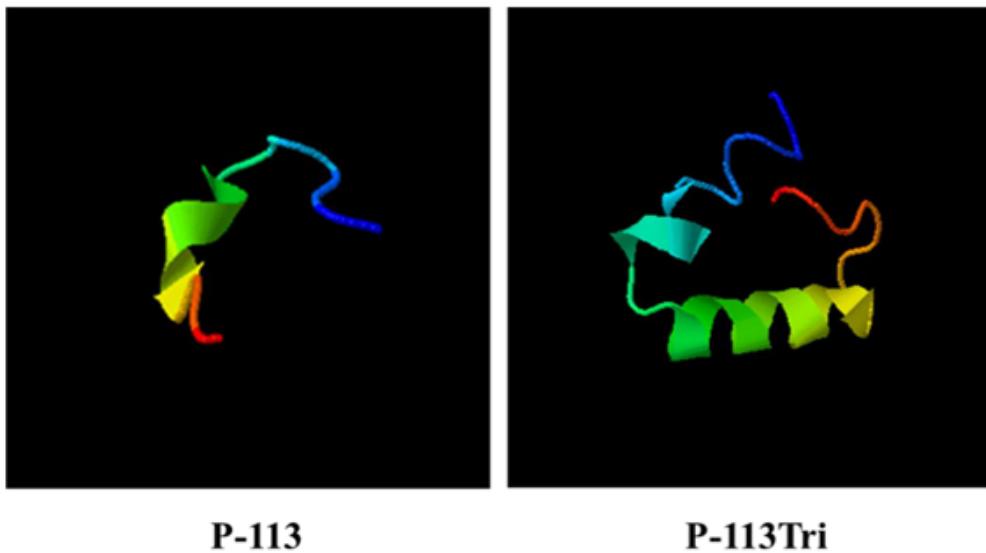


Figure S1. 3D structure of P -113 and P-113Tri predicted by I-TASSER (Iterative threading assembly refinement).

P-113 shows a coil structure, whereas P-113Tri exhibits an α -helical structure. I-TASSER on-line platform (<https://zhanglab.ccmb.med.umich.edu/I-TASSER/>).

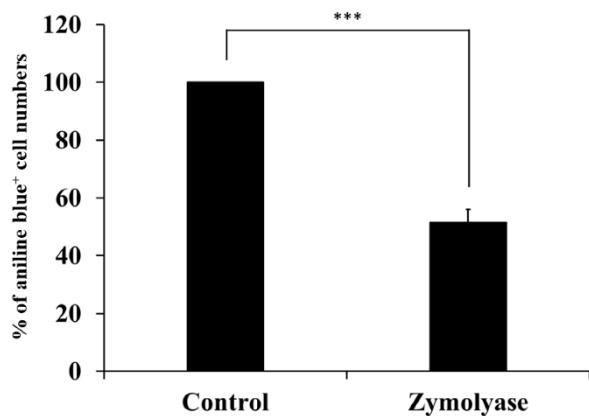


Figure S2. Aniline blue binding to zymolyase-treated *C. albicans*.

C. albicans cells were treated with Zymolyase (2.5 mg/ml) for 1 h at 37°C. Then, the cells were treated with aniline blue (500 μ g/ml) for 5 min in a black 96-well microplate. *, p<0.05; **, p<0.01; ***, p<0.001.