

Table S1 Sequence identities of *OfChtI*-CAD with other structure-known chitinases from different species.

Species	PDB	Identity (%)	RMSD
<i>Homo sapiens</i>	3fxy	43	0.90
	1guv	40	1.07
<i>Nicotiana tabacum</i>	3alf	30	1.79
<i>Bacillus circulans</i>	1itx	29	1.07
<i>Arabidopsis thaliana</i>	3aqu	27	1.56
<i>Serratia marcescens</i>	1e15	27	1.80
	1ctn	27	1.53
<i>Lactococcus lactis</i>	3ian	27	2.74
<i>Aspergillus fumigatus</i>	1w9p	27	1.56
	2y8v	26	2.87
<i>Vibrio harveyi</i>	3b8s	26	1.26
<i>Yersinia entomophaga</i>	3oa5	26	1.80
<i>Bacteroides thetaiotaomicron</i>	3co4	25	2.20
<i>Coccidioides immitis</i>	1d2k	25	1.16
<i>Arthrobacter tad20</i>	1kfw	24	1.52
<i>Clonostachys rosea</i>	3g6l	23	1.25
<i>Moritella marina</i>	4hmc	22	2.67
<i>Bacillus cereus</i>	3n15	22	2.67
<i>Crocus vernus</i>	3sim	21	2.47

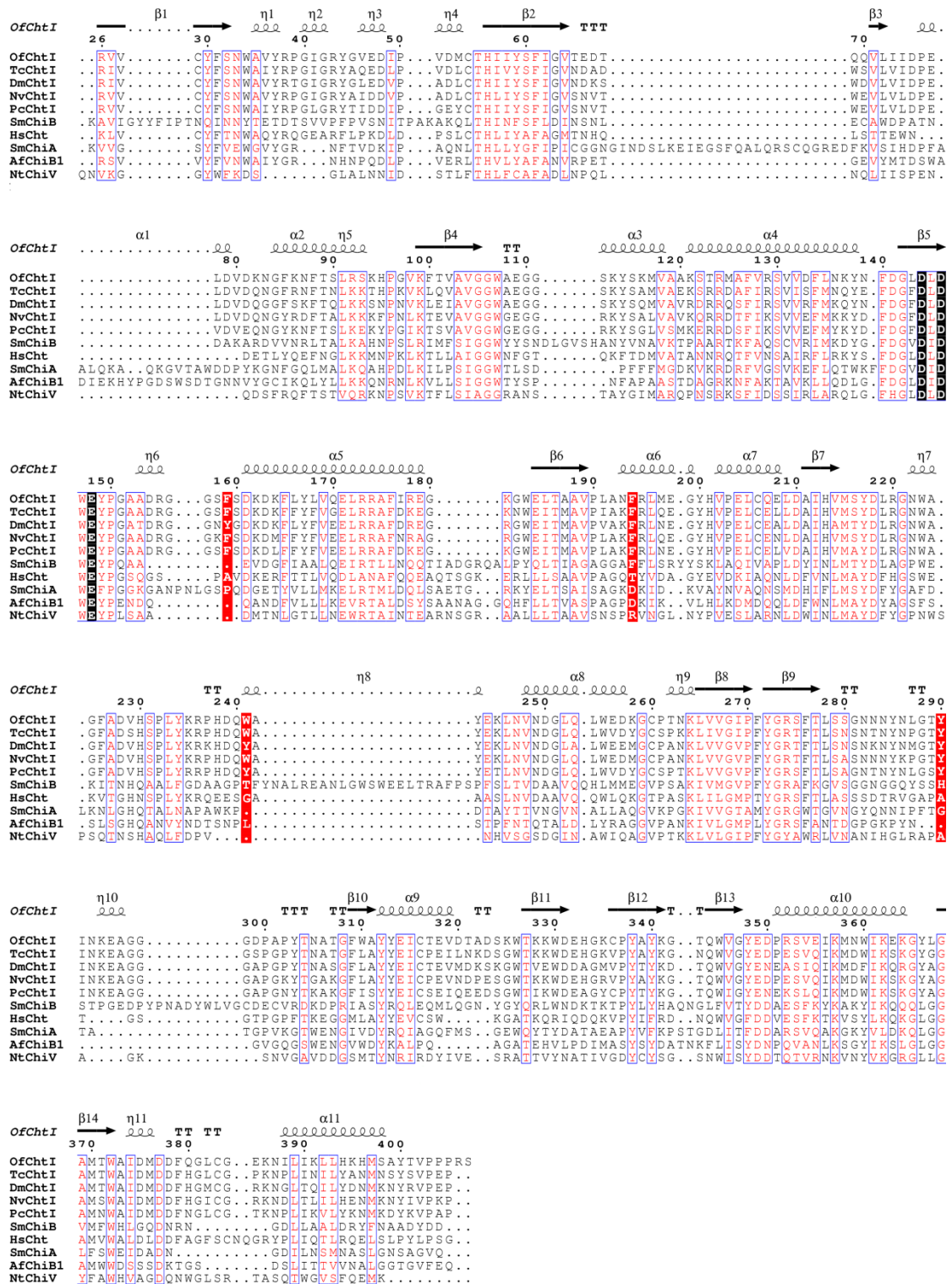


Figure S1

Structure-based sequence alignment of *OfChtI*-CAD with other GH18 chitinases from various species. The insect Group I chitinases include *OfChtI* from the lepidopteran insect *Ostrinia furnacalis* (Genbank ABB97081, PDB entry 3w4r), *PcChtI* from the hemipteran insect *Poophilus costalis* (Genbank AFW03959), *NvChtI* from the hymenopteran insect *Nasonia*

vitripennis (Genbank NP_001155084.1), *TcChtI* from the coleopteran insect *Tribolium castaneum* (Genbank AAV74190), and *DmChtI* from the dipteran insect *Drosophila melanogaster* (Genbank AAF54987). The other known chitinases include *SmChiA* (PDB entry 1ctn) and *SmChiB* (PDB entry 1e15) from *Serratia marcescens*, *HsCht* from *Homo sapiens* (PDB entry 1lg1), *AfChiB1* from *Aspergillus fumigatus* (PDB entry 1w9p), and *NtChiV* from *Nicotiana tabacum* (PDB entry 3alf). The secondary structure elements of *OfChtI*-CAD are shown and labeled. The conserved catalytic residues are labeled with black background blocks. The four aromatic residues, which form a hydrophobic plane on the surface of *OfChtI*-CAD, are highly conserved in the insect Group I chitinases but are absent in those from other species. They are labeled with red background blocks. The sequence alignment was generated by PROMALS3D (Pei *et al.*, 2008) and the picture was generated by ESPript (Gouet *et al.*, 2003).

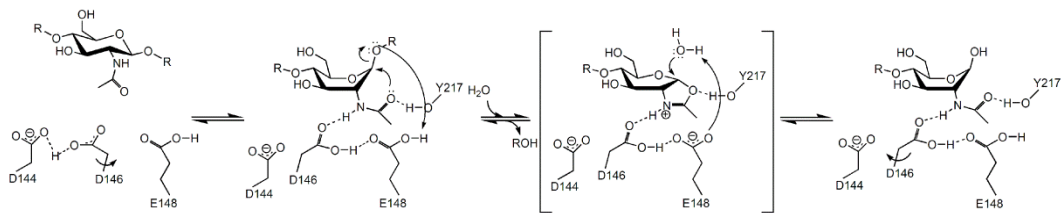


Figure S2

Proposed catalytic mechanism of family 18 chitinases, based on mechanism deduced by van Aalten *et al.* (2001).

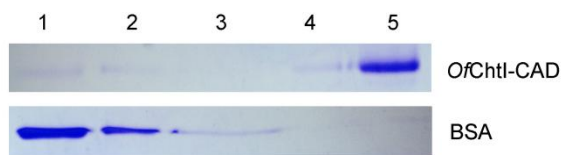


Figure S3

The binding of *OfChtI*-CAD to crystalline α -chitin. The mixtures (50 μ l) contained 10 μ g *OfChtI*-CAD and 1.0 mg crystalline α -chitin in 50 mM sodium phosphate buffer, pH 6.5. The mixtures were incubated at 303 K for 3 h and then centrifuged at 13 000g for 5 min. The supernatant was collected and the pellet was washed in 50 μ l of 50 mM sodium phosphate buffer, pH 6.5, 50 mM sodium phosphate buffer, pH 6.5, containing 1M NaCl, and 0.1 M acetic

acid in turn. After centrifugation of each wash, the supernatants were collected. Finally, the pellet and all collected supernatants were subjected to 10% SDS-PAGE. Lane 1, unbound fraction; lane 2, wash fraction in 50 mM sodium phosphate; lane 3, wash fraction in 50 mM sodium phosphate containing 1M NaCl; lane 4, wash fraction in 0.1 M acetic acid; lane 5, bound fraction. BSA was used as a control.