

Figure S1. CATs specifically interacted with HCPro.

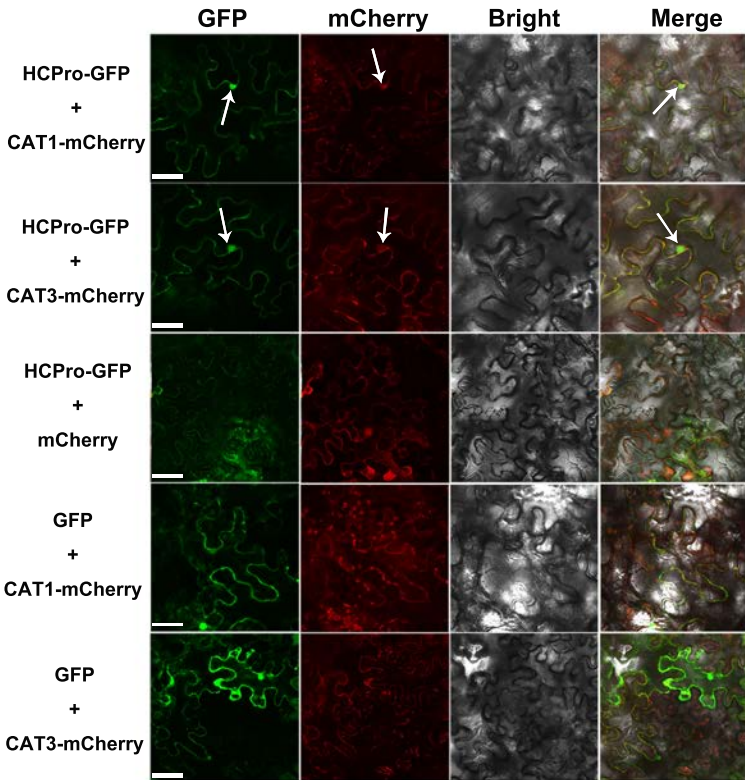


Figure S2. CAT1 and CAT3 colocalized to ChiVMV HCPPro in *N. benthamiana* leaves. Scale bars = 50 μ m.

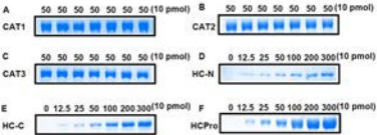


Figure S3. Coomassie brilliant blue staining of CAT1, CAT2, CAT3 and deletion mutants of HCPPro at the varying amounts used in this assay. HC-N (1–100 aa), HC-C (301–457 aa). The full length of HCPPro is marked as HCPPro.

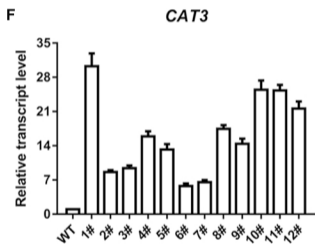
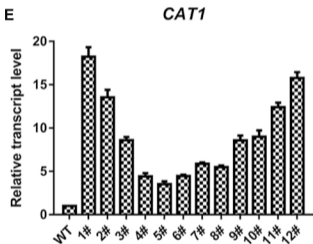
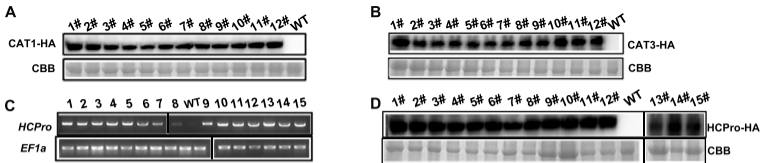


Figure S4. Identification of overexpression transgenic plants.

Western blot analysis of (A) CAT1 expression and (B) CAT3 expression by HA antibody. (C) Analysis of HCPPro expression by reverse transcription PCR and (D) Western blot. *EF1a* was used as loading control. qPCR analysis of (E) *CAT1* and (F) *CAT3* expression levels in T0 lines. Expression levels were standardized to *EF1a*, and result of WT was set at 1. Bars represent mean and standard deviation of values obtained from three biological repeats.

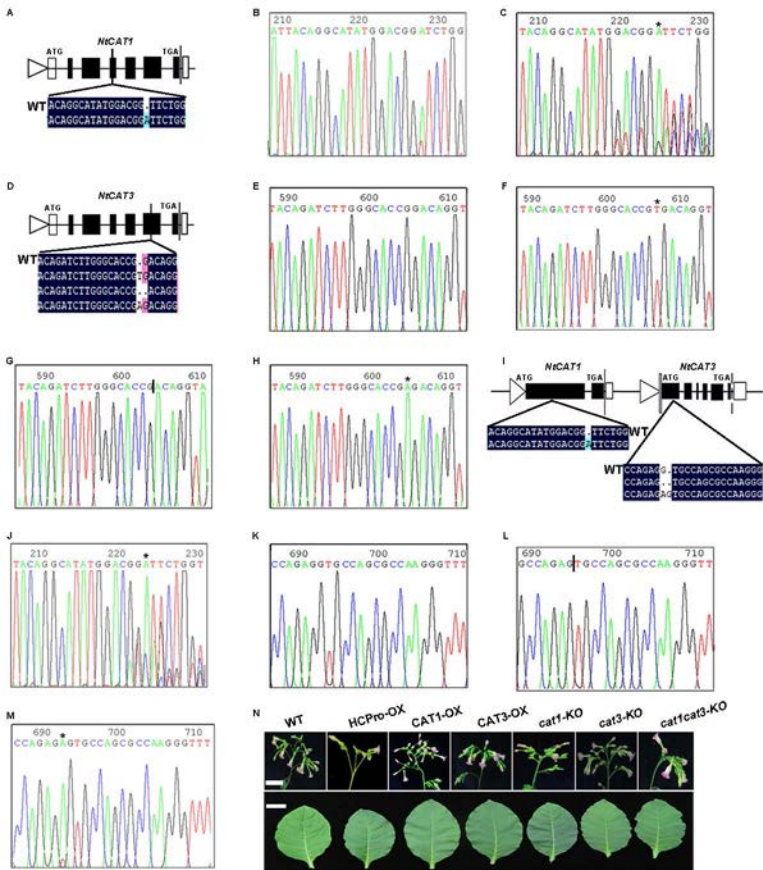


Figure S5. Phenotype and identification of *cat1*-KO, *cat3*-KO, *cat1cat3*-KO transgenic lines.

(A) Sequencing analysis of targeted region mutations of *CAT1* in representative T0 lines. Sanger

sequencing chromatogram of cloned *CAT1* in (B) WT and (C) *cat1*-KO plants. (D) Sequencing

analysis of targeted region mutations of *CAT3* in representative transgenic T0 lines. Sanger

sequencing chromatogram of cloned *CAT3* in (E) WT and (F), (G), (H) *cat3*-KO T0 plants.

(I) Sequencing analysis of targeted region mutations of *CAT1* and *CAT3* in *cat1cat3*-KO

representative T0 lines. (J) Representative Sanger sequencing chromatograms of target site

from the *CAT1* locus in *cat1cat3*-KO lines. Representative Sanger sequencing chromatograms

of *CAT3* target site from the *CAT3* locus (K) in WT and (L), (M) in *cat1cat3*-KO lines. Detected insert

base was marked in asterisk and delete base was marked by a vertical line. (N) Phenotypes of WT

and transgenic plant in T0 lines. Scale bars = 6 cm (upper panel) and 2.5 cm (lower panel).

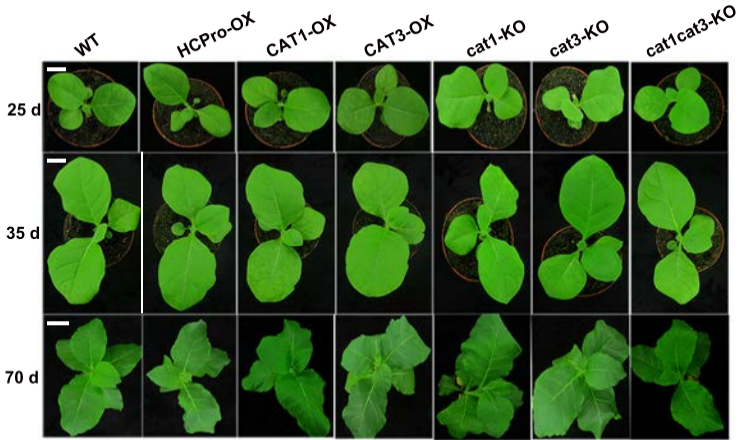


Figure S6. Phenotype of WT and transgenic lines at different growing stage in T1 lines.
Scale bars = 2.5 cm (upper and middle panel) and 8 cm (lower panel).

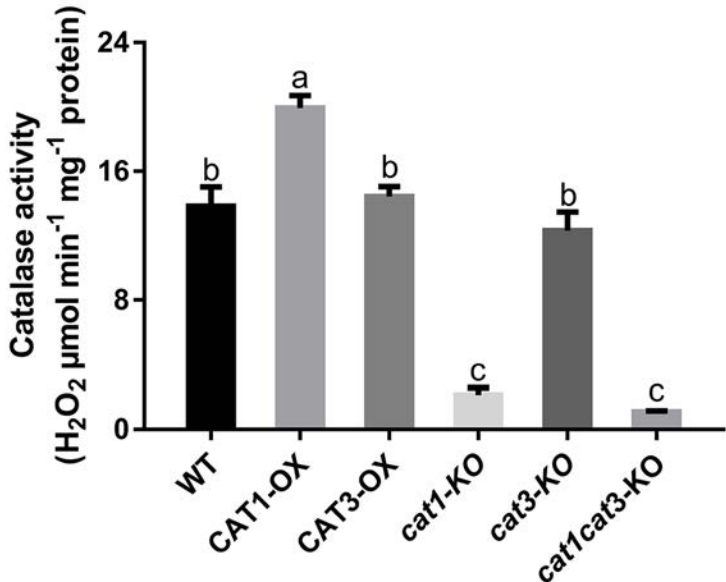


Figure S7. Catalase activity of WT and transgenic lines.

Bars represent mean and standard deviation of values obtained from three biological repeats. Significant differences ($P < 0.05$) are denoted by different lowercase letters.

Primer Names	Primer Sequences (5'-3')	Purpose
Yeast-CAT1-F	CCG GAATTC ATGGCAAGTG AAAAGTGGTT	Y2H
Yeast-CAT1-R	GCA CTGCAGT TACTGATGGGTGACTTCTC	Y2H
Yeast-CAT2-F	CCG GAATTC ATGGATCCCTCTAAGTTTCGA	Y2H
Yeast-CAT2-R	GCA CTGCAGT CACATTGTAGGCTTTAAAG	Y2H
Yeast-CAT3-F	CCG GAATTC ATGGATCCATAACAAGTATCG	Y2H
Yeast-CAT3-R	CGC GTCGACT CACATTGTGGGCCTTACAT	Y2H
Yeast-HCPro -F	CCG GAATTC ATGTCAGCAGGCGAGCTCTT	Y2H
Yeast-HCPro -R	CGC GGATCC CTAACCAACTCTGTACATT	Y2H
MBP-CAT1-F	CCG GAATTC ATGGCAAGTG AAAAGTGGTT	Pull down
MBP-CAT1-R	GCA CTGCAGT TACTGATGGGTGACTTCTC	Pull down
MBP-CAT2-F	CCG GAATTC ATGGATCCCTCTAAGTTTCGA	Pull down
MBP-CAT2-R	GCA CTGCAGT CACATTGTAGGCTTTAAAG	Pull down
MBP-CAT3-F	CCG GAATTC ATGGATCCATAACAAGTATCG	Pull down
MBP-CAT3-R	CGC GTCGACT CACATTGTGGGCCTTACAT	Pull down
GST-HCPro-F	CCG GAATTC ATGTCAGCAGGCGAGCTCTT	Pull down
GST-HCPro-R	CGC GTCGAC CTAACCAACTCTGTACATT	Pull down
GST-CAT1-F	CCG GAATTC ATGGCAAGTG AAAAGTGGTT	Pull down
GST-CAT1-R	GCA GTCGACT TACTGATGGGTGACTTCTC	Pull down
GST-CAT3-F	CCG GAATTC ATGGATCCATAACAAGTATCG	Pull down
GST-CAT3-R	CGC GTCGACT CACATTGTGGGCCTTACAT	Pull down
MBP-HCPro-F	CCG GAATTC ATGTCAGCAGGCGAGCTCTT	Pull down
MBP-HCPro-R	CGC GTCGAC CTAACCAACTCTGTACATT	Pull down
MBP-HC-N-F	CCG GAATTC ATGTCAGCAGGCGAGCTCTT	Pull down
MBP-HC-N-R	CGC GTCGAC GCTTTTAAGTAATTGTACAGCA	Pull down

MBP-HC-F	CCG GAATTC ATGCACTTTTTAAGCTTTAAA	Pull down
MBP-HC-R	CGC GTCGAC CGATCCATCATCATAAGTTACA	Pull down
MBP-HC-C-F	CCG GAATTC ATGCCAGTTTTGTCAGAATTCAA	Pull down
MBP-HC-C-R	CGC GGATCC CTAACCAACTCTGTACATT	Pull down
BiFC-CAT1 -F	TGCT TAGAA ATGGATCCATAACAAGTACCG	BiFC
BiFC-CAT1-R	CGC GTCGAC TATGCTTGGTCTCACA	BiFC
BiFC-CAT2-F	TGCT TAGAA ATGGATCCCTCTAAGTTTCG	BiFC
BiFC-CAT2-R	CGC GTCGAC CATTGTAGGCTTTAAAG	BiFC
BiFC-CAT3-F	TGCT TAGAA ATGGATCCATAACAAGTATCG	BiFC
BiFC-CAT3-R	CGC GTCGAC CATTGTGGGCCTTACAT	BiFC
BiFC-HCPro-F	CCG GAATTC ATGTCAGCAGGCGAGCTCTT	BiFC
BiFC-HCPro-R	CGC GTCGAC CTAACCAACTCTGTACATT	BiFC
CAT1-F	TCCACAAGATTACAGGCATA	qPCR
CAT1-R	AGCGGCAATAGAGTCATAG	qPCR
CAT2-F	CCAATTCCTTCTCGTGTCT	qPCR
CAT2-R	GTATCTGTCTTGCCTGTCA	qPCR
CAT3-F	AGGAGGAGCGAATCATAGT	qPCR
CAT3-R	TTCAATACCAAGCGACCAA	qPCR
EF1 α -F	TGCTGTAACAAGATGGATGC	qPCR
EF1 α -R	AGATGGGGACAAAGGGGATT	qPCR
Coat protein-F	AAACCCAGCCACAGTCTCGT	qPCR
Coat protein-R	ATCTCCGTCCATCATCACCC	qPCR
NbEF1 α -F	AGCTTTACCTCCCAAGTCATC	qPCR
NbEF1 α -R	AGAACGCCTGTCAATCTTGG	qPCR
GFP-F	ACATTATGGCAGACAAACAA	qPCR

GFP-R	TTACAAACTCAAGAAGCACC	qPCR
HCPPro - F	ATGTCAGCAGGCGAGCTCTT	RT-PCR
HCPPro - R	CTAACCAACTCTGTACATT	RT-PCR
RbohD-F	ACACGATCACATGGCTTCGAAAT	Pull down
RbohD-R	GTGTATTCCAACCCCAAGAGCA	Pull down
RbohF-F	AGTAAGCCTGGATACATAGAC	Pull down
RbohF-R	CAAGAAGGTGGTGTGAATAC	Pull down
35S-CAT1-F	CGC GTCGAC ATGGCAAGTG AAAAGTGGTT	Overexpression, RSS
35S-CAT1-R	TGCT CTAGAT ATGCTTGGTCTCACA	Overexpression, RSS
35S-CAT3-F	CGC GTCGAC ATGGATCCATAAAGTATCG	Overexpression, RSS
35S-CAT3-R	CGG GGTACC CATTGTGGGCCTTACAT	Overexpression, RSS
35S-HCPPro-F	CGC GTCGAC ATGTCAGCAGGCGAGCTCTT	Overexpression, RSS
35S-HCPPro-R	CGG GGTACC CTAACCAACTCTGTACATT	Overexpression, RSS
DT1- CAT1-BsF	ATATATGGTCTCGATTACAGGCATATGGACGGATCGTT	Knock out
DT1-CAT1-F0	TGACAGGCATATGGACGGATCGTTGTTTTAGAGCTAGAAATAGC	Knock out
DT2-CAT1-R0	ATATATGGTCTCGATTGATCCGTCCATATGCCTGTAAGAGTT	Knock out
DT2-CAT1-BsR	ATTATTGGTCTCGAAACGATCCGTCCATATGCCTGTAAGACAA	Knock out
DT1- CAT3-BsF	ATATATGGTCTCGATTGCAGATCTTGGGCACCGGACGTT	Knock out
DT1-CAT3-F0	TGCAGATCTTGGGCACCGGACGTTTTAGAGCTAGAAATAGC	Knock out
DT2-CAT3-R0	AACGTCCGGTGCCCAAGATCTGCAATCTCTTAGTCGACTCTAC	Knock out
DT2-CAT3-BsR	ATTATTGGTCTCGAAACGTCCGGTGCCCAAGATCTGCAA	Knock out
DT2-CAT3-R0	AACGAGGTGCCAGCGCCAAGGGCAATCTCTTAGTCGACTCTAC	Knock out
DT2-CAT3-BsR	ATTATTGGTCTCGAAACGAGGTGCCAGCGCCAAGGGCAA	Knock out
U626-IDF	TGTCCAGGATTAGAATGATTAGGC	Knock out
U629-IDR	AGCCCTCTTCTTTTCGATCCATCAAC	Knock out

Cas-CAT1-F	ACGTGTCCCT GAACGTGTTG	Knock out
Cas-CAT1-R	ACTTCACATAGGTTGATTC	Knock out
Cas-CAT3-F1	CTTGGGCCAAACTATCTGCA	Knock out
Cas-CAT3-R1	GACGCAATCTTCTGACCAAG	Knock out
Cas-CAT3-F2	ATGGATCCATAACAAGTATCGT	Knock out
Cas-CAT3-R2	GACGCAATCTTCTGACCAAG	Knock out
HPro-GFP-F	CGC GTCGAC ATGTCAGCAGGCGAGCTCTT	Subcellular location
HPro-GFP-R	CGC GGATCC CTAACCAACTCTGTACATT	Subcellular location
CAT1-mCherry-F	CCG GTCGAG ATGGATCCATAACAAGTACCG	Subcellular location
CAT1-mCherry-R	CGC GTCGAC TATGCTTGGTCTCACA	Subcellular location
CAT3-mCherry-F	CCG GTCGAG ATGGATCCATAACAAGTATCG	Subcellular location
CAT3-mCherry-R	CGC GTCGAC CATTGTGGGCCTTACAT	Subcellular location

Table S1. Primers used for construction of vectors and real-time PCR analysis.