

Figure S1

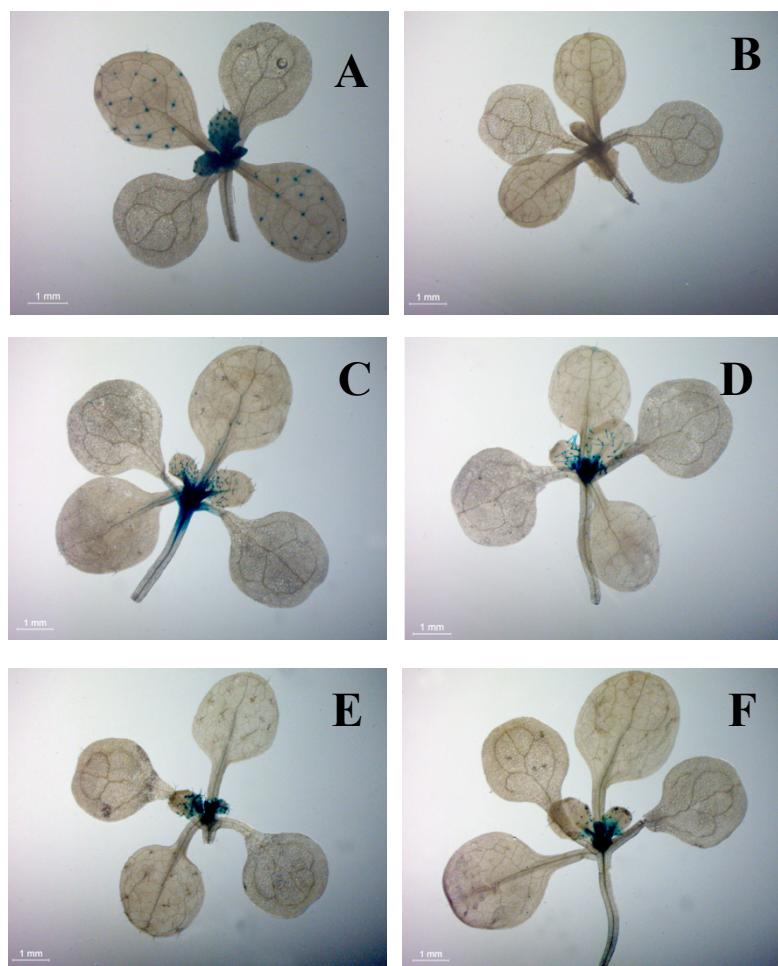


Figure S1: *TRY* but not *GL2* or *CPC* expression depends on *TTG2*

Expression pattern of the 5' regulatory regions of the *GL2*, *CPC* and *TRY* promoter in Ler and *ttg2* background as revealed by GUS reporter gene expression. Whole rosettes are shown. pTRY:GUS Ler (A), pTRY:GUS *ttg2* (B), pCPC:GUS Ler (C), pCPC:GUS *ttg2* (D), pGL2:GUS Ler (E), pGL2:GUS *ttg2* (F). Bars indicate the magnification of the images.

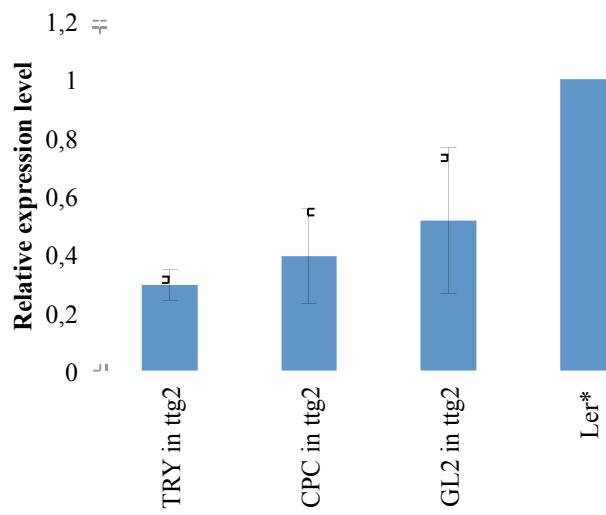


Figure S2: Analysis of TRY, CPC and GL2 expression by Real Time PCR.

TRY, CPC and GL2 expression in 10-day-old leaves three and four in *ttg2* mutants were analyzed by quantitative real-time PCR. *The expression levels were normalized to that of the respective genes in Ler. The 18s-RNA was used as an endogenous control. Error bars indicate the standard deviation of two biological replicas including three technical replicas. The expression of TRY, CPC and GL2 were significantly reduced compared to wild type (Student's T-test, $p<0,01$).

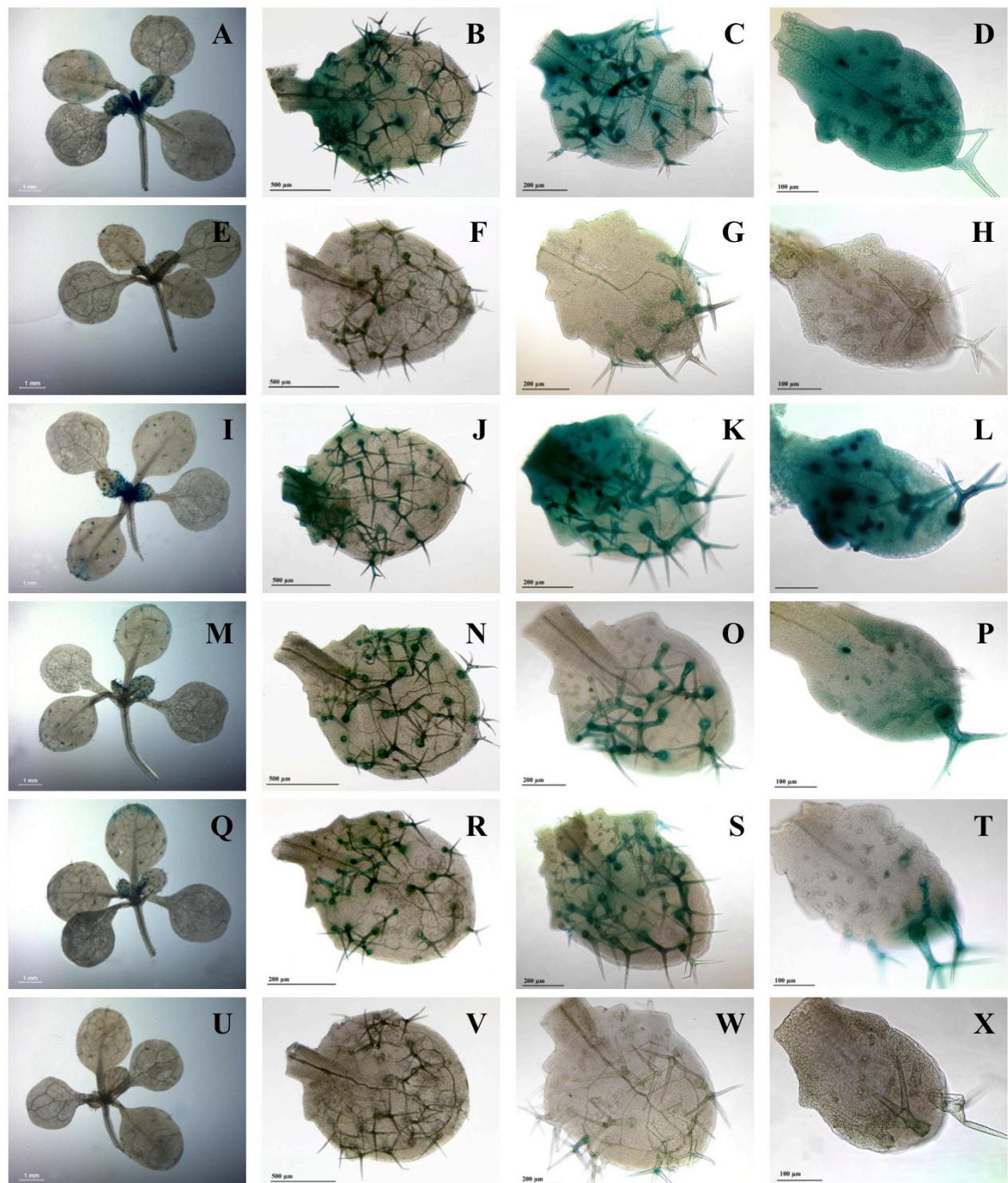


Figure S3: Expression analysis of *TRY* promoter fragments carrying mutations in W-boxes.

The expression pattern is shown for 10 days old plants: the whole rosette, and leaf number three to five to document different leaf developmental stages with leaf number three representing a mature leaf, leaf number four for an intermediate developmental stage and leaf number five to show a young leaf. A-D) pTRY-A3,B:GUS; E-H) pTRY-A3,B (Δ 234-176):GUS; I-L) pTRY-A3,B:GUS; M-P) pTRY-A3,B(mW1):GUS. The examples show leaves lacking the TRY expression at the leaf base. Q-T) pTRY-A3,B(mW2):GUS. These examples show trichome specific expression but no epidermal expression at all stages. U-X) pTRY-A3,B(mW1mW2):GUS.

Figure S4

W1W2	GAGT GTCAA CGACAAGTCTACACAAAGGGTAAGAG GTCAA CAAG
mW1mW2	GAGTCCC GGG CGACAAGTCTACACAAAGGGTAAGAG CCC GGG CAAG
mW1W2	GAGTCCC GGG CGACAAGTCTACACAAAGGGTAAGAG GTCAA CAAG
W1mW2	GAGT GTCAA CGACAAGTCTACACAAAGGGTAAGAG CCC GGG CAAG
W1	TTTGAGT GTCAA CGACAAG
mW1	TTTGAGT CCC GGG CGACAAG
W2	GTAAGAG GTCAA CAAGACC
mW2	GTAAGAG CCC GGG CAAGACC
1xW2*	TTATTCCAGCCATCAAAAG TTGAC CAATAAT
m19**	TTATTCCAGCCATCAAAAG TAGAC CAATAAT

Figure S4: Sequences of wild type and mutated W-boxes of the *TRY* promoter.

The region of the *TRY* promoter containing the wild-type and mutated sequences of the two W-boxes are shown. The W-Boxes are shown in red and the mutated W-boxes in a lighter grey. * The sequence 1xW2 was published previously and is identical to the region of the parsley *PR1-1* promoter that contains the W-Box W2 ([Ciolkowski et al., 2008](#)). ** m19 is a base substitution variant of 1xW2 that abolish binding specificity ([Ciolkowski et al., 2008](#)).

Figure S5

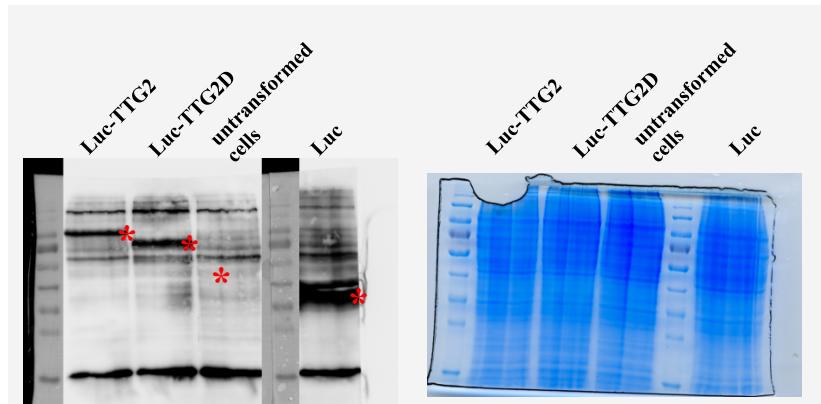


Figure S5: Integrity and functionality of Luc-TTG2 and Luc-TTG2D fusion proteins.
Western blot (left) and the corresponding Coomassie gel (right) showing the protein extract of HEC cells expressing Luc-TTG2 and Luc-TTG2D proteins. The luciferase antibody detects the two proteins at the expected size indicating that the presence of full-length fusion proteins. Asterisk (*) indicates the respective expressed protein band.

Figure S6

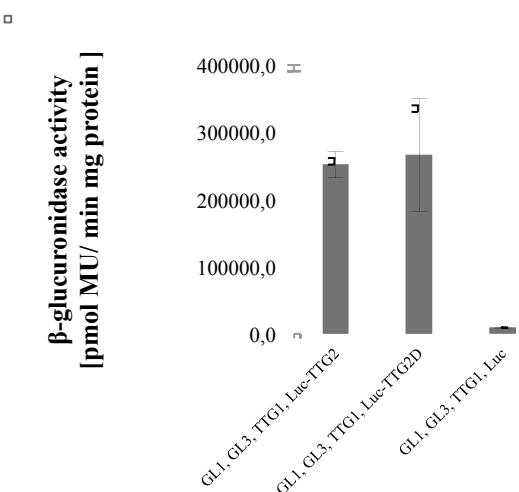


Figure S6: Expression levels of promoter:GUS constructs
Arabidopsis cell suspension cultures were transformed with the pTRY:GUS construct along with 35S:cDNAs of GL1, GL3, TTG1 and TTG2. The relative expression levels of the promoter:GUS constructs were determined by β -glucuronidase assays in three independent experiments. Error bars indicate the standard deviation.

Figure S7

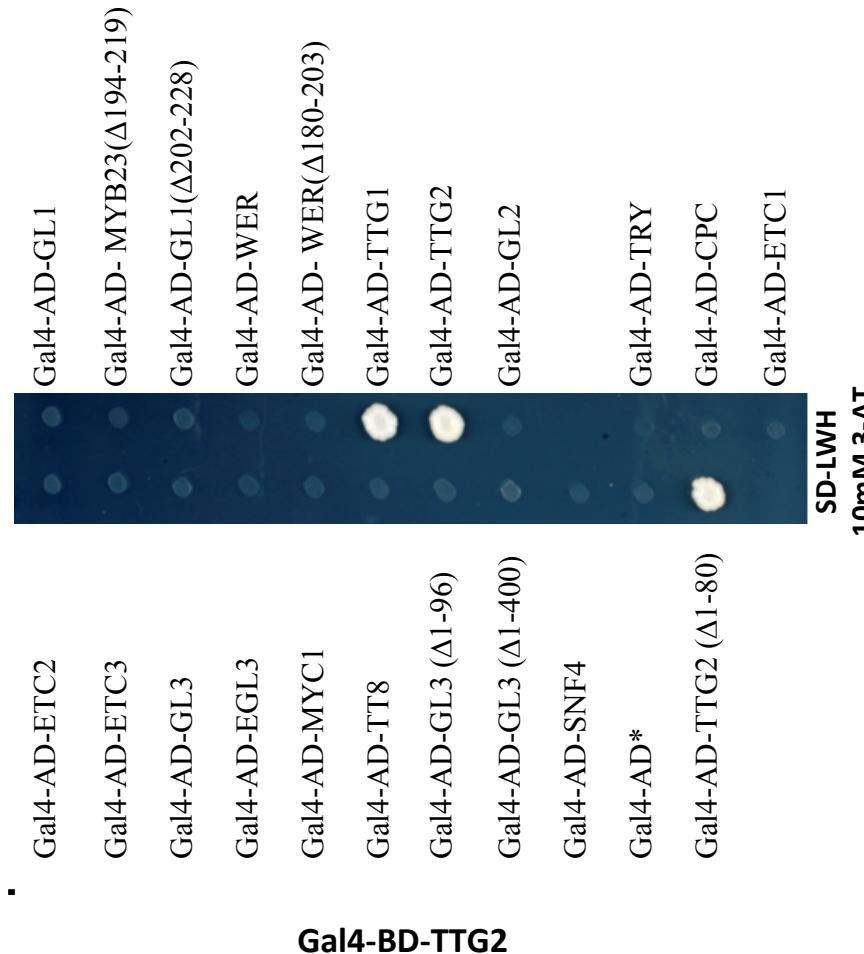
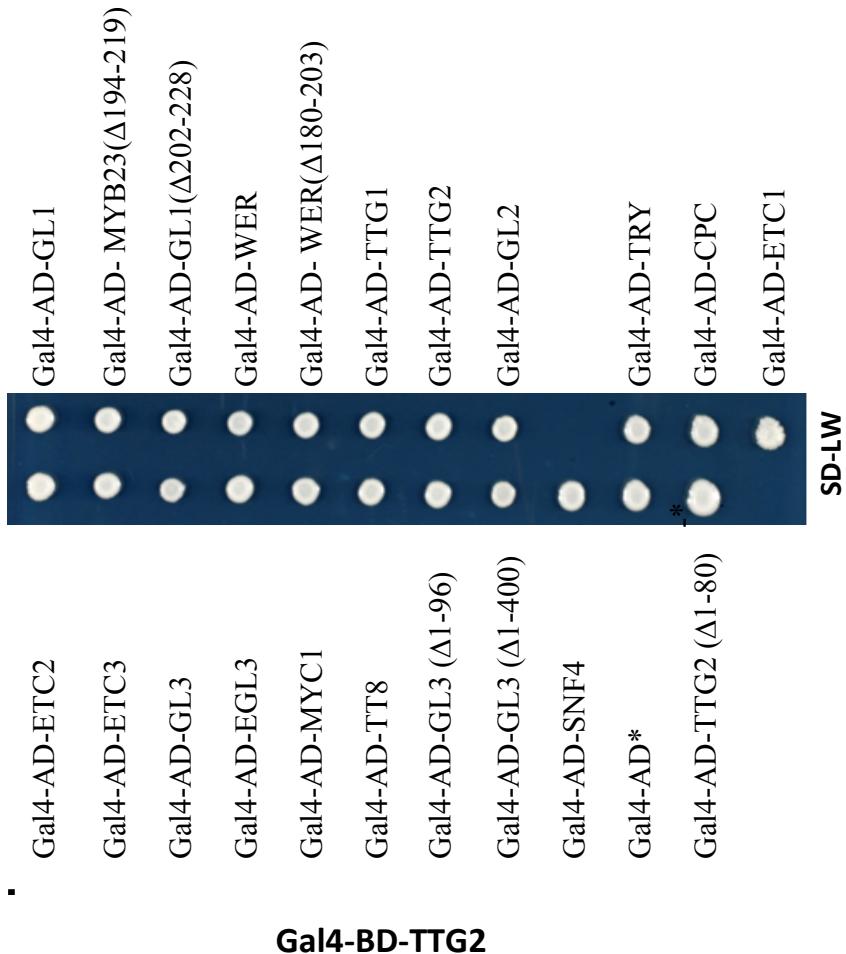


Figure S7: Yeast two hybrid interactions of TTG2-BD with other trichome patterning proteins.
Interaction of TTG2 fused to the DNA binding domain with other trichome and root hair patterning proteins
as indicated. Asterisk (*) indicates control construct without CDS fusion.

Figure S8

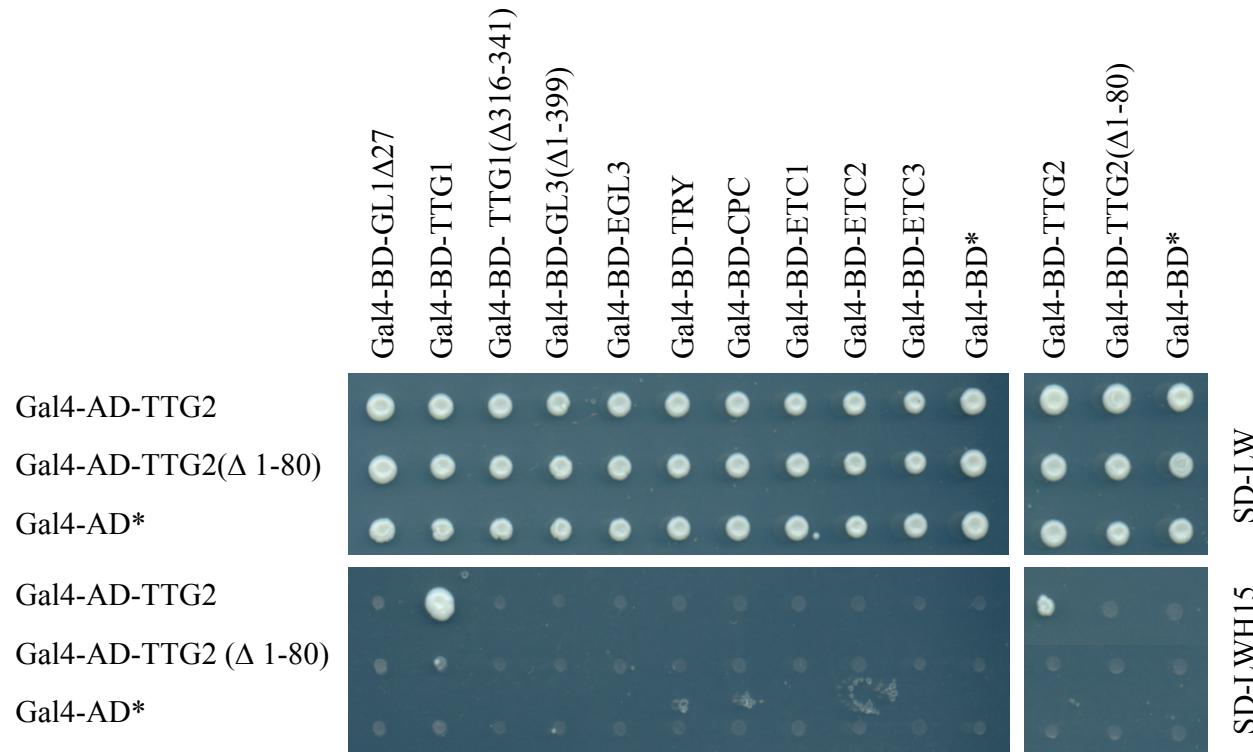


Figure S8: Yeast two hybrid interactions of TTG2-AD with other trichome patterning proteins.
Interaction of TTG2 fused to the activation domain with other trichome and root hair patterning proteins as indicated.

Figure S9

	template	fragment No.	Primer Code	Primer Name	
pTRY-A3(Δ234-176)	Pr15	Pr181	19 81	TRY-F3-sen-attB1 TRY-F4-asen-attB2 (ohne WRKYs)	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG GGGGACCACTTGTACAAGAAAGCTGGGTCTCAAATACACATGGAGATG
pTRY-A3(Δ195-176)	Pr15	Pr182	19 92	TRY-F3-sen-attB1 Try-Linker69-PCR-rev (mit W-Box2)	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG GGGGACCACTTGTACAAGAAAGCTGGGTGTGACCTCTAACCC
pTRY-A3-fragment I (mW1)	Pr15	Pr154	19 161	TRY-F3-sen-attB1 MP-Pr23-mutW-BOX1-Fr1-rev	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG
pTRY-A3-fragment II (mW1)	Pr15	Pr155	162 20	MP-Pr23-mutW-BOX1-Fr2-for TRY-F3-asen-attB2	GTATTIGAGTCGGCGGACAAGTCTACAAAAGGG GGGGACCACTTGTACAAGAAAGCTGGTTAAGAAGTGTGTGTGGTCT
pTRY-A3 (mW1)	Pr154+Pr155	Pr168	19 20	TRY-F3-sen-attB1 TRY-F3-asen-attB2	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG GGGGACCACTTGTACAAGAAAGCTGGTTAAGAAGTGTGTGGTCT
pTRY-A3-fragment I (mW2)	Pr15	Pr156	19 163	TRY-F3-sen-attB1 MP-Pr23-mutW-BOX2-Fr1-rev	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG GGTCTTGCCTGGCTTACCCCTTGTTAGACTTG
pTRY-A3-fragment II (mW2)	Pr15	Pr157	164 20	MP-Pr23-mutW-BOX2-Fr2-for TRY-F3-asen-attB2	CAAGTCTACACAAGGGTAAGAGCCCGGCAAGACC GGGGACCACTTGTACAAGAAAGCTGGTTAAGAAGTGTGTGTGGTCT
pTRY-A3 (mW2)	Pr156+Pr157	Pr169	19 20	TRY-F3-sen-attB1 TRY-F3-asen-attB2	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG GGGGACCACTTGTACAAGAAAGCTGGTTAAGAAGTGTGTGTGGTCT
pTRY-A3-fragment I (mW1mW2)	Pr168	Pr186	19 163	TRY-F3-sen-attB1 MP-Pr23-mutW-BOX2-Fr1-rev	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG MP-Pr23-mutW-BOX2-Fr1-rev
pTRY-A3-fragment II (mW1mW2)	Pr168	Pr187	164 20	MP-Pr23-mutW-BOX2-Fr2-for TRY-F3-asen-attB2	MP-Pr23-mutW-BOX2-Fr2-for GGGGACCACTTGTACAAGAAAGCTGGTTAAGAAGTGTGTGTGGTCT
pTRY-A3 (mW1mW2)	Pr186+Pr187	Pr192	19 20	TRY-F3-sen-attB1 TRY-F3-asen-attB2	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG GGGGACCACTTGTACAAGAAAGCTGGTTAAGAAGTGTGTGTGGTCT
pTRY-A3-B-fragment I (mW1mW2)	R260	Pr205	19	TRY-F3-sen-attB1	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG
pTRY-A3-B-fragment II (mW1mW2)	R260	Pr206	- -	Fusion 15+33-rev Fusion 15+33-for	GCAAAACTAATAGTAAGAAGTGTGTGTGG CCACACAACTCTTACTATAGTTTG
pTRY-A3-B (mW1mW2)	R260a + R260b	Pr207 = Pr195 (mW1mW2)	179	5'-TRY-179-rev-attB2	GGGGACCACTTGTACAAGAAAGCTGGTCAGCTTATTGAAGTAAGAAAAGAAAAATAGAGAG
pTRY-A3-B	genomic DNA Ler	Pr195	19 179	TRY-F3-sen-attB1 5'-TRY-179-rev-attB2	GGGGACAAGTTGTACAAAAAAGCAGGCTGTGAAACTAATTGTTACTATATGG GGGGACCACTTGTACAAGAAAGCTGGTCAGCTTATTGAAGTAAGAAAAGAAAAATAGAGAG

MP-WRKY-for-EcoRI catccatgt**GAATT**Cgagt
MP-WRKY-for-KpnI catccatgt**GGTACC**gagt
MP-WRKY-for-BamHI catccatgt**GGATCC**gagt
MP-WRKYrev-KpnI **eG**TACCAACT**A**T**A**T**G**T**A**AG**A**GT**T**TT**G**
MP-WRKYrev-BamHI **eG**GAT**C**CAACT**A**T**A**T**G**T**A**AG**A**GT**T**TT**G**
MP-WRKYrev-XbaI **cT**CT**G**AG**A**ACT**A**T**A**T**G**T**A**AG**A**GT**T**TT**G**

W1W2 GAGTGTCACGACAAGTCTACACAAAGGGTAAGAGGTCAACAAAG
mW1W2 GAGTCCC**G**CAACAGTCTACACAAAGGGTAAGAG**CCC**GG**C**AAG
mW1W2 GAGTCCC**G**CAACAGTCTACACAAAGGGTAAGAGGTCAACAAAG
W1mW2 GAGTGTCACGACAAGTCTACACAAAGGGTAAGAG**CCC**GG**C**AAG
W1 TTTGAGTGTCACGACAAG
mW1 TTTGAGT**CCC**GG**C**AAG
W2 GTAAAGGGTCAACAAGACC
mW2 GTAAAGAG**CCC**GG**C**AAGACC

MP-TTG2-WRKY-Domänen-for GGGGACAAGTTGTACAAAAAAGCAGGCTTAACGGGGATAGATCTTCT
MP-TTG2-WRKY-Domänen-rev GGGGACCACTTGTACAAGAAAGCTGGGTACTAGAGCAAATGATGATTATG

MP-Asl-Renilla-rev TTGGCGGCCATCCCCCTGCTCGTCTTC
MP-KpnI-Renilla-for TTGGTACCATGACAGCAAGGTACGAC

W18D-fwd CTG GTT CCG CGG GTG CCT ACT GAA ACA TCG GAC AC
W18D-rev TCC TCC GGT ACC TCA TGT AGC ATC CCC TTC AGA AGC ATT