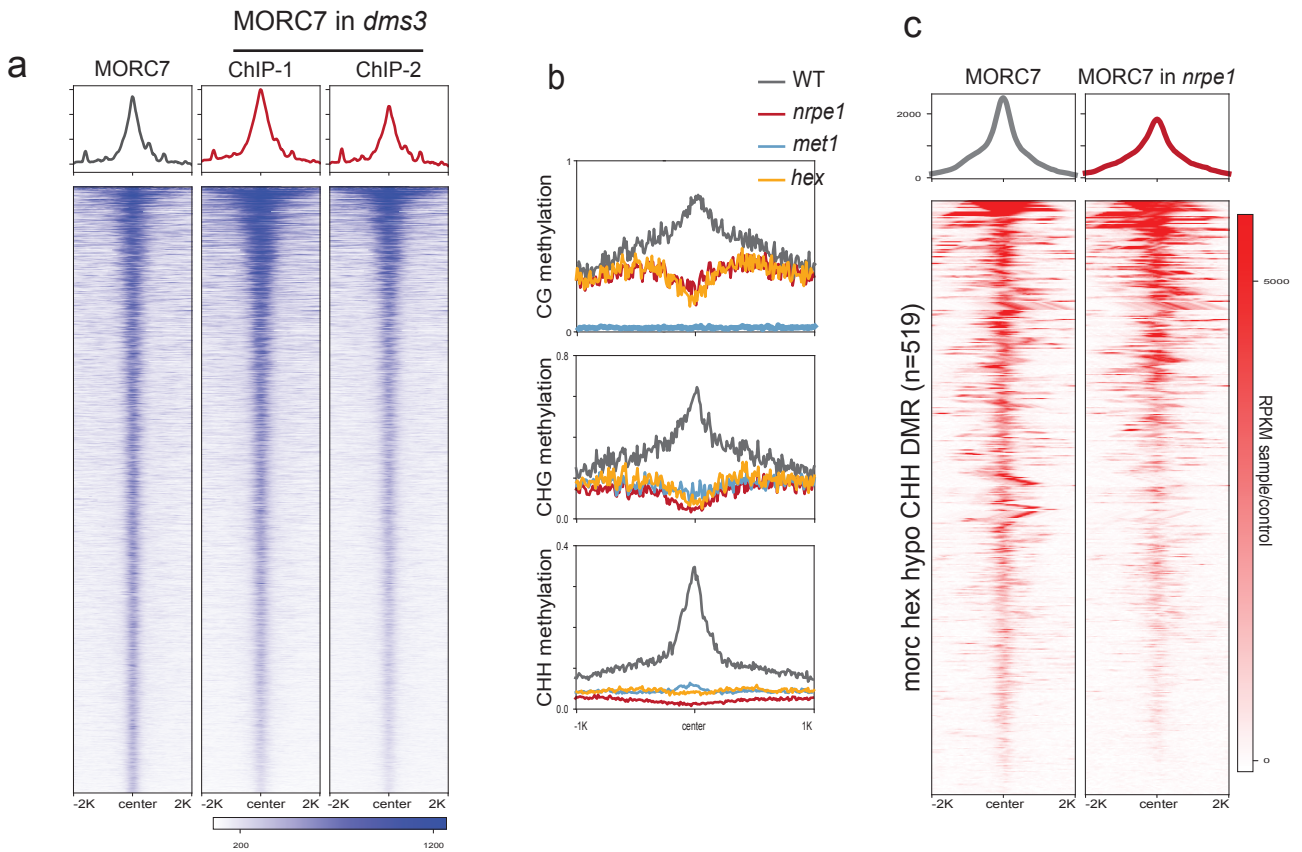
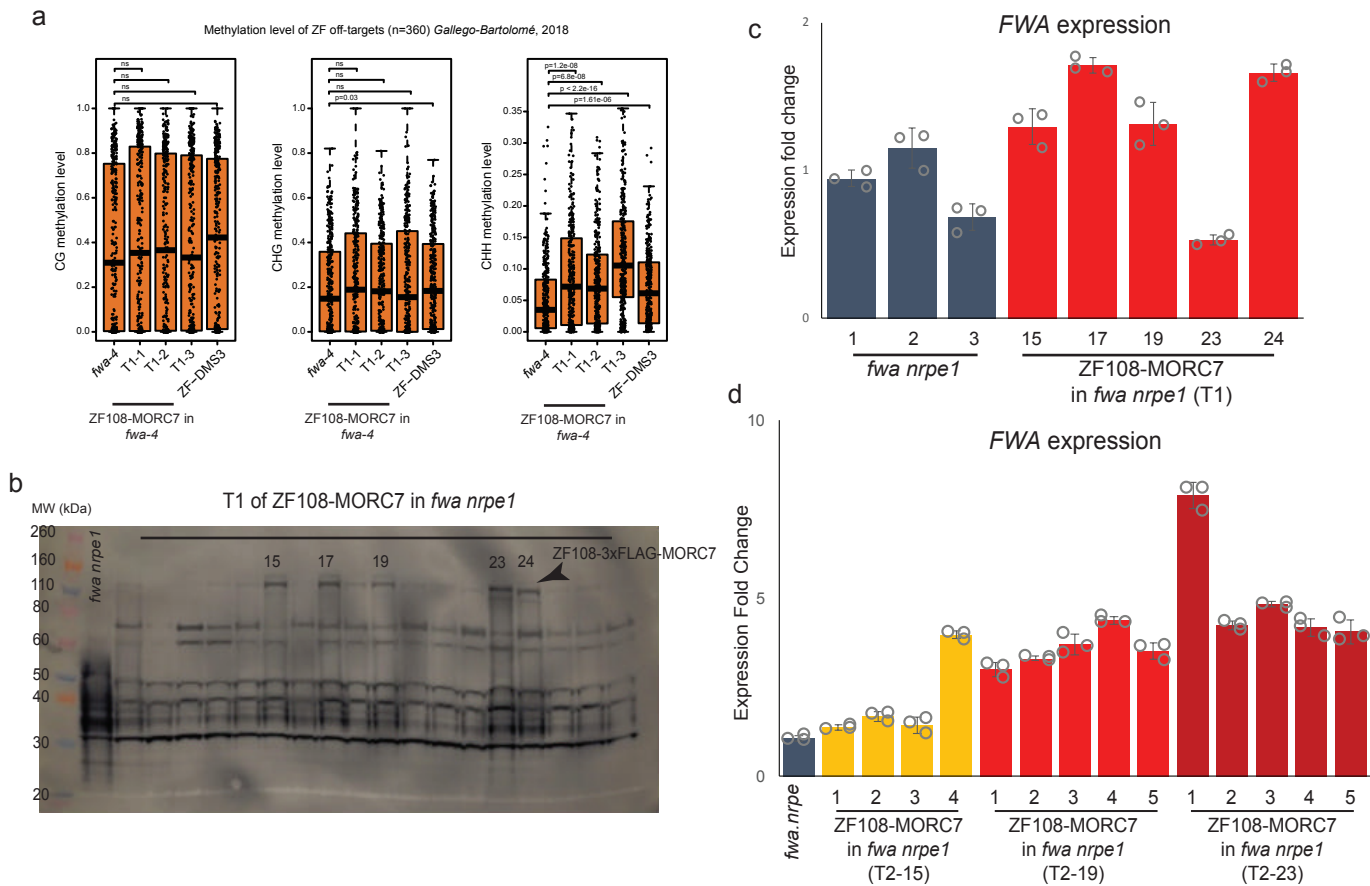


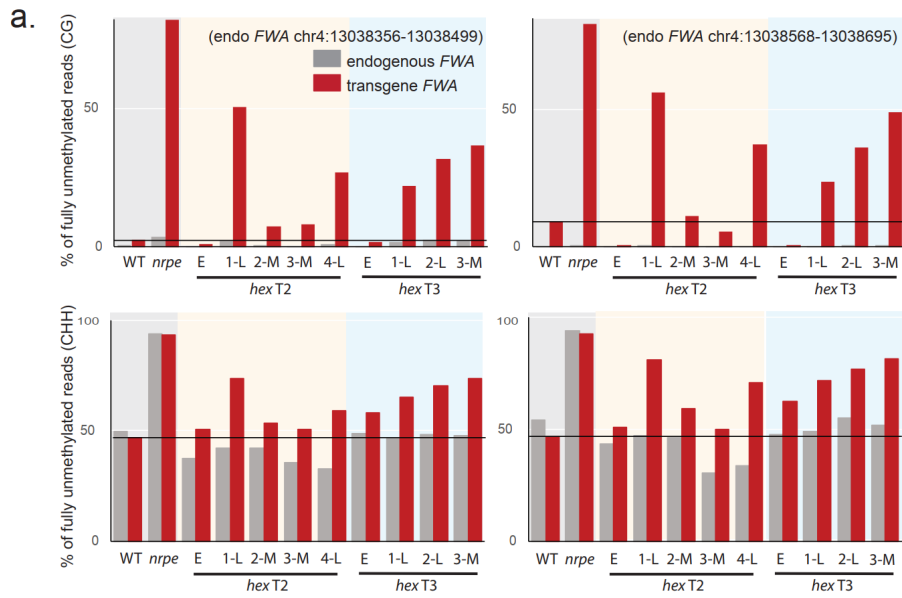
Supplementary Fig. 1. MORC4 and MORC7 chromatin localization, related to Figure 1. **a** Pearson correlation of MORC4-FLAG, MORC7-FLAG and NRPE1 ChIP-seq. **b** Overlap between MORC4 and MORC7 binding sites (top) ($p = 1.474663e-104e$, Hypergeometric test), enrichment of MORC4 and MORC7 ChIP signals (RPKM sample/control) over MORC7 and MORC4 specific regions respectively (bottom). **c** Enrichment of CG (left panel), CHG (middle panel) and CHH (right panel) DNA methylation over MORC7 unique (blue), MORC7-NRPE1 common (red) and NRPE1 unique (orange) peaks. **d** Screenshot of a representative locus of NRPE1 unique (left) and MORC7 unique peak (right). **e** Boxplot showing the distance of MORC7 unique, MORC7-NRPE1 common and NRPE1 unique peaks to the transcription start site (TSS) of the nearest gene. **f** Profile and heatmap of Pol II ChIP signals (RPKM sample/control) over MORC7 unique, MORC7-NRPE1 common and NRPE1 unique peaks. **g** Venn diagram showing regions that are upregulated in *hex*. Regions that have MORC7 bound within 1Kb (MORC7 bound) are in light blue (n=128) and regions that are MORC7 unbound are in dark blue (n=775). **h** DNA methylation over MORC7 bound regions (left) and MORC7 unbound regions (right). **i** Expression (log₂RPKM) over the MORC7 bound (left) and MORC7 unbound (right) regions in WT (blue), *hex* (red) and *nrpe1* (yellow).



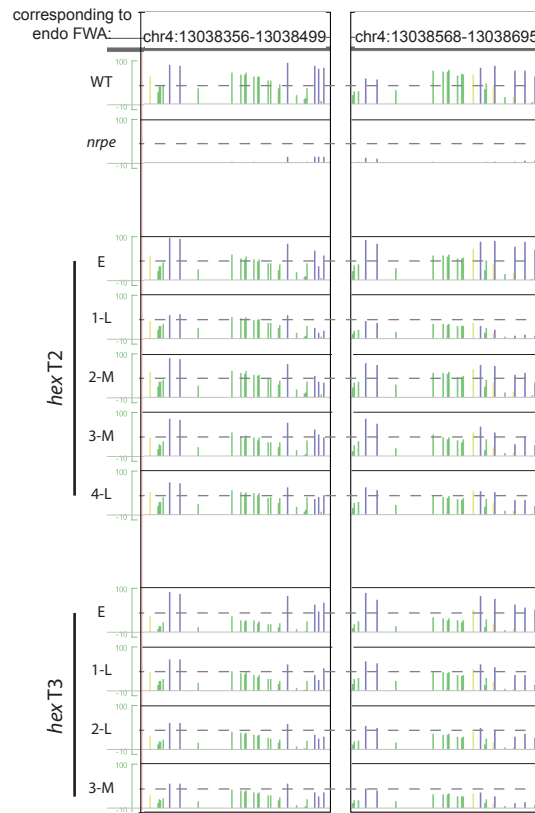
Supplementary Fig. 2. MORC7 binding in *dms3-4* and over *morc hex* hypo-CHH DMRs (fragile sites), related to Figure 2. **a Enrichment of MORC7 ChIP signals (biological replicate of Fig. 2d) (RPKM sample/control) in *dms3* (right) and its corresponding WT control (left). **b** Methylation level over the *hex* hypo-CHH DMRs in WT (grey), *hex* (orange), *nrpe1* (red) and *met1* (blue) (n = 519). **c** Enrichment of MORC7 ChIP signal (RPKM sample/control) over the same *hex* hypo-CHH DMR in **b**.**



Supplementary Fig. 3. MORC7 mediated repression on *FWA* is dependent on RdDM, related to Figure 4. **a** Boxplot showing 360 ZF ectopic sites that gained DNA methylation in the three independent T1 plants showed in Fig 5a. Wilcoxon test was used to determine significance. (ns: p value > 0.05). **b** anti-FLAG western blot showing the expression of ZF108-3xFLAG-MORC7 in the T1 plants displayed in **a**. Arrow indicates the band with expected size. T1 lines with their corresponding numbers were chosen for further analysis. **c** qRT-PCR analysis showing *FWA* expression level relative to *fwa nrpe1* in five independent T1 plants. Error bars represent the mean \pm s.d. of 3 technical replicates. **d** qRT-PCR analysis showing *FWA* expression level relative to *fwa nrpe1* in the T2 progenies of three T1 plants. Error bars represent the mean \pm s.d. of 3 technical replicates.

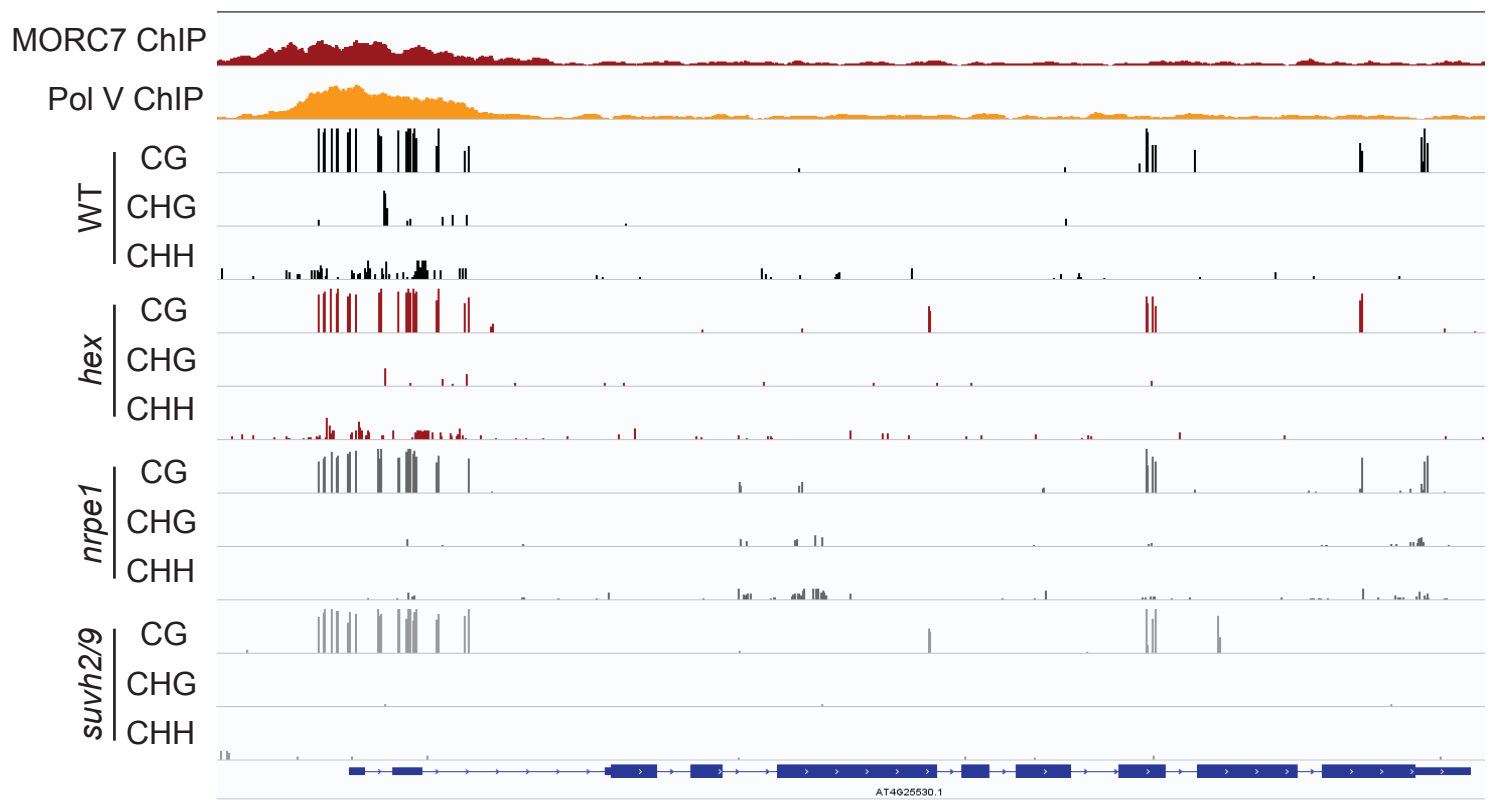


b.



Supplementary Fig. 4. Establishment of DNA methylation is attenuated over *FWA* transgene.

a Percentage of fully CG unmethylated reads (upper panels) or fully CHH unmethylated reads (lower panels) from the endogenous *FWA* (grey) and *FWA* transgene (red) over two regions within the promoter. The dark lines indicate the levels of fully unmethylated *FWA* transgene in WT. E: early flowering; M: Intermediate flowering; L: late flowering. **b** Methylation tracks of the *FWA* transgene over the same regions as in a. CG:blue; CHG, yellow; CHH, green.



Supplementary Fig. 5. Methylation over endogenous *FWA* promoter is not affected in *hex*, related to Figure 5. Screenshot of MORC7 ChIP seq (crimson), Pol V ChIP (orange) over *FWA* locus (AT4G25530) as well as DNA methylation level in WT (black), *hex* (crimson), *nrpe1* (dark grey) and *suvh2/9* (light grey) in CG, CHG and CHH context.

Supplementary Table 1: Primers used in this study

Genotyping primers:

<i>morc1-2</i> SAIL_893_B06	TTGCAGTTTGAACCAAATC	TGAGTTTTGACGACGATGATG
<i>morc2-1</i> SALK_072774 C	CTACTCAGAGCGTTGGCATTG	GTTGTAGCTGTATGGGGCTTG
<i>morc4-1</i> GK-249F08	TCAGGAAAGATTTACGAATTG	ACCTGCAGAACTTCCCAATC
<i>morc5-1</i> SALK_049050 C	GTTGGGATAGATAAGGCGACC	TGTCGAGAAATCGTTCCTTTG
<i>morc6-3</i> GABI_599B06	ACATCTTCCAATGGCTGAATC	GCTGGTGTCACTTCTTCATCC
<i>morc7-1</i> SALK_051729	GTCGAAAGGATGTGAGAAACG	TTCCATTCAATTGCTTGGTTC
<i>nrpe1-11</i> SALK_029919	ATTTCTTCTTTGATGGGGGAG	TGTCGTGGATATGACCATTG
<i>nrpd1-4</i> SALK_083051	TGGGTTTGCCATTTTCATATC	GCATGCTTGAGTAAAAGGTGC
<i>suvh2</i> SALK_079574	GCCCAGGTGTTGTTTCTGTC	CCATATCGCTCCTCCGTCTA
<i>suvh9</i> SALK_048033	CACACTAGTAGCTATTGGCTCCCCATTA	TCTCTGATAAACTCGTCAAACCCATTTCG
<i>dms3-4/idn1-1</i>	CTGCCTTGAAAACCTGAGGTAC	TCTGATAAGATGCATTCCATGG
<i>met1-3</i> CS16394	GATTGTGTCTCTACTACAGAGGC	GTAAAGCTCATTCATAGCCTTGC
<i>nrpe1-1</i>	CGTGGCATCATGAAACATTCACTGC	CCATCTGATGCTCTATCTTTGATTGAGATGT

BS PCR:

FWA region 1 chr4:13038356- 13038499	CTCATATATACCTTATCCCATTCAACATTCATA	AAGATYTGATATTTGGYTGGAAAAAAYAATAA TAAT
FWA region 2 chr4:13038568- 13038695	CRCTCTTTATCCCATTCAACATTCATAC	TTTGGTTGAAAAAATAATAAAAAATTTGATTGT YAGTAT

Cloning primers:

Primer	Description	Sequence
JP22356	N terminal ZF infusion into pDT Forward	AAAGACCGGTCAATTGGACGTCATGGTTGGGTACC AACAAATGG
JP22357	ZF-linker-DMS3 (overlap Reverse)	GATACATCCCACCAGAGCCTCCGAGTCCGGTGTGA GTCCTTTG
JP22358	ZF-linker-MORC7 (overlap Forward)	AAAGGACTCACACCGGACTCGGAGGCTCTGGTGG GATGGACAACAGTATTCACGT
JP22359	DMS3 infupDT infusion into pDT Reverse (no stop codon)	GAACGATTTAATTAATTAGACGTCTCATCTGGGTGT GTTCAATTGG

qPCR primers:

Description	Forward primer	Reverse primer
McrBC over <i>FWA</i> promoter	TTGGGTTTAGTGTTTACTTG	GAATGTTGAATGGGATAAGGTA
<i>FWA</i> expression	TTAGATCCAAAGGAGTATCAAAG	CTTTGGTACCAGCGGAGA
<i>IPP2</i> expression	GTATGAGTTGCTTCTCCAGCAAAG	GAGGATGGCTGCAACAAGTGT