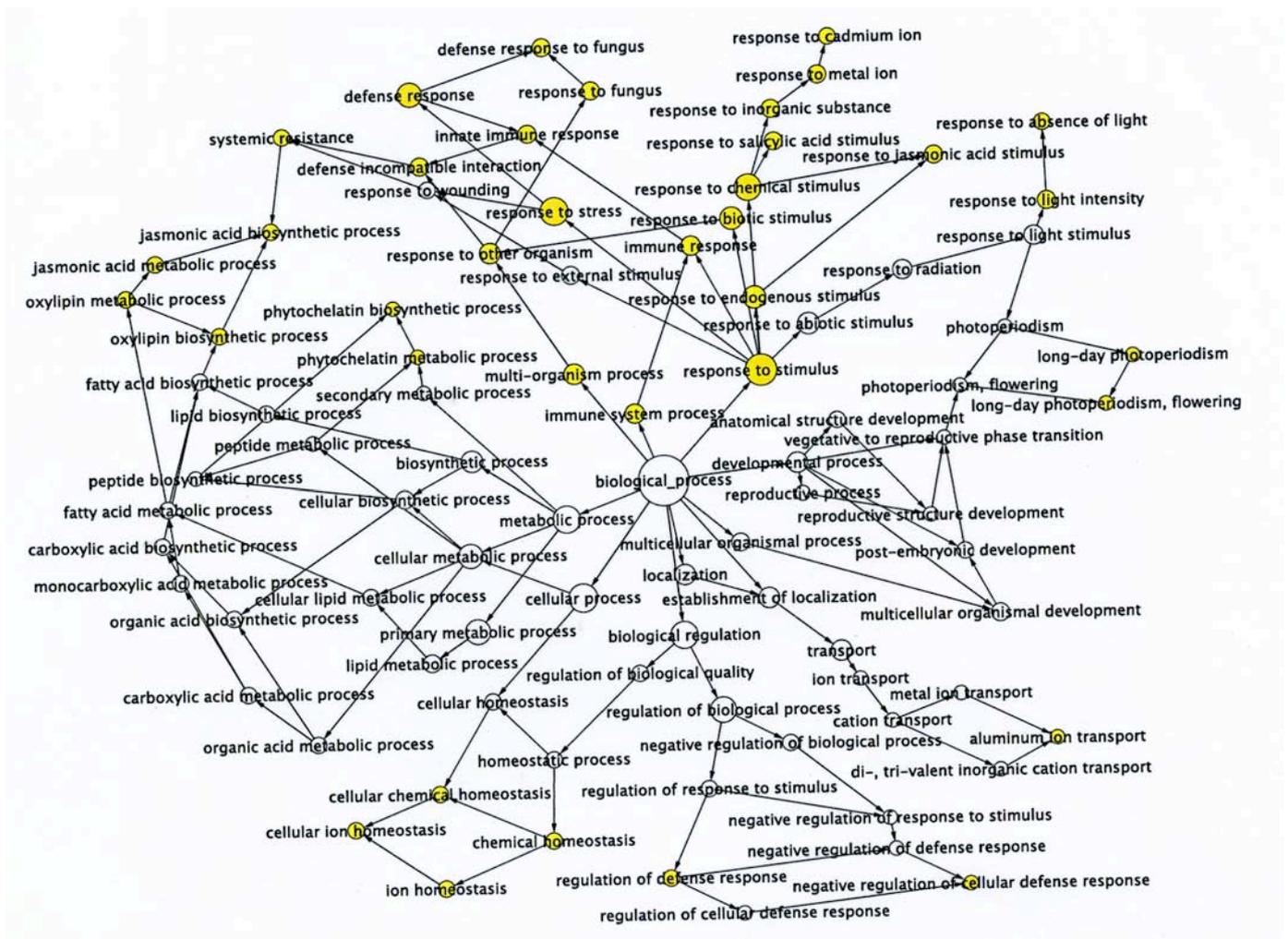
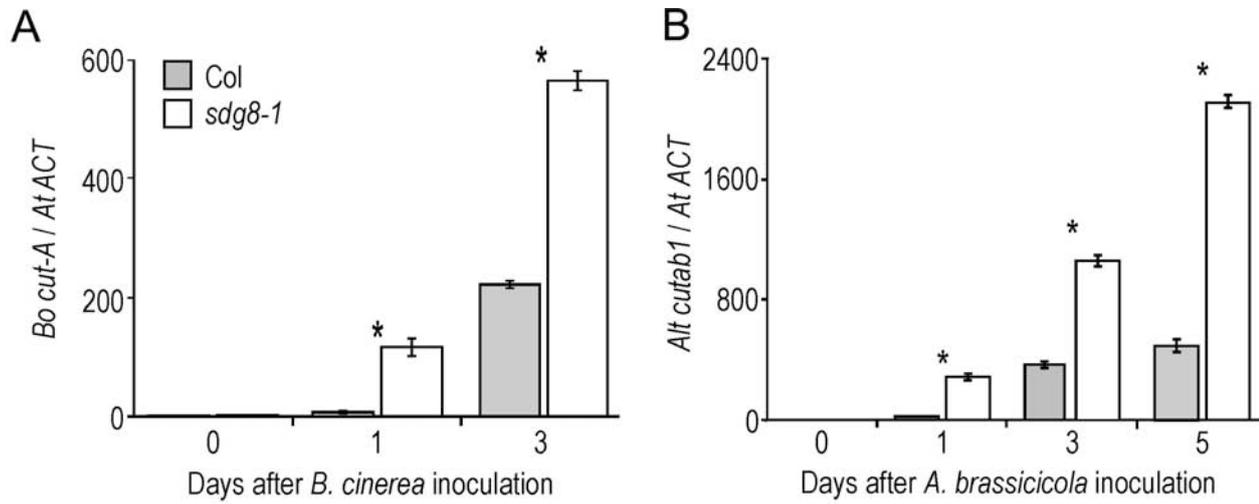


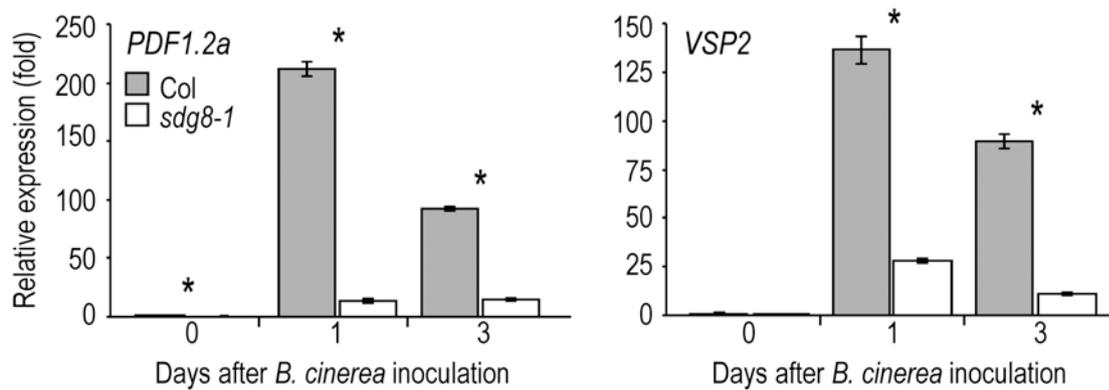
Supplemental Figure S1. Stress-induced expression of *SDG8p::GUS* in independent transgenic lines. Images show leaves wounded by slicing or inoculated with *B. cinerea* at 3 days post-inoculation from transgenic *Arabidopsis* plants containing the *35S::GUS* (control) or the *SDG8p::GUS* construct. Three independent transgenic lines are shown for *SDG8p::GUS*. Blue staining indicates GUS activity.



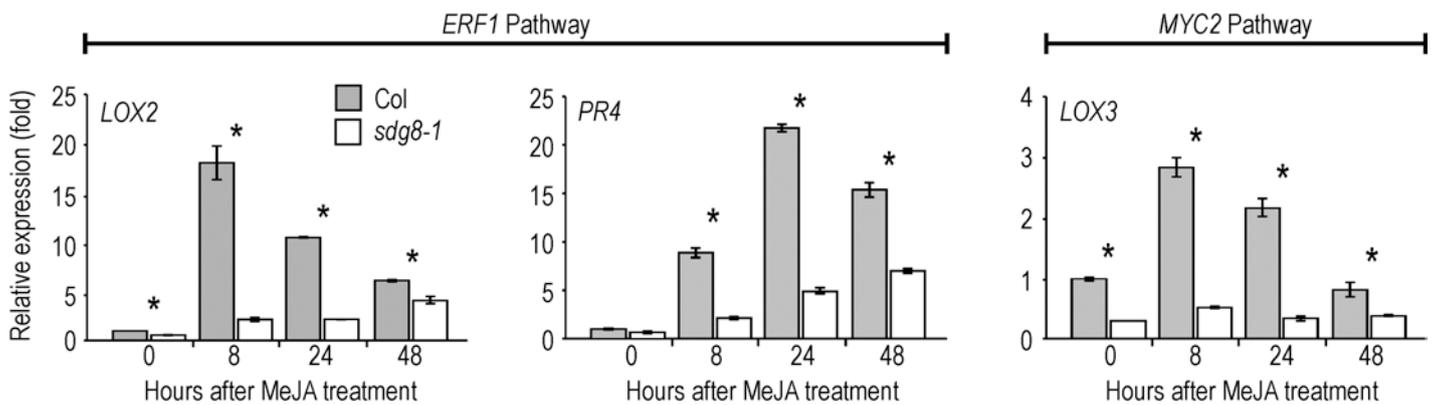
Supplemental Figure S2. Gene ontology analysis of differentially expressed genes in *sdg8-1* seedlings. Genes were identified by microarray analysis previously reported in Xu et al. (2008). Color of the circles corresponds to the level of significance of the overrepresented GO category from $P \leq 0.05$ and below, according to a multiple t -test with false discovery rate-corrected p -value. Yellow tones denote decreasing P values, and the size of the circle is proportional to the number of genes in each category.



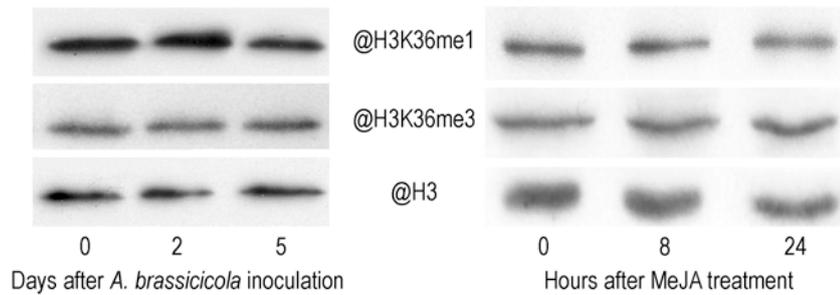
Supplemental Figure S3. Comparison of fungal pathogen growth in infected leaves of mutant *sdg8-1* and wild-type Col plants. Results from an independent experiment are shown, which confirms that *B. cinerea* (A) and *A. brassicicola* (B) multiplication is enhanced in *sdg8-1* compared to Col, as described in Figure 2. Data represent the mean \pm SD of triplicates. Asterisks indicate significance at $P < 0.05$ using *t*-test.



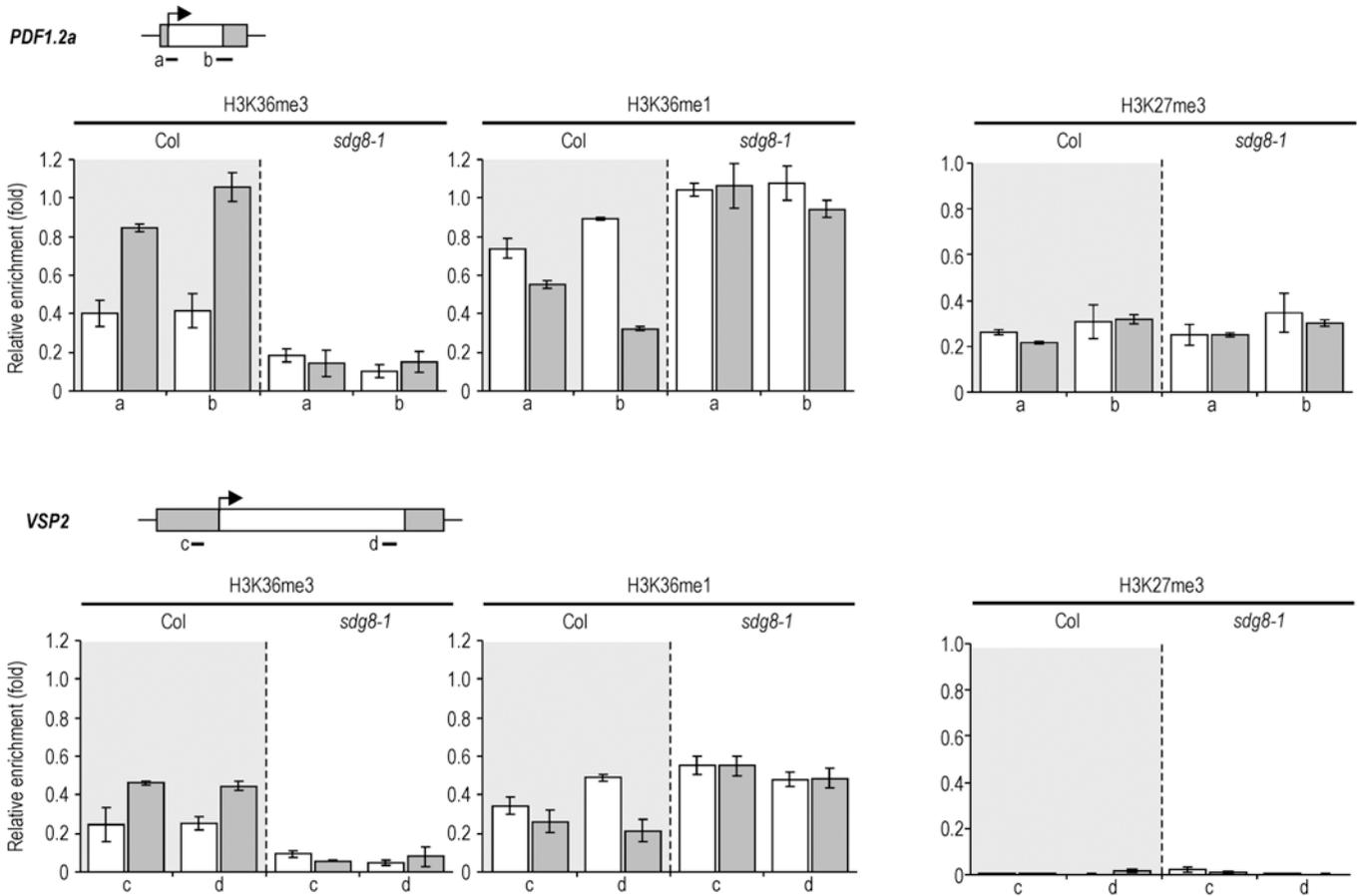
Supplemental Figure S4. Comparison of defense gene expression in response to *B. cinerea* infection in mutant *sdg8-1* and wild-type Col plants. Expression levels of *PDF1.2a* (left) and *VSP2* (right) in *sdg8-1* and Col leaves after *B. cinerea* inoculation are presented relative to average wild-type levels at time point 0 (set as 1). Data represents the mean \pm SD of triplicates. Similar results were obtained in two independent experiments (refer to Supplemental Table S3). Differences in fold induction of defense marker genes in *sdg8-1* and Col are listed in Table S3. Asterisks indicate a significant difference between Col and *sdg8-1* at $P < 0.05$ (two sided *t*-test).



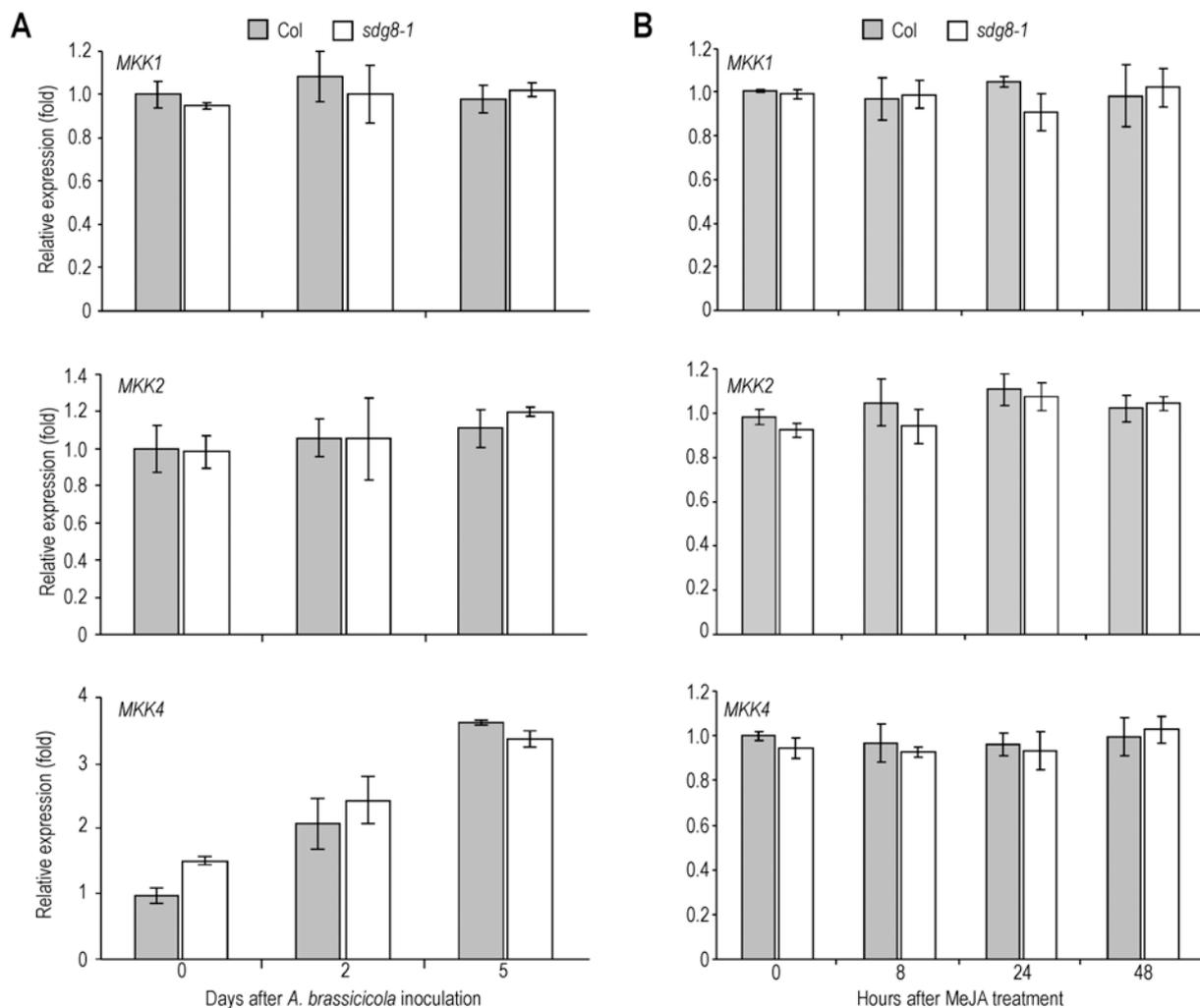
Supplemental Figure S5. Comparison of JA/ET pathway gene expression in response to MeJA treatment in mutant *sdg8-1* and wild-type *Col* plants. Expression levels of JA/ET-responsive genes (*LOX2*, *PR4* and *LOX3*) in *sdg8-1* and *Col* leaves after treatment with exogenously applied MeJA are presented relative to average wild-type levels at time point 0 (set as 1). Data represents the mean \pm SD of triplicates. Similar results were obtained in two independent experiments (refer to Supplemental Table S4). Differences in fold induction of defense marker genes in *sdg8-1* and *Col* are listed in Table S4. Asterisks indicate a significant difference between *Col* and *sdg8-1* at $P < 0.05$ (two sided t -test).



Supplemental Figure S6. Western blot analysis of global H3K36 methylation in response to *A. brassicicola* inoculation or MeJA treatment. Histone-enriched protein extracts of wild-type Col leaves inoculated with *A. brassicicola* (left) or treated with exogenous MeJA (right) were immunoblotted using antibodies that specifically recognize the indicated histone forms. Histone H3 total protein was used as a loading control.



Supplemental Figure S7. Histone methylation at ET/JA pathway defense genes in response to MeJA treatment in mutant *sdg8-1* and wild-type Col plants. Chromatin immunoprecipitation analysis was used to determine the relative levels of H3K36me3, H3K36me1 and H3K27me3 before (white bar) and 8 h after (black bar) exogenous MeJA treatment of 6-week-old Col and *sdg8-1* plants at indicated regions of *PDF1.2a* and *VSP2*. Data represents the mean \pm SD of triplicates. Similar results were obtained in two independent experiments (refer to Supplemental Table S7). Amplified regions (named a to d) are indicated below each gene, which is represented by a white box for the coding region and grey boxes for the 5' and 3' untranslated regions.



Supplemental Figure S8. Expression of *MKK* genes in response to *A. brassicicola* inoculation and MeJA treatment in mutant *sdg8-1* and wild-type Col plants. Expression levels of *MKK1*, *MKK2* and *MKK4* after *A. brassicicola* inoculation and exogenous MeJA treatment are presented relative to average wild-type levels at time point 0 (set as 1). Data represents the mean \pm SD of triplicates. Similar results were obtained in two independent experiments (refer to Supplemental Tables S2 and S4). Note that no significant difference was observed between Col and *sdg8-1*.