Supplementary information inventory

- Figure S1: Comparison of gene expression in *kdm5* mutant larvae and adults, related to Figure 1. Differences in KDM5 targets are shown, in addition to quality control data for RNA-seq analyses.
- 2. Figure S2: Analyses of gene expression levels and KDM5 binding in normal and oxidative stress conditions, related to Figure 1. Scatter plots related to RNA-seq data of paraquat treated flies are shown, in addition to quality control data regarding these transcriptome data.
- 3. Figure S3: Quality control analyses for ChIP-seq data, related to Figure 1. Data showing quantitation of KDM5 ChIP-seq binding data with respect to genomic features are shown. In addition, data comparing ChIP-seq data from normal and paraquat treated animals is shown.
- Figure S4: KDM5 binding is strongly correlated with H3K9 acetylation (H3K9ac) and weakly with H3K27 acetylation (H3K27ac) and H3K4 monomethylation (H3K4me1). Related to Figure 1. Figure contains heat maps showing distribution of KDM5, H3K9ac, H3K4me1 and H3K27ac relative to TSS.
- 5. FigureS5: Preventing KDM5-mediated binding to H3K4me0 does not affect promoter recruitment, related to Figure 6. Figure shows the generation and characterization of a fly strain harboring a mutation in PHD1 that abrogates its ability to bind to H3K4me0.
- 6. Figure S6: KDM5 binding regions of mitochondrial function genes show motif enrichment, related to Figure 6. Figure shows MEME-ChIP analyses of KDM5-regulated genes.
- 7. Supplementary Figure legends: Describes supplementary figures.
- 8. **Table S1: Summary of RNA-seq mapping data, related to Figure 1.** Quantitation of the number of RNA-seq reads that were generated and mapped is shown.
- 9. **Table S2: Gene expression levels of TCA cycle enzymes, related to Figure 3.** Table shows expression of genes involved in the TCA cycle in addition to whether they were identified as direct targets in ChIP-seq analyses.
- 10. **Table S3: Metabolic analyses in kdm5 mutant flies, related to Figure 3.** Table has extensive analyses of the metabolites (amino acids, fatty acids, antioxidants) in *kdm5* mutant flies.
- **11. Table S4: Primers used for RT-PCR and ChIP analyses, related to Figures 2, 4, 5, 6.** Lists all primer sets used for real-time PCR and ChIP-PCR analyses.
- **12. Supplementary methods:** This section contains extensive additional information, particularly regarding the analyses of RNA-seq and ChIP-seq experiments carried out.
- **13. Supplementary references:** Bibliography of references cited in supplementary section.