

Promoter				
Histone Modification	Rank Percentile	Trend	P-value	Ranked Gene Lis
H3K4 Di-Methylation	36.9 %	Depleted	< 10-11	
H3K36 Di-Methylation	39.2 %	Depleted	< 10-7	
H2AK7 Acetylation	26.7 %	Depleted	< 10-12	
H2AZK14 Acetylation	34.6 %	Depleted	< 10-10	
H2BK11 Acetylation	27.6 %	Depleted	< 10-11	
H2BK16 Acetylation	26.5 %	Depleted	< 10-12	
H3Nterm Acetylation	31.0 %	Depleted	< 10-23	0 = 1
H3K9 Acetylation	29.7 %	Depleted	< 10-9	
H3K14 Acetylation	22.7 %	Depleted	< 10 ⁻¹⁶	
H3K18 Acetylation	23.4 %	Depleted	< 10 ⁻¹⁵	
H3K23 Acetylation	26.4 %	Depleted	< 10 ⁻¹²	=1
H3K27 Acetylation	30.3 %	Depleted	< 10 ⁻⁸	
H3K56 Acetylation	48.6 %	none	0.5017	
H4Nterm Acetylation	29.7 %	Depleted	< 10 ⁻²⁶	
H4K8 Acetylation	33.9 %	Depleted	< 10 ⁻⁶	1
H4K12 Acetylation	29.5 %	Depleted	< 10-9	
H4K16 Acetylation	41.4 %	none	0.0080	
H2A Occupancy	55.7 %	none	0.0065	
H2B Occupancy	61.8 %	Enriched	< 10-9	
H3 Occupancy	62.2 %	Enriched	< 10 ⁻⁹	
H4 Occupancy	61.1 %	Enriched	< 10 ⁻⁷	
H2AZ Occupancy	44.0 %	none	0.0031	8 -

Figure S4. Genes repressed by the H2A R78 sprocket arginine residue have high levels of promoter histone occupancy and low levels of active histone post-translational modifications. (A) Histone occupancy and histone modification levels in the promoter regions of 290 genes repressed by H2A R78 (i.e., genes with induced expression in the H2A R78A mutant). Analysis was performed on published ChIP-chip data sets using the web-based database ChromatinDB (O'CONNOR AND WYRICK 2007). Note, histone modification data were normalized by nucleosome occupancy levels. (B) Statistical analysis of the data shown in part A showing enrichment or depletion of histone occupancy or post-translational modifications.

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