SUPPLEMENTAL MATERIAL

TABLES

Table 1. Primers sequences used in ChIP assays performed on the EP300 gene.

		relation to +1
EP300 Promoter	GAGGAGGTGAGTGTCTCTTG	-68
EP300 Promoter	CGCCATCAACCTCTTCCTC	32
EP300 Exon 3	AGATGGGAATGATGAACAACC	33260
EP300 Exon 3	ACTCACCATGTTGGGCATTC	33444
EP300 Exon 6	GAGCACCCGTTGGACTTG	38798
EP300 Exon 6	GGTTCTTTGCTTGCACCTG	38974
EP300 Exon 31	ACCAGGCACTGTGTCTCAAC	85235
EP300 Exon 31	AGGGATGGGTTGTGGATTAG	85412
EP300 3' UTR	TTACCACCAGCCTTTCTTCC	86570
EP300 3' UTR	ATGTCAACCATCTGCACCAG	86753
EP300 3' FL1	GCAGGTACCAGCTAACAGTC	87659
EP300 3' FL1	GGACTTAGGCAGATATGATCC	87812
EP300 3' FL2	TATGCAAGGAATGCAAGAGGG	88994
EP300 3' FL1	AGCTGCTCCCTAGCAGAATG	89160
	EP300 Promoter EP300 Exon 3 EP300 Exon 3 EP300 Exon 6 EP300 Exon 6 EP300 Exon 6 EP300 Exon 31 EP300 Exon 31 EP300 3' UTR EP300 3' FL1 EP300 3' FL1 EP300 3' FL1 EP300 3' FL2	EP300 PromoterCGCCATCAACCTCTTCCTCEP300 Exon 3AGATGGGAATGATGAACAACCEP300 Exon 3ACTCACCATGTTGGGCATTCEP300 Exon 6GAGCACCCGTTGGACTTGEP300 Exon 6GGTTCTTTGCTTGCACCTGEP300 Exon 31ACCAGGCACTGTGTCTCAACEP300 Exon 31AGGGATGGGTTGTGGATTAGEP300 3' UTRTTACCACCAGCCTTTCTTCCEP300 3' UTRATGTCAACCATCTGCACCAGEP300 3' FL1GGACTTAGGCAGATATGATCCEP300 3' FL2TATGCAAGGAATGCAAGAGGG

FIGURE LEGENDS

Supplementary figure 1. Cofilin 1 preferentially occupies gene coding region. ChIP assays. Lysates from cross-linked cells were incubated with the indicated antibodies and coprecipitated DNA was analyzed by PCR with primers amplifying promoter and coding regions of the S19, β -tubulin and GAPDH genes. Non-specific rabbit IgGs were used as negative control. All experiments were reproduced at least in triplicates.

Supplementary figure 2. Effect of latrunculin A, cytochalasin D and jasplakinolide on the occupancy of EP300 gene promoter, proximal, distal exons and 3' flanking region by phosphorylated pol II, actin, H3, H3K9ac and cofilin 1. **(A)** ChIP analysis on lysates from crosslinked cells treated with latrunculin A, cytochalasin D or jasplakinolide, incubated with the indicated antibodies. Co-precipitated DNA was analyzed by PCR using primers amplifying promoter, proximal exon 6, distal exon 31 and 3' gene flanking region FL2. Densitometric quantification of relative occupancies determined by ChIP experiments on chromatin isolated from cells treated with **(B)** latrunculin A, **(C)** cytochalasin D or **(D)** jasplakinolide. In all cases the bars diagrams represent average values calculated from analysis of promoter, coding regions (exon 6 and exon 31) and 3' FL2 over five independent experiments. All values are normalized against the signals obtained with an anti-histone H3 antibody.



