

Figure W1. Coomassie-stained membranes used to ensure equal quantity of protein sample loading for Western blot analysis shown in Figure 2*A*.

Table W1. Primer Sequences Used in the Study.

	Sequence			
ChIP-PCR				
ADAM19 promoter	F: TCACCTCTGAGTGTCACCCAAG	133	60	
1	R: GATTTGTGGTGGACGCGG			
ADAM19 exon	F: TTTGTTCCCACGTTCTGCG	150	60	
	R: GAACGGCGGAAAGAGAAGC			
ADAM19 intron	F: GCTCCATGGGAGCAGTATTCAT	141	60	
	R: AGGCCAGGCTAGAGATATGCTG			
RT-PCR				
SMAD4	F: GTCTTTGATTTGCGTCAGTGTCAT	151	60	
	R: CAGCTGACAGACTGATAGCTGGAG			
TGFβR1	F: AGTTAAGGCCAAATATCCCAAACAG	101	59	
	R: CCTAGCTGCTCCATTGGCAT			
TGFβR2	F: ATTCCCAGCTTCTGGCTCAAC	123	60	
	R: CTCACTGAAGCGTTCTGCCAC			
ADAM19	F: CAAGCTGAGGCAACAGTTCAGT	132	60	
	R: CGCAGGATTTCCGGAGTGT			
GAPDH	F: CCCCTTCATTGACCTCAACTACAT	135	60	
	R: CGCTCCTGGAAGATGGTGA			
Bisulfite sequencing				
ADAM19 promoter	F: AGAGGGGTTTAGTTTAATTTTTAGAAG	340	60	
	R: CTCAACCATACCTACCCACTACCC			
ADAM19 exon	F: GGGTAGTGGGTAGGTATGGTTGAG	347	60	
	R: CRA CCTCTACCACCTCCCAAAAC			
MSP				
ADAM19	MF: AAGGCGTTTGGTATAGTGC	101	60	
	MR: AACGCGACCTATAAAAAATCG			
	UF: TTTAAGGTGTTTGGTATAGTGT	101	60	
	UR: AACACAACCTATAAAAAATCAAAA			
SMAD4 mutation				
Exon 9	F: TATTAAGCATGCTATACAATCTG	330	58	
	R: CTTCCACCCAGATTTCAATTC			
	seq: TTTTGACAACAAATAGAGCTTTAAGTC			
Exon 11	F: CCAAAAGTGTGCAGCTTGTTG	554	58	
	R: CAGTTTCTGTCTGCTAGGAG			
	seq: TATTTTGTAGTCCACCATC			

Table W2. Mutations in SMAD4 in Ovarian Cancer Cell Lines.

	IOSE	SKOV3	MCP3	MCP2	A2780	CP70
Exon 9	Ν	Ν	Ν	Ν	Codon 404 (TA insertion)	Ν
Exon 11	Ν	Intron/T to C	Ν	Ν	Ν	Ν



Figure W2. Chromatin immunoprecipitation–PCR analysis of *ADAM19* in ovarian cancer cells. Chromatin immunoprecipitation assays were performed with antibodies directed against DNA methyltransferase 1 (DNMT1; primers used to amplify two regions of *ADAM19* are shown in Figure 24). The relative binding of each antibody to the corresponding region was measured by quantitative PCR. Error bar indicates SD calculated from triplicates. *P < .05.