Figure S1. Transfection efficiency of p53-siRNAs. Data are presented as the mean \pm SD (n=3). *P<0.05 and **P<0.01 vs. control. siRNA, small interfering RNA.



Figure S2. Effect of dioscin treatment (2.9, 5.8 and 11.6 μ M) for 24 h on colony formation, migratory and invasive properties in A431 cells. Data are presented as the mean ± SD (n=5). **P<0.01 vs. dioscin group. siRNA, small interfering RNA.



Figure S3. Effects of dioscin and caspase inhibitor (Z-VAD-FMK) on the expression levels of cleaved-caspase 3/9. **P<0.01 vs. dioscin group.



Table SI. Primary antibodies used in the present study.

Antibody	Source	Dilution ^a	Company	Cat. no.
Rho	Rabbit	1:1,000	ProteinTech Group, Inc.	10749-1-AP
Cdc42	Rabbit	1:1,000	ProteinTech Group, Inc.	10155-1-AP
GAPDH	Rabbit	1:1,000	ProteinTech Group, Inc.	10494-1-AP
p53	Rabbit	1:1,000	ProteinTech Group, Inc.	10442-1-AP
Bax	Rabbit	1:1,000	ProteinTech Group, Inc.	50599-2-Ig
Bcl-2	Rabbit	1:1,000	ProteinTech Group, Inc.	60178-1-Ig
Cleaved/pro caspase-3	Rabbit	1:1,000	ProteinTech Group, Inc.	19677-1-AP
Cleaved/pro caspase-9	Rabbit	1:1,000	ProteinTech Group, Inc.	10380-1-AP
Cleaved/pro PARP	Rabbit	1:1,000	ProteinTech Group, Inc.	13371-1-AP
MMP-2	Rabbit	1:1,000	ProteinTech Group, Inc.	10373-2-AP
MMP-9	Rabbit	1:1,000	ProteinTech Group, Inc.	10375-2-AP
ATM	Rabbit	1:500	Wuhan Boster Biological Technology, Ltd.	bs-1370R
p-ATM	Rabbit	1:500	Wuhan Boster Biological Technology, Ltd.	bs-12545R

^aDilutions used for western blotting. MMP, matrix metalloproteinase; p-ATM, phosphorylated ataxia telangiectasia-mutated; PARP, poly (ADP-ribose) polymerase.