Supplementary Table 1 Expression levels of the selected genes in vehicle-treated versus cisplatin-treated cells.

GenBank No.	Gene name	Fold change	P-value
NM_003383	VLDLR	7.230	0.0095
NM_032387	WNK4	6.237	0.0042
<u>NM_000201</u>	<u>CD54</u>	<u>5.687</u>	<u>0.0013</u>
NM_002571	PAEP	4.371	0.0243
NM_002548	OR1D2	4.150	0.0492
NM_172160	KCNAB1	3.837	0.0198
NM_032375	AKT1S1	3.632	0.0365
NM_152291	MUC7	3.564	0.0318
NM_012128	CLCA4	3.416	0.0423
NM_014941	MORC2	3.307	0.0435
NM_014376	CYFIP2	3.282	0.0127
NM_147148	GSTM4	3.254	0.0235
NM_020889	PHF12	3.176	0.0472
NM_013278	IL17C	3.129	0.0497
NM_005082	TRIM25	3.019	0.0056
NM_024889	C10orf81	3.002	0.0143

Supplementary Table S2 The tumor initiation capabilities in CD54⁺ versus CD54⁻ PCa cells.

Gro	oup	No. cells / injection	No. tumors / No. injections
PC3		10,000	7/10
	CD54	100,000	9/10
	CD54⁻	10,000	0/10
		100,000	2/10
		10,000	6/10
	CD54	100,000	8/10
LINCAP	CD54⁻	10,000	0/10
		100,000	1/10
DU145 ·	CD54 ⁺	10,000	3/10
		100,000	5/10
	CD54⁻	10,000	0/10
		100,000	0/10
22RV1	CD54 ⁺	10,000	3/10
		100,000	4/10
	CD54⁻	10,000	0/10
		100,000	0/10

Supplementary Table 3

Relationship between the expression levels of CD54 and clinicopathological features in prostate cancers

			Aberrant expression of		
				CD54	
Total		Patients	High	Low	Р
Age (years)	Mean	62.5	63.7	61.2	0.375
Stage (cases)	T1	32	13	19	< 0.001
	T2	70	32	38	
	Т3	89	58	31	
	T4	48	39	9	
Grade (cases)	G1 or 2	142	50	92	< 0.001
	G3	97	83	14	
Lymphatic invasion (cases)	Negative	79	36	43	<0.005
	Positive	57	46	11	
	Unknown	103	58	45	
Chemo (cases)	Primary	24	6	18	< 0.001
	Recurrent	12	9	3	
Follow-up (months)	Mean	73.2	72.5	73.9	0.576

Supplementary Table 4

The tumor initiation number in primary (1°) and recurrent (2°) $CD54^+$ versus $CD54^-$ PCa xenograft mice.

Group		No. Cells/Injections	No. Tumors/Injections
1°	CD54+	10,000	3/10
		100,000	5/10
	CD54-	10,000	0/10
		100,000	0/10
2°	CD54+	10,000	4/10
		100,000	7/10
	CD54-	10,000	0/10
		100,000	0/10

Supplementary Figure Legends

Figure S1. Tumor formation (photon counts) was monitored in the xenografts of cells from DDP-treated and vehicle-treated prostate cancer.

Figure S2. The expression of CD54 in PC3 cells as determined by FACS. **(A)** Knockdown or/and exogenous expression of CD54 in cancer cells. **(B)** Knockdown and exogenous expression of FLAG-labeled CD54 in cancer cells.

Figure S3. The mRNA expression of basal or progenitor markers in CD54⁺ versus CD54⁻ PC3 cells.

Figure S4. The tumor weight in primary (1°) and recurrent (2°) CD54⁺ versus CD54⁻ cancer cells xenografted into mice.

Figure S5. The representative and quantitative data of sphere formation capabilities in primary (1°) and recurrent (2°) PC3 cells. oe, overexpressed CD54; CD54-DAPT, cells treated with the CD54 inhibitor DAPT.

Figure S6. HE and IHC staining in the representative animals from the patient-derived xenograft mouse model.

Figure S7. Flow cytometry analysis of CD54. (A, B) Upon culture for 15 days, FACS-sorted CD54+ PC3 cells gradually reconstituted the original proportion of CD54-expressing cells.













Supplementary Figure 7

