



Supplementary

RGL2 drives the metastatic progression of colorectal cancer via preventing the protein degradation of β -catenin and KRAS

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Table S1. Cox univariate and multivariate analyses under the condition of overall survival probability in association with RGL2 mRNA expression levels and pathological stage derived TCGA cohort with colon cancer.

Overall survival (n = 184)				
Variables	Crude HR (95%CI)	P	Adjusted HR (95%CI)	р
Age				
< 64	1	NA	1	NA
≥ 64	1.24 (0.65-2.36)	0.522	2.10 (1.02-4.34)	0.044
Gender				
female	1	NA	1	NA
male	1.03 (0.54-1.03)	0.925	0.71 (0.35-1.46)	0.353
pT				
T1-T2	1	NA	1	NA
T3-T4	4.27 (1.03-17.8)	0.046	1.90 (0.42-8.67)	0.409
pΝ				
N0	1	NA	1	NA
N1-N2	2.49 (1.28-4.82)	0.007	0.49 (0.13-1.85)	0.293
pM				
M0	1	NA	1	NA
M1	4.95 (2.58-9.51)	< 0.001	2.98 (1.30-6.85)	0.010
stage				
I-II	1	NA	1	NA
III-IV	3.34 (1.65-6.74)	0.001	3.31 (0.65-16.9)	0.149
RGL2 expression				
low	1	NA	1	NA
high	4.73 (1.45-15.4)	0.010	3.67 (1.08-12.4)	0.037

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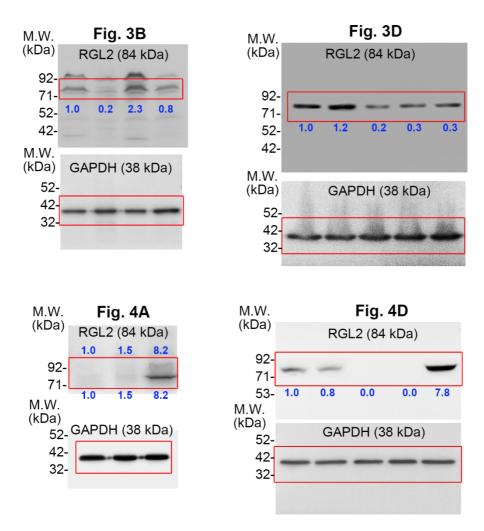


Figure S1. Uncut blots for Figure 3 and 4. The abbreviations M.W. and kDa denote molecular weight and kilodalton, respectively. The inserted values indicate the ratios of protein intensities which were normalized to the protein levels of their respective GAPDH control.

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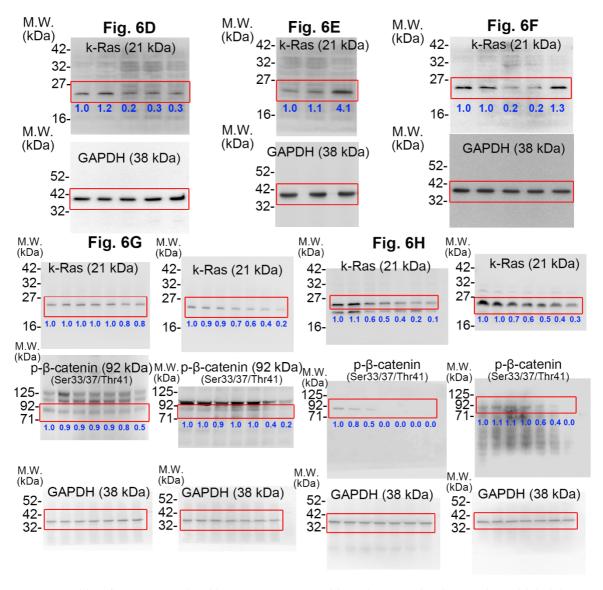


Figure S2. Uncut blots for Figure 6. The abbreviations M.W. and kDa denote molecular weight and kilodalton, respectively. The inserted values indicate the ratios of protein intensities which were normalized to the protein levels of their respective GAPDH control.