RIN1 promotes renal cell carcinoma malignancy by activating EGFR signaling through Rab25

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Renal cell carcinoma (RCC) is one of the most common genitourinary neoplasms and accounts for approximately 3% of all malignancies worldwide with global incidence rates increasing by 2–3% per year.1,2 The clear cell renal cell carcinoma (ccRCC) is the main subtype of RCC and accounts for approximately 75% of all renal tumors.3 Approximately 30–40% of patients possibly experience disease metastasis after primary radical surgery.4–6 The prognosis of metastatic ccRCC patients remains generally dismal and its 5-year survival rate is approximately 10%.7 Therefore, understanding the molecular mechanisms during the initiation and the development of ccRCC could be helpful for identifying therapy strategies in ccRCC.

Ras and Rab interactor 1 (RIN1), a RAS effector protein, was first reported to activate ABL tyrosine kinases,8,9 Overexpression of RIN1 has been reported to be associated with poor prognosis in non-small cell lung cancer,10 bladder urothelial carcinoma,11 gastric adenocarcinoma12 and melanoma.13 Recently, we identified the important role of RIN1 expression in the prognosis of ccRCC.14 RIN1 interacts with Abl kinases to regulate cell adhesion and migration.9 RIN1 can also act as a Rab5-directed guanine nucleotide exchange factor (GEF) protein,15 thereby regulating EGFR signaling.16,17 To date, however, the impact of RIN1 expression on ccRCC cells and its potential oncogenic role and molecular mechanisms in ccRCC have not been elucidated.

Rab25 (also termed Rab11c) are key regulators in endosomal trafficking and recycling back to the plasma membrane of EGFR.18,19 Rab25 have been shown to play important roles in receptor recycling (including EGFR recycling) in kidney cells.20,21 By analyzing the transcriptional levels of 52 Rab GTPases in RCC tissues, Rab25 was the most significantly upregulated one.22 We have previously demonstrated Rab25 plays an oncogene role in urothelial carcinoma.23 The relationship between RIN1 and Rab25, as well as their role in EGFR signaling in ccRCC are not clear.

In the current study, we found that ectopic overexpression of RIN1 in ccRCC cells promotes tumorigenesis in ccRCC both in vitro and in vivo. RIN1 plays an important role in the activation of EGFR signaling in ccRCC through interacting with Rab25. Our results provide functional and mechanistic links...
between the putative oncogene RIN1 in the aggressive nature of ccRCC.

Materials and Methods

Cell lines and clinical samples. 769-P, 786-0, A498, ACHN, HK-2, Caki-1, Caki-2 and NC65 were bought from the American Type Culture Collection and were maintained in appropriate medium. OS-RC-2 cell was bought from the National Platform of Experimental Cell Resources for Sci-Tech (Wuhan, China) and was maintained in T-medium supplemented with 5% FBS. A panel of 30 fresh ccRCC tissues and matched adjacent non-tumor normal tissues were collected from 30 patients with ccRCC treated at the First Affiliated Hospital of SYSU between January 2014 and August 2014 and stored in liquid nitrogen until further use. The patient study was conducted in accordance with the Declaration of Helsinki, and the study protocol was approved by the Ethics Committee of the First Affiliated Hospital of Sun Yat-sen University. Informed consent was obtained from all patients.

RNA isolation and quantitative RT-PCR. Extraction of total RNA and measurement of mRNA quantity were performed as described previously. RNA was extracted from cells using TRIzol (Invitrogen, Carlsbad, CA, USA) following protocols supplemented with the manufacturer’s instructions. First-strand cDNA was then reverse transcribed by MMLV transcriptase (Promega, Madison, WI, USA) using random primers. Real-time RT-PCR was performed on a CFX96 real-time PCR detection system (Bio-Rad), and a Roche SYBR FAST Universal qPCR Kit (Roche Molecular Biochemicals, Indianapolis, IN, USA) was used for gene detection. Primers for RIN1 and GAPDH are listed as follows: RIN1-specific primer: forward 5'-GCACCTGGCGAGAAAG-3' and reverse 5'-TAGATTTCCGCACGAGGAACG-3'.

Vector construction and lentivirus packaging and transduction. Flag-tagged Rab25 (LPP-T4697-Lv121-200), Flag-tagged RIN1 (LPP-T2755-Lv121-200) and their corresponding control vectors were purchased from GeneCopoeia (Rockville, MD, USA). Lentiviral shRNA targeting RIN1 (LV3-pGLV-h1-GFP-puro) and control empty vector were synthesized at GenePharma (Shanghai, China). The target sequence of RIN1 used to construct a lentiviral shRNA was 5'-AGCTTCCTCCTGGCG-GAAATCT-3'. Vectors were packaged in 293T cells using ViraPower Mix (GeneCopoeia). After culturing for 48 h, lentiviral particles in the supernatant were harvested and filtered by centrifugation at 5000 g for 10 min, and then transfected into ccRCC cells. The cells were cultured under puromycin (2 μg/mL) selection for 2 weeks, at which point real-time PCR was used to determine the level of RIN1. The siRab25 (5'-GGAGCUCAUUGACCAUGCU-3') oligonucleotides were synthesized at GenePharma. Transfection of oligonucleotides was performed using Lipofectamine 2000 reagent (Invitrogen) according to the manufacturer’s instructions. EGFR inhibitor AG1478 was purchased from Abcam (Shanghai, China).

Western blot and immunoprecipitation. For western blots, total cellular protein was extracted from cells and separated by sodium dodecyl sulfate–polyacrylamide gel electrophoresis. For immunoprecipitation, cells were transfected with the Flag-tagged Rab25 or with Flag-tagged RIN1 vectors. Cells were solubilized in lysis buffer. The whole-cell lysates obtained by centrifugation were incubated with 2 μg of specified antibody bound to either protein A or Protein G Sepharose beads or with Streptavidin Sepharose beads (Amersham Biosciences, Pittsburg, PA, USA) for 1 h at 4°C. The immunocomplexes were then applied to SDS-PAGE. The following antibodies were used: anti-RIN1 (Abcam, Cambridge, MA, USA), phospho-EGFR (Tyr1173), anti-EGFR (both from Santa Cruz Biotechnology, Santa Cruz, CA, USA), anti-Phospho-AKT (Thr308), anti-AKT, anti-phospho-ERK (Thr202/Tyr204), anti-ERK and anti-Rab25 (all from Cell Signaling Technology, Beverly, MA, USA); and anti-β-actin and anti-Flag (both from Sigma, St. Louis, MO, USA). HRP-conjugated anti-mouse and anti-rabbit secondary antibodies were obtained from Promega.

MTT assay. Cell viability was measured using a 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) assay (Sigma). Briefly, cells were seeded in 96-well plates and cultured. Cell viability was examined by following standard procedures. Experiments were performed in triplicate.

Wound healing and invasion assays. Cell migration was assessed by measuring the movement of cells into a scraped, a cellular area made by a 200-μL pipette tube. Wound closure was observed after 24 h. Invasion assays were performed with 24-well BioCoat Matrigel Invasion Chambers (BD) according to the manufacturer’s instructions. Briefly, 2 × 10⁵ cells were seeded into 8-μm porous inserts in triplicate wells and incubated for 24 h. The invaded cells in lower filters were fixed in methanol and stained in crystal violet (Sigma) before being counted under a microscope.

In vivo experiments. All experiments were performed in accordance with China Public Health Service Guide for the Care and Use of Laboratory Animals. Experiments involving mice and protocols were approved by the Institutional Animal Care and Use Committee of Sun Yat-sen University. Female BALB/c nu/nu athymic mice (4–5 weeks old), purchased from Shanghai SLAC Laboratory Animal (Shanghai, China), were kept under specific-pathogen-free conditions. For the xenograft tumor growth assay, 786-O-shRIN1 or 786-O-Con cells were injected subcutaneously into the right flank of the mice (5 mice per group), and this was performed in triplicate. Two weeks after inoculation, tumor size was measured every 3–4 days until the tumors grew to a diameter of 20 mm or when the tumor burden exceeded 10% of the body weight, at which time the mice were killed by cervical dislocation. Tumor volume was calculated by the formula V = ab²/2, where a = longest axis and b = shortest axis. In the tumor metastasis analysis, 10 four-week-old BALB/c nude mice in each experimental group were injected with 786-O-shRIN1 or 786-O-Con cells, respectively. Briefly, 2 × 10⁵ cells were injected intravenously through the tail vein into each mouse in a laminar flow cabinet. Six weeks after injection, the mice were sacrificed and examined.

Immunohistochemical staining. In brief, paraffin-embedded sections were deparaffinized and incubated in retrieval buffer solution for antigen retrieval. Protein expression was visualized using a DAKO Real Envision Kit (K5007; Dako, Glostrup, Denmark) after staining with the primary antibody. Staining intensity was scored manually by two independent experienced pathologists as: 0 = no staining, 1 = weak staining, 2 = moderate staining and 3 = strong staining. Tumor cells in five fields were selected randomly and scored based on the percentage of positively stained cells (0–100%). The final immunohistochemistry (IHC) score was calculated by multiplying the intensity score by the percentage of positive cells.

TCGA data. For the TCGA set, clinical data and mRNA expression (level 3 data, RNA-seq Version 2 Illumina) were downloaded from the TCGA data portal (https://cancergenome.nih.gov/) on 1 June 2016. Global mRNA expression profiles of a subset of TCGA ccRCC specimens for which RIN1 expression data were available were subject to GSEA to identify the association of RIN1 with EGFR signaling pathways. For
Fig. 1. RIN1 is overexpressed in clear cell renal cell carcinoma (ccRCC) cell lines and tissues. (a) Western blotting analysis of RIN1 protein expression in pairs of matched ccRCC (T) and adjacent normal tissues (N). (b) RT-qPCR analysis of RIN1 mRNA expression in 30 pairs of ccRCC tumor tissues and adjacent normal tissues. Patients with blue lines indicate increased RIN1 expression in the tumors, while orange lines indicate decreased RIN1 expression in the tumors. (c) Relative expression of RIN1 in renal carcinoma cell lines. (d) Western blotting analysis showing RIN1 knockdown in 786-O and Caki-1 cell lines and RIN1 overexpression in NC65 cell line.

Fig. 2. Loss of RIN1 attenuates clear cell renal cell carcinoma (ccRCC) cell growth, migration and invasion in vitro. (a) Suppression of RIN1 reduced the proliferative ability of 786-O and Caki-1 ccRCC cells, as determined by MTT assay. (b) RIN1 knockdown markedly reduced the foci-formation ability of the cells as determined by the foci-formation assay. (c) Suppression of RIN1 reduced the migration ability of the cells as determined by a wound healing assay. (d) Suppression of RIN1 reduced the cells’ invasion ability as determined by Boyden chamber invasion assay. Each bar represents the mean ± SEM derived from three independent experiments. A two-tailed Student’s t-test was used for statistical analysis (*P < 0.05).
GSEA, RIN1 expression was treated as a numeric variable. GSEA was performed using GSEA 2.0.9 software (http://www.broadinstitute.org/gsea/). We analyzed the reverse-phase protein array (RPPA) data of the TCGA cohort using the cBioPortal web application from the Memorial Sloan Kettering Cancer Center.\(^{(25)}\)

**Statistical analysis.** The Mann–Whitney U-test was used for comparing expression levels in malignant and non-malignant samples, and the Wilcoxon test was used for pairwise comparison. Comparisons between groups were performed using Student’s t-test. All error bars represent the mean ± SEM derived from three independent experiments. For survival analysis, the Kaplan–Meier method was used, and the log-rank test was used for comparing cumulative survival. Statistical analyses were carried out using IBM SPSS Statistics 20.0 (IBM, Armonk, NY, USA). Statistical significance was set at 0.05.

**Results**

**RIN1 is overexpressed in clear cell renal cell carcinoma cell lines and tissues.** In this study, western blotting was conducted for RIN1 protein expression in six pairs of ccRCC and adjacent normal tissues. Five (83.3%) ccRCC tissues showed upregulated RIN1 expression compared with adjacent normal tissues (Fig. 1a). The western blotting results were confirmed by real-time PCR in 30 pairs of ccRCC and adjacent normal tissues. Twenty-three (76.7%) ccRCC tissues showed upregulated RIN1 mRNA expression compared with adjacent normal tissues (Fig. 1b). The expression levels of RIN1 were then examined by real-time PCR in eight renal cell carcinoma cell lines (Fig. 1c). Subsequently, we infected 786-O and Caki-1 cells with lentivirus carrying shRNA targeting RIN1 (shRIN1) or control shRNA (shCon). We transfected NC65 cells, which showed almost undetectable expression of endogenous RIN1, with an RIN1 expression vector or control empty vector. Western blotting results showed high-efficiency RIN1 gene knockdown or overexpression in indicated ccRCC cells (Fig. 1d).

**Loss of RIN1 attenuates clear cell renal cell carcinoma cell growth, migration and invasion in vitro.** MTT assay showed that knockdown of RIN1 dramatically reduced the proliferation of both 786-O and Caki-1 cells (Fig. 2a). Using foci-formation assays, we found that RIN1 depletion reduced foci-formation ability of 786-O and Caki-1 cells compared with the control cells (Fig. 2b). Knockdown of RIN1 caused an apparent suppression of cell migration in both 786-O and Caki-1 cell lines using a wound-healing assay (Fig. 2c). Matrigel invasion assays also demonstrated that ablation of endogenous RIN1 reduced the invasive capacity of both 786-O and Caki-1 cell lines (Fig. 2d). Taken together, our data suggest that loss of RIN1 attenuates ccRCC cell growth, migration and invasion in vitro.

**Depletion of endogenous RIN1 inhibits tumor growth and metastasis of clear cell renal cell carcinoma cells in vivo.** We further studied the in vivo impact of RIN1 on ccRCC cell growth and metastasis by injecting 786-O cells containing either a control or RIN1-shRNA vector into BALB/c nude mice, either subcutaneously or via the tail vein. RIN1-silenced cells showed weakened tumorigenicity and slower growth, forming smaller tumors than control cells (Fig. 3a–c). In the mouse metastasis model, we injected 786-O-con and 786-O-shRIN1 into the lateral tail vein of athymic nude mice (10 mice per group). As shown in Figure 3d, the mice injected with 786-O-shRIN1 cells formed fewer nodules per lung than the mice injected with control shRNA cells (2.3 ± 1.8 vs 9.5 ± 3.6, \(P < 0.001\), Fig. 3).
Mann–Whitney test). Histological studies confirmed that the lesions were caused by extravasation and subsequent tumor growth of 786-O cells into the lungs (Fig. 3e).

**RIN1 activates EGFR signaling in clear cell renal cell carcinoma.** To investigate the mechanism underlying the robust effect of RIN1 on tumorigenesis in ccRCC cells, we first sought to identify potential RIN1 regulate signaling by employing GSEA analyses. By performing GSEA in the TCGA ccRCC cancer dataset, we found that the RIN1 mRNA level was positively correlated with EGF-activated gene signatures (Fig. 4a). Phosphorylation of EGFR, AKT and ERK, well-known downstream molecules in the EGFR signaling pathway, were also inhibited by RIN1 knockdown in both 786-O and Caki-1 cells, and the opposite results were observed in the RIN1 overexpressing NC65 cells (Fig. 4b). Next, we analyzed the different downstream signaling activity (as measured by phosphorylation) in the reverse-phase protein array (RPPA) data of the same TCGA cohort. In this proteomic analysis, the expression of p-AKT (PT308) and p-MAPK/ERK kinase 1 (PS217_S221) in the RIN1 mRNA altered group (upregulation) are significantly higher than in the RIN1 mRNA unaltered group (Fig. 4c,d). Thus, these results illustrate the potential importance of the EGFR signaling axis in RIN1-induced tumor malignancy in ccRCC.

**Rab25 is responsible for RIN1-induced EGFR signaling and tumor malignancy in clear cell renal cell carcinoma cell.** To investigate whether Rab25 and EGFR signaling are required for RIN1-induced ccRCC cell proliferation and invasiveness, NC65-RIN1 cells were treated with siRNA targeting Rab25 and EGFR inhibitor (AG1478, 10 µM), respectively. After siRab25 and AG1478 treatment, RIN1-induced EGFR signaling was inhibited, as evidenced by the decreased expression of p-EGFR, p-AKT and p-ERK in NC65-RIN1 cells (Fig. 5a). The silencing of Rab25 and inhibition of EGFR strikingly reversed the ability of RIN1-overexpressing 786-O cells for growth, migration and invasion in vitro (Fig. 5b–d). These data, taken together, provide evidence that Rab25 is responsible for the RIN1-induced tumor malignancy and/or the activation of EGFR signaling in ccRCC cells.

**RIN1 interacts with Rab25 in clear cell renal cell carcinoma cells.** We validated the interaction of RIN1-Rab25 in vivo through co-immunoprecipitation experiments. Endogenous RIN1 was found to be associated with exogenously expressed Flag-tagged Rab25 and conversely endogenous Rab25 was found to be immunoprecipitated with RIN1 (Fig. 6a). These results were confirmed by IHC analyses of RIN1 and Rab25 expression in 20 fresh ccRCC specimens, in which a positive correlation between RIN1 levels and the expressions of Rab25 was observed ($r = 0.21$, $P < 0.05$, Fig. 6b). Rab25 is overexpressed in ccRCC tissues, and overexpression of Rab25 was seen in endogenous RIN1 high expression 786-O and Caki-1 cells (Fig. S1). To confirm the important roles of RIN1 and Rab25 in ccRCC, we analyze the prognostic significance of RIN1 and Rab25 by using the ccRCC dataset from TCGA. Kaplan–Meier survival analysis showed that high mRNA expression level of RIN1 or Rab25 predicts poor overall survival (OS) in 533 ccRCC patients (Fig. 6c,d).

**Discussion**

Previous studies of ccRCC confirmed the important role of the EGFR signaling pathway in renal carcinogenesis. Understanding the regulation of EGFR signaling in ccRCC is of great value for future development of novel therapeutic strategies. We demonstrate that RIN1 overexpression substantially activates EGFR signaling in ccRCC through binding to Rab25. These observations suggest that RIN1 is involved in EGFR signaling transduction pathways and might provide a direct link between RAS and tyrosine kinase-mediated signals.
Overexpression of EGFR in RCC has been shown in various research, ranging from 50% to 90%. (29–33) EGFR intracellular tyrosine kinase domain is autophosphorylated and results in activation of several downstream signaling pathways, including the RAS-RAF-MEK-ERK and the PI3K-PTEN-AKT pathways. Our data show that p-AKT and p-ERK were simultaneously upregulated by RIN1. The RPPA ccRCC data from TCGA also confirmed the higher expression of activated AKT and activated MAPK/ERK kinase 1 in RIN1 mRNA upregulated ccRCC patients than those with unaltered mRNA expression level. Furthermore, antagonizing RIN1 caused multilevel inactivation of EGFR signaling and had obvious inhibitory effects on tumorigenesis. These results suggest that RIN1 may be a therapeutic agent that can target EGFR signaling in ccRCC to suppress tumorigenesis.

RIN1 is a RAS-effector protein that can affect Ras signaling at different levels: (34) first, by competing with RAF1 protein for binding to activated Ras; (35) second, by enhancing signaling from ABL1 and ABL2, which regulate cytoskeletal remodeling; (36,37) and, third, by activating Rab5A, possibly by functioning as a guanine nucleotide exchange factor for Rab5A, by exchanging bound GDP for free GTP, and facilitating Ras-activated receptor endocytosis. (38) In our study, we found that silencing Rab25 inhibited EGFR signaling and strikingly reversed the ability of RIN1-overexpressing ccRCC cells to proliferation, migration and invasion in vitro. Co-immunoprecipitation data show that RIN1 binds to Rab25 in ccRCC cells. These results suggest that Rab25 might be responsible for RIN1-induced EGFR signaling and tumor malignancy in ccRCC cells.

Rab25 (Catx-8, Rab11c) is a member of the Rab family that plays a key role in governing apical and late endosomal recyling routes. (20) Increased Rab25 is associated with poor patient outcome in ccRCC, likely through effects on metastatic pathways. (22,39) Mechanistically Rab25-decorated vesicles transport integrins, (40) PI3K (41) and EGFR, (18,19,42,43) facilitating aggressive tumor growth and metastasis. We speculate that the role of Rab25 in EGFR trafficking and recycling to the plasma membrane might contribute to its role in RIN1-induced EGFR signaling in ccRCC. Clearly, further work is needed to clarify the mechanisms of Rab25 regulating EGFR signaling in detail.

Our current study revealed that overexpression of RIN1 in ccRCC cells activated the EGFR signaling pathway, thus promoting proliferation, migration and invasion of ccRCC. Meanwhile, antagonizing RIN1 caused multilevel inactivation of EGFR signaling and had obvious inhibitory effects on
These results suggest that RIN1 may be a novel prototype therapeutic agent that can target EGFR signaling in ccRCC to suppress tumorigenesis.

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**Disclosure Statement**

The authors have no conflicts of interest to declare.
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Supporting Information

Additional Supporting Information may be found online in the supporting information tab for this article:

Fig. S1. Rab25 is overexpressed in clear cell renal cell carcinoma (ccRCC) cell lines and tissues. (A) Western blotting analysis of Rab25 protein expression in pairs of matched ccRCC (T) and adjacent normal tissues (N). (B) Western blotting analysis showing relative expression of Rab25 in renal cell carcinoma cell lines.