

# Evaluation of the novel gene Rabl3 in the regulation of proliferation and motility in human cancer cells

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**Abstract.** Rab is a large subfamily of small GTPases which play an important role in multiple processes relating to cellular transportation and modulation of the cytoskeleton. Novel gene Rab-like 3 (Rabl3), which we originally reported as a new member belonging to this subfamily, may participate in other processes in human cancer cell lines based on our research. In order to investigate the function of Rabl3 and the basic mechanism which regulates cancer cell survival and motility, we constructed the Rabl3-pcDNA3.1 fusion plasmid, also, the specific siRNA against Rabl3 was used. We detected cellular viability and motility changes in Rabl3 overexpressed or si-Rabl3 transfected cancer cell lines. Overexpression of Rabl3 resulted in the enhancement of cell proliferation, inhibition of apoptosis and paclitaxel resistance in human cancer cell lines. Rabl3 also promoted cell motility activity. Next, we silenced Rabl3 in order to determine its exact physiological function. We found that processes which are associated with tumor formation and metastasis were inhibited. These included an increased incidence of apoptosis, abrogated cellular proliferation and mobility. Furthermore, Western blot analysis revealed that the function of Rabl3 was closely associated with focal adhesion kinase (FAK) phosphorylation, both in HeLa and MDA-MB-231 cell lines at the sites of Tyr 397/576/577. Our results suggested that Rabl3 may be a novel oncogene which regulates the oncological behavior of human cancer cells. As a consequence, Rabl3 can be considered as a novel candidate in the control of tumorigenesis and as a new target for anti-tumor treatment.

## Introduction

The Rab subfamily, which is the largest group in the Ras superfamily, comprises small G-proteins which represent the

guanine nucleotide binding proteins (1). More than 60 Rabs have been discovered to exist ubiquitously in various kinds of eukaryocyte organisms, ranging from yeast to mammals (2). The sequences of the Rab subfamily members are highly conserved, with molecular weights ranging between 20 and 26 KD. Most of the Rab members can bind to GTP or GDP, functioning as low-molecular weight GTPases and contributing to GDP/GTP conversion.

At the physiological level, they are involved in various kinds of cellular processes or signaling transductions which are crucial for cellular membrane function, cytoskeleton regulation and protein recycling. Therefore, Rabs play key roles in secretion, endo-/exo-cytosis and intra-/inter-trafficking events (3,4). Moreover, the present study provides additional evidence in support of the participation of Rabs in signaling transduction, modification and the crosstalk between signal cascades (5,6).

At the pathological level, the dysfunction of G-proteins results in abnormalities in many central events and can even initiate tumorigenesis, tumor cell invasion or metastasis in some cases (7-9). Rab family members, such as Rab25, Rab20 and Rab2, are highly expressed in cancer cells and have close relationships with the progression and aggressiveness of different types of cancer (10,11). The impact of Rabs on tumor cells, in terms of their modulation of the cytoskeleton and intracellular transportation are well established, but there is limited evidence related to their effects on functional changes in tumor cell viability and motility. As a consequence, efforts to investigate and reveal the multiple biological functions and mechanisms related to these proteins will potentially be of great benefit to clinical practice, and will provide a better understanding of this gene family.

Due to continuing progress in the field of bio-molecular technology and applications developed as a result of the Human Genome Project (HGP), there are increasing numbers of novel Rab genes being identified and cloned. Their structures and functions are distinct from typical Rabs. Some of them are being named Rab-like proteins. Novel gene Rabl3 is one such protein. To date, studies have revealed few functions for this gene, and there are no reports regarding the relationship between Rabl3 and tumorigenesis or tumor cell metastasis.

In the present study, we investigated the functional characterization of Rabl3 as a candidate oncogene and its

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engagement in multiple processes which influence viability, motility and drug resistance in human cancer cells.

## Materials and methods

**Reagents and materials.** Restricted endonucleases: *EcoRI* was purchased from Takara (DaLian, China); polyclonal antibody against Rab13 was obtained from ProteinTech Group Inc. (USA); monoclonal antibody against  $\beta$ -actin was purchased from Sigma (USA); IRDTMye 800-conjugated secondary IgG antibodies against rabbit or mouse were obtained from LICOR Bioscience (USA); paclitaxel was purchased from Haikou Pharmaceutical Factory Co., Ltd., China; DNA plasmid extraction kit was purchased from Qiagen (USA); FITC-conjugated Annexin V, and propidium iodide (PI) were purchased from Biosea Biotechnology Co., (Beijing, China); RMPI-1640 was purchased from Invitrogen (Carlsbad, CA, USA).

**Cell culture and transfection.** Human breast cancer cell line MDA-MB-231 and HeLa cell lines were cultured in RPMI-1640 medium containing 10% fetal bovine serum (FBS) and L-glutamine (2 mM). Other cell lines which were used for the analysis of the Rab13 expression profile were purchased from the American Type Culture Collection (ATCC). All of the cell lines were cultured in a humid incubator at a temperature of 37°C containing 5% CO<sub>2</sub>. Plasmid transfection of MDA-MB-231 or HeLa cells was performed using an ECM830 electroporation system (BTX, USA) at a voltage of 130 V, 20 ms, single pulse. Plasmid (10  $\mu$ g) was used for 1-3x10<sup>6</sup> transfected cells without FBS in 2-mm gap cuvettes.

**cDNA cloning and vector construction.** The full length of the Rab13 gene was obtained from the cDNA library of HeLa cell lines through PCR technology with the primer p1: 5'-ATGGCGTCCCTGGATCGGGT-3' and P2: 5'-CATCT A GATGAGCTGTGAAAACTGCC-3'. The PCR product was ligated into the T-easy vector and then cut off the insert through *EcoRI*. Subsequently the released Rab13 insert fragment was subcloned into the pcDNA3.1myc/hisB(-) *EcoRI* polycloned site, to form the pcDNA3.1myc/his(-)-Rab13 fusion plasmid. The plasmids mentioned above were confirmed by DNA sequencing.

**Rab13 siRNA synthesis.** The Rab13 specific siRNA was designed and synthesized by a method which targets the sequence of 5'-CAAGAGCAUAUCUACAATT-3'. The mock siRNA was designed with a sequence of 5'-UUCUCCGA ACGUGUCACGUTT-3'. Both siRNAs mentioned above underwent PAGE purification in an environment free of RNase contamination. In accordance with the manufacturer instructions, the siRNA was dissolved at the appropriate concentration (20  $\mu$ M). Dissolved siRNA (10  $\mu$ l) was used for 1-3x10<sup>6</sup> transfected cells in 400  $\mu$ l of FBS free RPMI-1640 during the electroporation procedure.

**RT-PCR analysis.** The total mRNA was extracted from the distinct cell lines by TRIzol reagent (Invitrogen, USA) and retranscribed using a iScript™ c-DNA synthesis kit (Bio-

Red, CA, USA) as indicated in the manufacturer's instructions. Total mRNA (1  $\mu$ g) was used for the RT procedure to synthesize the cDNA library at 42°C for 30 min. The cDNA libraries from indicated cell lines were then analyzed using Rab13 specific primers P3: 5'-TTGGGAGACTCAGGTGT TGGGAAA-3' and P4: 5'-CAGTTGGCACCAAATCCC TGTTGA-3' and the PCR products were separated by agarose electrophoresis. The GAPDH specific primers were also used as the internal control.

**Cell apoptosis and proliferation assay.** As the key index for apoptosis detection, Annexin V-PI staining was performed to measure phosphatidylserine externalization in MDA-MB-231 cells. Briefly, cells transfected with different plasmids were harvested at the indicated time points. Trypsinized cells were washed twice and resuspended in 200  $\mu$ l of binding buffer (10 mM HEPES, pH 7.4, 140 mM NaCl, 1 mM MgCl<sub>2</sub>, 5 mM KCl, 2.5 mM CaCl<sub>2</sub>). FITC-conjugated Annexin V (10  $\mu$ l) was then added to give a final concentration of 0.5  $\mu$ g/ml. The staining sample was incubated at room temperature for 20 min, without light. Subsequently, 5  $\mu$ l of PI was added to the samples (final concentration of 1  $\mu$ g/ml) and 10000 cells were immediately analyzed using a FACSCalibur flow cytometer (Becton-Dickinson). Results were calculated as a percentage of apoptotic cells. Cell proliferation detection was performed using a Cell Counting Kit-8 assay. According to the manufacturer's instructions, cells were paved into 96-well plates with 4000 cells per well. Cell proliferation intensity was calculated as the OD450 absorbent value for each well.

**Protein extraction and Western blot analysis.** The cells transfected with different plasmids or siRNAs were harvested at the indicated time points and washed twice with pre-cooled PBS. Then, an appropriate volume of lysis buffer (20 mM Tris HCl, pH 7.4, 150 mM NaCl, 1 mM EDTA, 1 mM EGTA, 1 mM PMSF and 1% Triton X-100) was added, sufficiently suspended, and reacted for 30 min at 4°C. After that, the supernatant was collected and assayed using the BCA protein quantification kit (Pierce, USA).

The same amounts of protein lysates were separated by SDS-PAGE gel and electroblotted into an NC membrane (Amersham Pharmacia, UK). Then the membrane was incubated in 5% BSA solution, dissolved using TBS-T (Tris-buffered saline containing 0.1% Tween-20) for 2 h at room temperature. Subsequently, specific primary antibodies were used and incubated overnight at 4°C, and secondary antibodies were incubated for 2 h at room temperature in the dark. The membranes were washed twice using TBST and scanned using an Odyssey Infrared Imager (LI-COR Bioscience) system.

**Wound healing assay.** The wound healing assay was used to detect cellular mobility. In our experiments, transfected cells were paved into 6-well plates and the cell layer was scratched using yellow tips. The scratched cells were washed twice with medium and cultured at 37°C. The wounds were photographed at indicated time points (0, 36 h) using phase contrast microscopy to monitor the wound healing process.

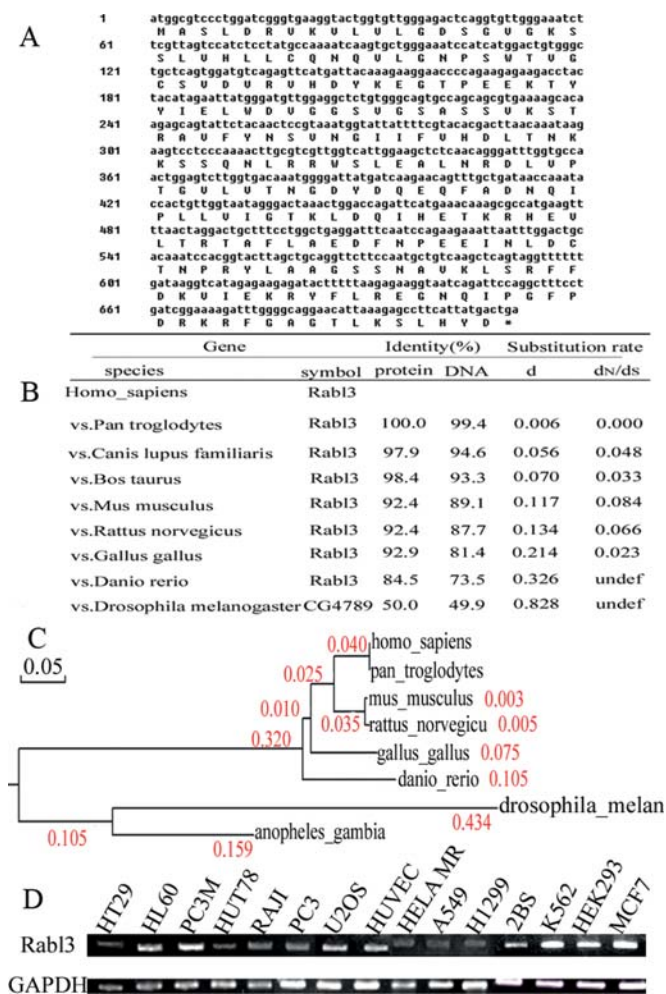


Figure 1. The novel Rab13 gene. (A) Nucleotide sequences and encoded amino acid sequences of human gene Rab13. (B) Sequence alignments of the Rab13 gene. The homology alignment results were indicated using identity degrees. (C) Phylogenetic analysis of Rab13. (D) The expression profile of the Rab13 gene analyzed using RT-PCR in various kinds of human cell lines originated from normal or cancer tissues.

**Cellular detachment assay.** The cellular detachment assay was performed to assess cellular adhesion ability. Transfected cells were paved into 6-well plates and cultured for 36 h. Cells were then treated with 0.005% trypsin in  $Mg^{2+}/Ca^{2+}$  free PBS. The trypsinized cells were photographed at indicated time points (0, 25 or 50 min). Cellular adhesive strength was assayed by determining relative sensitivity to trypsin.

**Statistical analysis.** All the experiments mentioned above were repeated at least 3 times. The results shown in each figure were calculated as the arithmetic means  $\pm$  SD. Statistical analysis was performed using the T-test.  $P < 0.05$  was considered as statistically significant.

## Results

**Identification and expression pattern of Rab13.** In the present study, the full-length coding region of Rab13 was directly isolated and amplified from the HeLa cell cDNA library by PCR using the specific primers P1 and P2. A 711 bp

complete cDNA product was obtained and found to encode 236 amino acids with the predicted molecular weight of 26.307 KD. The cloned ORF sequence of the Rab13 gene and encoded amino acid sequence are shown in Fig. 1A.

Based on the basic sequence analysis, human Rab13 is located on 3q13.3, and is composed of 8 exons and 7 introns. Upstream of the putative ATG start codon there exists an in-frame stop codon. The translation products of the Rab13 gene have the isoelectric point of 6.31. By searching the GeneBank database and aligning sequences among different eukaryotes, it was found that Rab13 is highly conserved in *Homo sapiens*, *Pan troglodytes*, *Mus musculus*, *Rattus norvegicus*, *Gallus gallus*, *Danio rerio*, *Drosophila melan* and *Anopheles gambia* (Fig. 1B and C).

Furthermore, the RT-PCR analysis was performed to identify the expression level in different cell lines. As shown in Fig. 1D, RT-PCR results clearly revealed that human gene Rab13 was expressed ubiquitously and abundantly in serial cell lines originating from normal tissues, as well as cancer tissues.

**Effects of Rab13 overexpression on cell motility in cancer cells.** It is known that cell motility plays a key role in tumor cell development and metastasis. To further evaluate the role of Rab13 in tumorigenesis, the effect of this gene on cell mobility and adhesion was studied using F-actin staining, and functionally assessed using the wound healing and cell detachment assays.

For the wound healing assay, cells were transfected with Rab13 and control plasmid and scratch wounding was carried out using a yellow pipette tip. Phase contrast images of the wound healing process were photographed at the indicated times. As shown in Fig. 2A, identical wound widths healed at different speeds in the different experimental groups. Human breast cancer cells overexpressed with Rab13 exhibited strong healing activity. In contrast, control cells exhibited relatively slow healing activity towards the middle of the wound. This indicated that Rab13 was crucial for promoting cell migration and suggested that it had an important role to play in the mobility and metastasis of tumor cells.

Furthermore, using the cell detachment assay, the results showed clearly that cells overexpressing Rab13 demonstrated strong adhesion abilities. They were more resistant to trypsin, as compared with the negative controls (Fig. 2B and C). This finding further demonstrated the key role played by Rab13 in the regulation of cell motility.

In addition, F-actin staining clearly revealed differences in morphological changes between Rab13 overexpressed MDA-MB-231 and control cells (Fig. 2D). F-actin in Rab13 transfected cells exhibited stronger staining with a cortical distribution, as compared with negative control cells. This finding also potentially supports the positive effects of Rab13 on cell motility.

**Effects of silencing gene Rab13 on cell viability, motility and apoptosis in MDA-MB-231 cells.** Since detection of functional changes in Rab13 downregulated cells was considered necessary for our physiological studies, siRNA against Rab13 was designed, synthesized and then used to silence the expression of Rab13 in cancer cells. Non-silencing



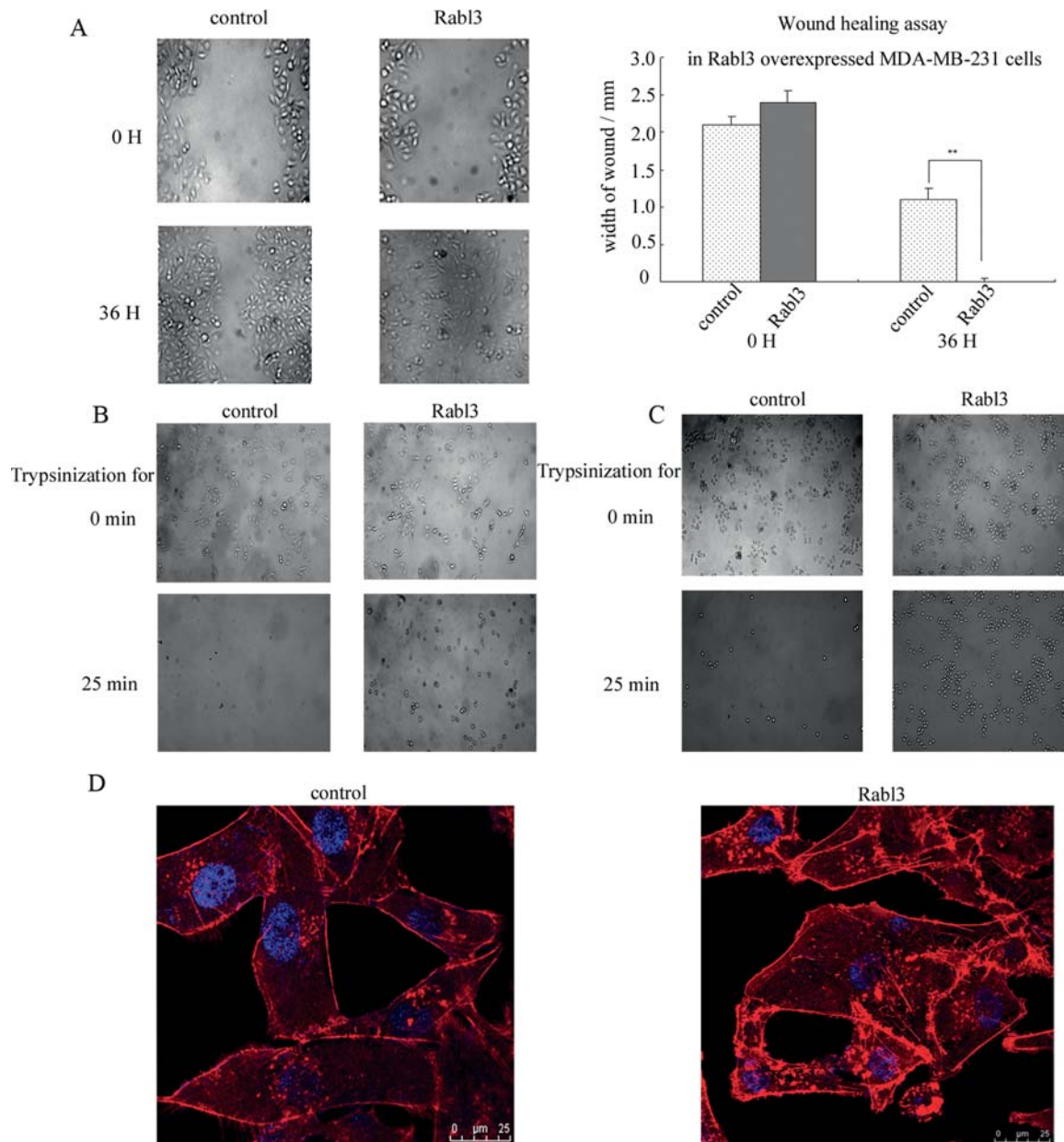


Figure 2. Detection of cellular motility changes in Rab13 overexpressed cells. (A) Wound healing assay in Rab13 overexpressed MDA-MB-231 cells. Phase contrast photographs showed that overexpression of Rab13 enhanced the mobility of MDA-MB-231 cells at 36 h after transfection. The wound width can be used as the index reflecting the ability of cell migration between different experiment teams. (B) Cell detachment assay in Rab13 overexpressed MDA-MB-231 cells. Cells were trypsinized using 0.005% trypsin in  $Mg^{2+}/Ca^{2+}$  free PBS at 36 h after transfection. Phase contrast photographs were taken at the indicated time points after trypsinization. The density of adherent cells remaining in the plate represents adherent ability in different experimental groups. (C) Cell detachment assay in Rab13 overexpressed HeLa cells. (D) F-actin staining with Phalloidin analyzed by confocal microscopy (Olympus, Tokyo, Japan). The pictures revealed that the Rab13 overexpressed MDA-MB-231 cells exhibited a stronger signal of F-actin with cortical distribution.

siRNA termed 'si-non-RNA' was also used at the same time as the negative control.

RT-PCR and Western blot analysis confirmed that the Rab13 specific si-RNA was efficient in blocking both the Rab13 transcriptional and expressional procedures for 24 to 48 h after electroporation. In contrast, si-non-RNA had no effect on the Rab13 expression (Fig. 3A).

To confirm that downregulation of Rab13 could inhibit cell proliferation and initiate apoptosis, the CCK-8 assay and Annexin V-PI staining were performed. Compared with the previous results obtained using Rab13 overexpressed cells, the OD 450 value for Rab13 downregulated cells was signi-

ficantly lower ( $P < 0.05$ ) than that for the si-non-negative control group (data not shown). Next, we evaluated whether Rab13 downregulation could restore or even enhance PCD condition. As shown in Fig. 3B, the findings from phosphatidylserine staining strongly supported this hypothesis, in that the fraction of apoptotic cells in si-Rab13 transfected cells was obviously higher than in si-non cells. These results provided more evidence that Rab13 functions as a key growth stimulator during tumor cell proliferation.

In addition, the relative migration abilities of si-Rab13 and si-non transfected cells were compared. The results clearly demonstrated that a lack of Rab13 expression caused

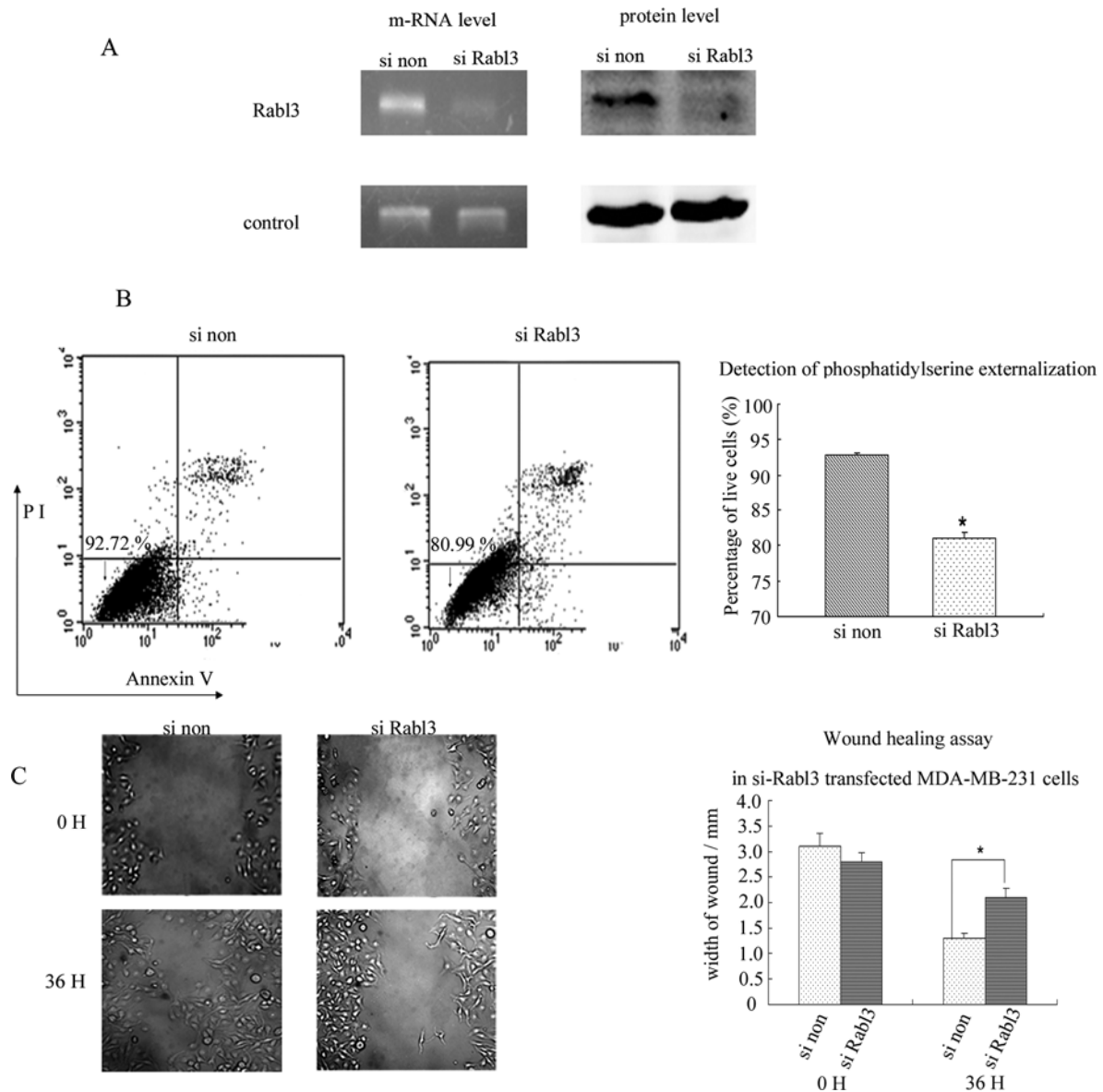


Figure 3. Detection of cellular viability and motility changes in Rabl3 siRNA transfected cancer cell lines. (A) Identification of siRNA against Rabl3 both in mRNA level and protein level. Cells were harvested at 36 h after transfection. (B) Detection of PS externalization in non-silencing and siRabl3 transfected MDA-MB-231 cells. Cells were harvested at 48 h after transfection. The percentages of Annexin V or PI single positive and Annexin V/PI double positive cells were calculated as the apoptotic intensity in different cells. The results were further quantified as the percentage of live cells shown in the diagram below. (C) Wound healing assay in siRNA transfected MDA-MB-231 cells. The width of the wound represents the wound healing ability in different experimental groups.

significant downregulation of migration activity in MDA-MB-231 cells. The wound healing ability of these cells appeared lower than that of negative controls (Fig. 3C). Therefore, these data suggested that Rabl3 may be involved in both cell survival and migration.

*Evaluation of FAK phosphorylation in Rabl3 overexpressed cancer cells.* It is widely accepted that complicated regulation is involved in tumorigenesis and that there is a loss of the tight control on cell growth or apoptosis. To investigate how Rabl3 modulates cellular proliferation and migration processes, we screened and analyzed various signal pathways. We found a high phosphorylation level of FAK in Rabl3 overexpressed MDA-MB-231 cells. As is known, FAK is a non-receptor tyrosine kinases (NRPTK) which

is localized in focal adhesions, multi-molecular structures involved in connections between the cytoskeleton and extra-cellular matrix (12). FAK is also a potential candidate signaling molecule. It is capable of receiving and integrating multiple signals transduced from integrin and regulating diverse signaling pathways downstream of FAK (13). Therefore, FAK mediates fundamental effects associated with complicated biological process, such as cell proliferation, migration and adhesion (14). In particular, for many kinds of cancers, FAK is a key signaling trigger which has close connection with cancer cell invasion and metastasis (15). To date, five phosphorylation sites have been identified on the FAK molecule, including Tyr397/577/861/925. Phosphorylation of these sites activates FAK into its fully functional state (16). In particular, the phosphorylation sites of Tyr

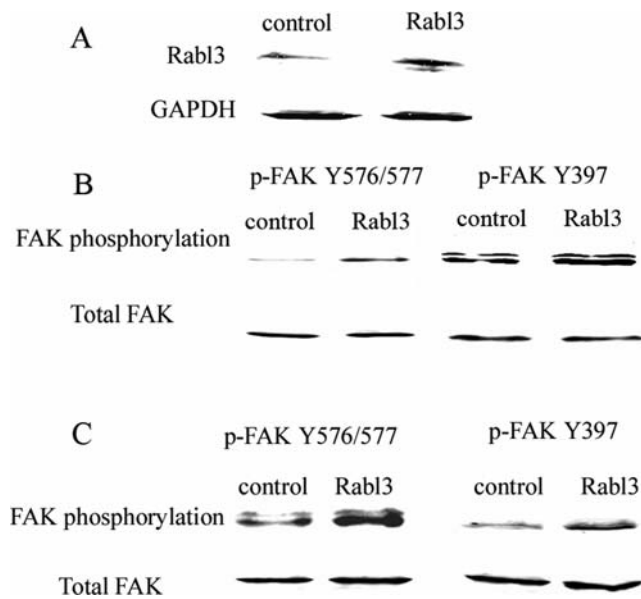


Figure 4. Rabl3 overexpression was accompanied by FAK phosphorylation. (A) Rabl3 overexpression identification in HeLa cells using anti-Rabl3 specific antibody at 36 h after transfection. (B) Rabl3 overexpression induced the increased FAK phosphorylation at the sites of Y397/576/577 in HeLa cells at 36 h after transfection. T-FAK was also detected as the internal control. (C) Western blot analysis of FAK phosphorylation in MDA-MB-231 cells.

576/577 endow the molecule with the strongest kinase activity.

Using Western blot analysis, we found that after efficient Rabl3 overexpression (Fig. 4A), the phosphorylation level of FAK Y397/576/577 was clearly higher than the level in empty vector transfected control cells. At the same time, the level of total FAK was found to be constant (Fig. 4B and C). This finding indicated that Rabl3 could strengthen FAK phosphorylation and then activate FAK, thus displaying the ability to activate proliferation and motility functions in cancer cells.

**Sensitization of MDA-MB-231 cells to paclitaxel by inhibition of the Rabl3 gene.** Over the years, there has been increasingly extensive use of the potent chemotherapeutic agent, paclitaxel. This agent is the first member of the taxel family (17) and is effective in treating certain types of cancer (18). Paclitaxel has been shown to stabilize microtubule polymers, leading to mitotic arrest and apoptosis (19). However, drug-resistance is a major but unsolved problem which limits the clinical efficacy of paclitaxel in approximately 50% of patients with breast cancers or other solid tumor types (20). In the case of patients undergoing long-term treatment with paclitaxel, the occurrence of drug-resistance is the primary obstacle to successful therapy (21). The precise mechanisms by which paclitaxel exerts its cytotoxic effects in tumor cells and the basic mechanism causing resistance to the drugs are still under wide discussion.

Our functional research related to Rabl3 revealed important influences on paclitaxel resistance in human breast cancer MDA-MB-231 cell lines. First of all, we tested

paclitaxel sensitivity in MDA-MB-231 cells. As shown in Fig. 5A, paclitaxel cytotoxicity against MDA-MB-231 cells was concentration dependent. Photographs clearly showed that cells exhibited evidence of extensive cytoplasm vacuolization, blebbing and increased numbers of apoptotic cells detached from the cultural matrix after paclitaxel treatment. These findings indicated that MDA-MB-231 cells had high sensitivity to paclitaxel. Annexin V-PI staining clearly revealed that in overexpressed Rabl3 cells, the fraction of apoptotic cells induced by 7  $\mu$ mol/l paclitaxel was relatively low at 24 h after the agent administration. However, we found that transfecting MDA-MB-231 cells with siRNA against Rabl3 resulted in the induction of significant numbers of apoptotic cells compared with the negative control, using paclitaxel at a concentration of 7  $\mu$ mol/l (Fig. 5B and C). These changes in the proportion of apoptotic cells induced by paclitaxel, due to manipulation of the novel Rabl3 gene, indicate that there could be potential clinical benefits in targeting this gene.

In summary, the initial findings of our studies suggest that Rabl3 plays an active role in the generation of paclitaxel resistance. We demonstrated that silencing this gene can substantially restore, or even reverse, tumor cell resistance to paclitaxel.

## Discussion

In the present study we investigated a novel human gene, Rabl3, which is highly conserved in a range of animal species. This gene is a member of the Rab subfamily. Rabs are mostly noted for the key roles that they play in regulating the formation of the cytoskeleton and dynamic transportation. It is clear that Rabs, although not usually considered oncogenic products, could profoundly affect the proliferation, survival and invasiveness of cancer cells. The Rab family loss or gain-of-function resulting from deletion or gene amplification could therefore promote cell transformation and/or enhance tumor cell motility.

The findings of the present study extend our understanding of the functional scope of this gene subfamily, and have contributed new evidence that Rabl3 functions as a positive regulator in connection with the processes of tumor cell viability and motility.

Revealing the signaling molecules involved in the Rabl3 oncological effects is a crucial area for further investigation. We have demonstrated for the first time that Rabl3 related signaling proteins play important roles in cellular proliferation and migration. Western blot analysis clearly showed that FAK phosphorylation levels were elevated as a result of Rabl3 overexpression at the sites of Y397/576/577. This finding implies a close relationship between Rabl3 and FAK. Since the FAK/Erk (Extracellular Regulatory Kinase) interaction is known to regulate many signaling transductions, and the functions of Erk in cell migration and survival are well documented (22,23), we hypothesize that Rabl3 is a component in the FAK-Erk signaling pathway. Rabl3 have also been linked to cancer cell viability and migration, however, the underlying mechanism is unclear. Further studies need to be performed to reveal the exact biological behavior and interaction of FAK or other proteins.



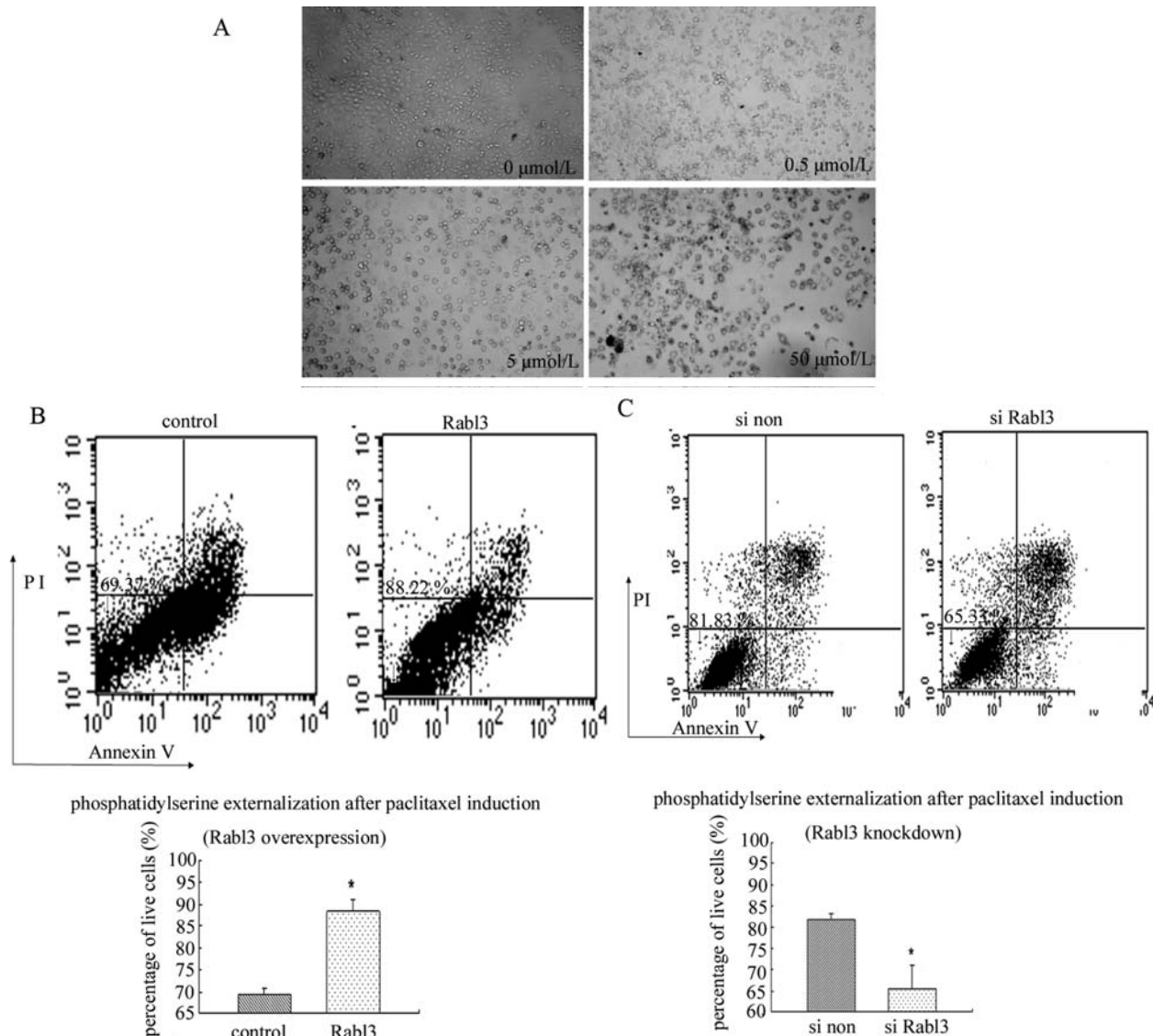


Figure 5 Sensitization of MDA-MB-231 cells to paclitaxel by inhibition of the Rab13 gene. (A) The identification of paclitaxel sensitivity in MDA-MB-231 cells. The cytotoxicity induced by paclitaxel was both concentration and time-dependent in MDA-MB-231 cells. (B) Overexpression of Rab13 induced paclitaxel resistance in MDA-MB-231 cells. The numbers of PI/Annexin V single positive and Annexin V/PI double positive cells were calculated as the measurement of apoptotic cells. The diagram shown below was utilized as the measurement of live cells within the total amounts of  $1 \times 10^4$  detected cells. (C) Silencing of Rab13 expression increased paclitaxel induced apoptosis in MDA-MB-231 cells.

Anti-tumor agent resistance and non-sensitive treatment has become the most concentrated obstacles for cancer therapy; therefore, finding a new effective target of anti-drug resistance is very important. We have found that the novel human gene Rab13 showed clear activity relating to paclitaxel sensitivity in breast cancer cells. Our results demonstrated that overexpression of Rab13 could prevent tumor cells from undergoing paclitaxel-induced apoptosis, and as a consequence elevate resistance to paclitaxel. Silencing of Rab13 expression could increase paclitaxel induced apoptosis in MDA-MB-231 cells. Given the importance of RAB GTPases in regulating many cellular functions, it is not surprising that altered expression or mutation of Rab proteins and their interacting partners may cause human diseases. However, as different kinds of PCD induced by paclitaxel can be triggered through a range of different mechanisms (24-27), the exact mechanism

underlying Rab13 mediated paclitaxel resistance remains unknown. Further studies with regard to this problem are needed.

The studies represent an important role of Rab13 in cancer treatment. Based on recent research, substantial evidence indicated that some RAB gene members might be useful for prognostic and/or diagnostic markers in cancer therapy practice. Progress in this area is likely to improve our understanding of not only the role of RAB GTPase in the biology of cancer but also the essential role of cellular trafficking in the maintenance of a number of normal physiological processes. As the underlying mechanism of Rab-mediated tumorigenesis comes to light, it is certain that Rab protein and their interaction molecules will represent a novel therapeutic target.

In conclusion, we have demonstrated that Rab13 can enhance cell proliferation and migration activity, and inhibit

apoptosis in human cancer cell lines exogenously. We also showed that silencing the Rab13 gene can generate wholly opposite changes in cell activity. Moreover, we found that the biological functions of Rab13 are largely connected with FAK phosphorylation at the sites of Y397/576/577. Importantly, our findings have potential clinical implications. Loss of Rab13 expression will cause an elevated sensitivity to paclitaxel in breast cancer cells, therefore, implicating Rab13 might also be new potential targets for therapeutic strategies against breast cancer in viability and metastasis.

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