Tumor Cell Migration and Invasion Are Enhanced by Depletion of Rap1 GTPase-activating Protein (Rap1GAP)

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Background: Although widespread, the significance of Rap1GAP down-regulation in human tumors is unknown.

Results: Genetic silencing of Rap1GAP in human colon cancer cells confers a mesenchymal mode of cell migration and enhances invasive behavior.

Conclusion: Rap1GAP regulates the mechanism of cell motility.

Significance: Down-regulation of Rap1GAP endows tumor cells with more aggressive properties.

The functional significance of the widespread down-regulation of Rap1 GTPase-activating protein (Rap1GAP), a negative regulator of Rap activity, in human tumors is unknown. Here we show that human colon cancer cells depleted of Rap1GAP are endowed with more aggressive migratory and invasive properties. Silencing Rap1GAP enhanced the migration of confluent and single cells. In the latter, migration distance, velocity, and directionality were increased. Enhanced migration was a consequence of increased endogenous Rap activity as silencing Rap expression selectively abolished the migration of Rap1GAP-depleted cells. ROCK-mediated cell contractility was suppressed in Rap1GAP-depleted cells, which exhibited a spindle-shaped morphology and abundant membrane protrusions. Tumor cells can switch between Rho/ROCK-mediated contractility-based migration and Rac1-mediated mesenchymal motility. Strikingly, the migration of Rap1GAP-depleted, but not control cells required Rac1 activity, suggesting that loss of Rap1GAP alters migratory mechanisms. Inhibition of Rac1 activity restored membrane blebbing and increased ROCK activity in Rap1GAP-depleted cells, suggesting that Rac1 contributes to the suppression of contractility. Collectively, these findings identify Rap1GAP as a critical regulator of aggressive tumor cell behavior and suggest that the level of Rap1GAP expression influences the migratory mechanisms that are operative in tumor cells.

Altering the cytoskeleton, cell/cell adhesion, and integrin activation are critical nodes in the transition from benign to invasive carcinomas. Stable expression of activated Rap enhanced metastasis in prostate cancer cells and the infiltration of breast cancer cells into the vasculature (25, 26). However, the significance of these studies to human tumors is unclear in that activating mutations in Rap have not been reported (27). Down-regulation of Rap1GAP is widespread in human tumors (28–33). Overexpression of Rap1GAP in human tumor cells impaired cell migration and invasion in vitro (28, 29, 31, 33–35) and metastasis in vivo (36, 37). Intriguingly, the expression of ectopically expressed Rap1GAP was lost from disseminated tumors but retained in those that formed at the sites of subcutaneous injection (36). This supports the existence of selective pressure to decrease Rap1GAP expression, which appears to be operative in human tumors where the expression of Rap1GAP decreases with tumor progression (30, 31, 33, 35).

The cellular processes that are sensitive to the levels of Rap1GAP are unknown. Importantly, whether the widespread down-regulation of Rap1GAP observed in human tumors alters the behavior of tumor cells has not been determined. We previously reported that silencing the expression of Rap1GAP in human colon cancer cells weakened cell/cell adhesion and enhanced spreading on collagen, changes that are reminiscent of those that take place during the early stages of tumor cell dissemination (32). We demonstrate that silencing Rap1GAP endows cells with a Rap- and Rac1-dependent mechanism of cell motility that was inactive in parental cells. Invasive behavior was profoundly up-regulated in Rap1GAP-depleted cells. Collectively, these findings suggest that down-regulation of Rap1GAP in human tumors harbors the potential to increase migratory and invasive behaviors that promote tumor progression.

EXPERIMENTAL PROCEDURES

Cell Culture and Transfection—HT29 and LoVo cells were cultured in RPMI 1640 medium containing 10% fetal bovine serum (FBS). The isolation of the Rap1GAP-depleted HT29 cell lines was described previously (32). In brief, SMARTvector

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2 The abbreviations used are: Rap1GAP, Rap1 GTPase-activating protein; MLC, myosin 2 light chain.
Silencing Rap1GAP accelerates wound closure. A, photomicrographs of Rap1GAP-depleted cell lines (E11, C10, C4, F3) and a control (Con) cell line selected to express a nonspecific shRNA immediately after wounding (0) and after stimulation with FBS for 20 h are shown. Scale bar, 20 μm. B, total cell lysates prepared from Rap1GAP-depleted and control cells were subjected to Western blotting for Rap1GAP and actin to confirm equal protein loading. C, increased wound closure in Rap1GAP-depleted cells was statistically significant (***, p < 0.0001). Error bars, S.E. D, photomicrographs show parental HT29 and control cells in wound closure experiments conducted in the absence and presence of the Src inhibitor PP2 (10 μM, gray bars) or inactive analog PP3 (10 μM, black bars). Inhibitors were added for 60 min prior to the addition of serum-supplemented medium. Scale bar, 20 μm. E, results from five wound closure experiments comparing control and HT29 cells are shown. F, photomicrographs show Rap1GAP-depleted (E11) and control (Con) cells at the wound border. Asterisks show examples of elongated protrusions on Rap1GAP-depleted cells. Scale bar, 20 μm.

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human Rap1GAP and nonspecific control were purchased from Thermo Scientific (Dharmacon). Viruses expressing two different Rap1GAP-directed shRNAs were used, and multiple clones expressing each were isolated and analyzed. Cell lines E11 and C10 were isolated using shRNA#2 and the C4 and F3 lines isolated using shRNA#3. Control cell lines (Con) were isolated from cells expressing a nonspecific shRNA (32). All experiments were conducted in three or more Rap1GAP-deficient cell lines, including one made with each shRNA with similar results and compared with control and parental HT29 cells. For acute silencing, cells were transfected with Rap1/2 or Rap1GAP-directed and nonspecific siRNAs using Amaza-mediated electroporation as described previously (32).

Wound Closure Assays—Cells were grown to confluence on 6-well plates marked with a line down the center and coated with 2 μg/ml collagen IV. Cells were starved for 24 h, and wounds (5–6/well) were made perpendicularly to the line. After wounding, cells were stimulated with serum-supplemented medium. Images were acquired using a Nikon Eclipse TE2000 microscope and Northern Eclipse software (Empix Imaging). The distances between the intersection of the mark and the border of the wound were measured using Multi Gauge Imaging). The distances between the intersection of the mark and the border of the wound were measured using Multi Gauge software (Fujifilm). The distance between the wound borders at 20 h was subtracted from that at 0 h and the difference expressed as percentage closure. The percentage closure from 5–6 wounds/dish was averaged in each experiment.

Phagocytic Tracking Assays—Fluorescent beads (Cellomics Cell motility kit; Thermo Scientific) were plated on collagen IV-coated glass slips. Starved cells (10⁶ cells/ml) were plated under conditions where the cells were subconfluent, allowed to attach for 2 h, and then stimulated with serum-supplemented media. After wounding, cells were stimulated with serum-supplemented medium. Images were acquired using a Nikon Eclipse TE2000 microscope and Northern Eclipse software (Empix Imaging). The distances between the intersection of the mark and the border of the wound were measured using Multi Gauge software (Fujifilm). The distance between the wound borders at 20 h was subtracted from that at 0 h and the difference expressed as percentage closure. The percentage closure from 5–6 wounds/dish was averaged in each experiment.

Results from three experiments comparing track length in Rap1GAP-depleted cells with control cells. Track length in control cells was set to 1.0. B, photomicrographs of control (Con) and Rap1GAP-depleted (C10) cells in phagocytic tracking assays. Additional Rap1GAP-depleted cells (E11, C4) are shown in Fig. 6F. C, Rap1GAP-depleted and control cells transiently transfected with siRNAs directed to Rap1 and Rap2 (open bars) or nonspecific (NS) siRNAs (filled bars). Phagocytic tracking assays were performed at 48 h after transfection. Silencing Rap significantly (*, p < 0.05) decreased migration in Rap1GAP-depleted cells. Error bars, S.E. D, photomicrographs of siRNA-transfected cells in tracking assays. E, total cell lysates prepared from the cells used in the tracking assays analyzed for Rap1/2 expression by Western blotting. Western blotting for actin confirmed equal protein loading. F, control and Rap1GAP-depleted (C4) cells transiently transfected with siRNAs directed to Rap1/2 (open bars) or nonspecific (NS) siRNAs (filled bars) and analyzed in wound closure experiments. Silencing Rap significantly (*, p < 0.05) impaired wound closure in Rap1GAP-depleted cells. G, total cell lysates prepared from the cells used in the wound closure assays analyzed for Rap1/2 expression by Western blotting. Western blotting for actin confirmed equal protein loading.
medium for 24 h. Where inhibitors were used, cells were plated in the absence of inhibitor and stimulated with inhibitor-supplemented medium. Cells were fixed and stained for F-actin, and the tracks of individual cells were analyzed on a Zeiss Axio-phot fluorescence microscope using Zeiss Axiovision software.

**Time Lapse Microscopy**—Cells (20,000/well) were plated on collagen IV-coated chamber slides. Phase contrast images of cells in serum-supplemented medium were acquired every 10 min over 16 h using a Deltavision Deconvolution Microscope (Applied Precision, Inc.). Tracks of randomly selected single cells were analyzed by manual tracking using ImageJ software (National Institutes of Health). The distance, velocity, and directionality (calculated as net displacement/total path length) of migration were determined using chemotaxis software.

**Immunostaining**—Cells plated on collagen IV-coated coverslips were starved overnight and stimulated with 10% FBS for varying times. Cells were fixed in 3.7% formaldehyde in PBS and stained with rhodamine-phalloidin or antibodies to FAK-Y397p followed by an Alexa Fluor 488-conjugated secondary antibody. Within an experiment, images for a given antibody were acquired for the same times.

**Antibodies and Western Blotting**—Rap1 (sc-65), Rap1GAP (sc-28189), and actin antibodies were obtained from Santa Cruz Biotechnology. Rap2 (610215) and FAK-Y397p (611723) anti-

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**FIGURE 3. Migration distance, velocity, and directionality are enhanced by Rap1GAP depletion.** A and B, graphical representations show tracks made by seven individual (A) HT29 and (B) Rap1GAP-depleted (E11) cells as assessed by time lapse microscopy. C–E, the migration of 21 HT29 and E11 cells was analyzed for distance (C), velocity (D), and directionality (E) as described under “Experimental Procedures.” Distance (**, \( p < 0.005 \)), velocity (**, \( p < 0.005 \)), and directionality (*, \( p < 0.05 \)) were significantly increased in Rap1GAP-depleted cells. Error bars, S.E.
bodies were from BD Transduction Laboratories. MLC2-T18/S19p antibody was from Cell Signaling. For total cell lysates, cells were disrupted in Triton buffer (20 mM Tris-HCl, pH 7.4, 100 mM NaCl, 1% Triton X-100, 2 mM EDTA, 1 mM NaF, plus protease inhibitors). Protein concentrations were determined by DC protein assay (Bio-Rad) and equal micrograms of protein analyzed.

**Invasion Assays**—Cells (100,000) were mixed with 100 μl of ice-cold growth factor-reduced Matrigel (BD 356230), plated onto Transwell filters (Corning, 8 μm, 3422) and incubated for 30 min at 37 °C until hardened. Filters were placed into a well containing 600 μl of growth medium (± inhibitors) and 100 μl of serum free medium (± inhibitors) added to the cells in the upper chamber. After 48 h, Matrigel was removed from the upper well with a cotton swab, and cells on the bottom of the filter were fixed in 3.7% formaldehyde, stained with 0.1% crystal violet, washed with water, and counted. For outgrowth in Matrigel, cells (10,000) were mixed with 160 μl of Matrigel and plated in 35-mm Mattek dishes containing a glass coverslip. After hardening, cells were overlaid with 2 ml of growth medium and incubated at 37 °C for up to 11 days.

**Statistical Analysis**—All experiments were repeated at least three times. The bar graphs shown represent the combined results from at least three experiments unless otherwise noted. Statistical significance was determined by Student’s t test.

**RESULTS**

**Down-regulation of Rap1GAP Enhances Cell Motility**—To explore the consequences of Rap1GAP down-regulation on tumor cell motility, wound closure assays were conducted in four independent lines of Rap1GAP-depleted HT29 cells (32). Confluent monolayers of cells, plated onto collagen IV to mimic...
adhesion to the basement membrane, were grown to confluence, starved for 18 h, wounded, and stimulated with serum-containing medium. Images were acquired immediately after wounding and after 20 h in serum-supplemented medium. Down-regulation of Rap1GAP significantly accelerated wound closure compared with cells selected to express a nonspecific shRNA (control) (Fig. 1, A–C). Wound closure in control cells was unchanged in the absence or presence of inhibitors (Src inhibitor, PP2 shown here) compared with parental HT29 cells (Fig. 1, D and E).

We previously observed that silencing Rap1GAP impaired cell/cell adhesion (32). Indeed, Rap1GAP-depleted cells along the border of the wound were more dispersed than control cells, which maintained cell/cell contacts (Fig. 1F). Rap1GAP-depleted cells elaborated elongated membrane protrusions that were not observed in control cells (examples marked by asterisks in Fig. 1F). These findings prompted us to investigate whether depleting Rap1GAP altered migratory mechanisms.

Because weakened cell/cell adhesion could enhance wound closure, we determined whether the migratory activity of single cells was enhanced by Rap1GAP depletion using phagocytic tracking assays. Cells were plated on a lawn of beads on collagen IV, and migration in the presence of serum was analyzed. The tracks made by all four lines of Rap1GAP-depleted cells were longer (Fig. 2A) and less circuitous (C10 shown in Fig. 2B; E11 and C4 in Fig. 6F) than those made by control cells. Rap1 and Rap2 activity are durably increased in Rap1GAP-depleted cells (32). Therefore, we determined whether enhanced migration was a consequence of increased Rap activity. Silencing Rap1 and Rap2 expression significantly impaired the migration of Rap1GAP-depleted cells in phagocytic tracking (Fig. 2, C–E) and in wound closure assays (Fig. 2, F and G). The migration of control cells was not affected by silencing Rap (Fig. 2, C and F). The selective requirement for Rap activity in Rap1GAP-depleted cells implies that cells acquire additional or altered means of cell motility when endogenous Rap activity is increased. This is the first demonstration that down-regulation of Rap1GAP, an event widely observed in human tumors, enhances migratory behavior.

Individual cell migration was further analyzed in time lapse microscopy experiments. HT29 cells migrate in a circuitous fashion, characterized by frequent alterations in the direction of cell migration (Fig. 3A). Rap1GAP-depleted cells made fewer turns and migrated over greater distances (Fig. 3B). Migration distance, velocity, and directionality were increased in Rap1GAP-depleted cells (Fig. 3, C–E).

To confirm that increased migration was due to depletion of Rap1GAP rather than secondary events that occurred during the isolation of stable cell lines, Rap1GAP expression was acutely silenced in HT29 cells. Cells were transfected with Rap1GAP-directed or nonspecific siRNAs and cell migration analyzed at 48 h after transfection. The Rap1GAP siRNA targeted a region different from either of the two shRNAs used to isolate the Rap1GAP-depleted cell lines. Cells in which Rap1GAP expression was acutely silenced exhibited an increase in the distance, velocity, and directionality of migration (Fig. 4, A–F). Therefore, enhanced migration is likely a primary consequence of Rap1GAP depletion.

Enhanced Invasive Activity in Rap1GAP-depleted Cells—Rap1GAP-depleted cells exhibited weakened cell/cell adhesion, enhanced spreading on collagen (32) and increased migratory activity, changes that resemble those seen in invasive cancer cells. To explore the consequences of Rap1GAP depletion on invasive behavior, cells were plated in Matrigel in the upper well of Transwell chambers and exposed to serum in the lower chamber. Control and HT29 cells were poorly invasive (Fig. 5, A and B). Invasion was significantly increased in Rap1GAP-depleted cells.

To explore effects on invasive activity under conditions that more closely mimic those in vivo, cells were embedded in Matrigel, and their ability to invade the matrix in three dimensions was analyzed. Control and HT29 cells formed compact, dense spheres under these conditions (Fig. 5, C–E). The boundaries of the colonies were smooth, indicative of robust cohesive activity. Single cells or dispersed colonies were rarely seen. In contrast, Rap1GAP-depleted cells actively invaded the surrounding matrix (Fig. 5, C and D). Single cells and small clusters of cells separate from the large colonies were routinely observed. In conclusion, depletion of Rap1GAP endows cells with enhanced migratory and invasive activity. These results suggest that tumors in which Rap1GAP expression is down-regulated harbor the potential for more aggressive behavior.
Loss of Rap1GAP Suppresses ROCK-mediated Actomyosin Contractility—Cell migration and invasion entail dramatic reorganization of the actin cytoskeleton. To explore the consequences of Rap1GAP depletion on cytoskeletal dynamics under the same conditions used to monitor cell motility, the acute effects of serum on the actin cytoskeleton were investigated in cells plated on collagen IV. Serum induced profound membrane blebbing in control (Fig. 6, A and D) and HT29 (Fig. 6, B and C) cells. Membrane blebbing was most pronounced at 5 min and maintained for up to 120 min after serum stimulation. Membrane blebbing was suppressed in Rap1GAP-depleted cells (Fig. 6, A and B). Similar results were observed when Rap1GAP was acutely silenced in HT29 cells (Fig. 6 C).

Cancer cells are endowed with multiple mechanisms of cell motility. The migration of single tumor cells has been characterized as amoeboid (also called rounded) or mesenchymal (38–41). Amoeboid/rounded cell migration is driven by high levels of ROCK-mediated contractility that underlie profound membrane blebbing. To assess whether membrane blebbing reflected actomyosin-mediated contractility, cells were exposed to inhibitors of ROCK (Y27632) and myosin ATPase (blebbistatin) activity. Membrane blebbing in control and parental HT29 cells required ROCK and myosin ATPase activity (Fig. 6, D and E). To assess whether migration was ROCK-dependent, the effects of Y27632 on individual cell migration were analyzed. The migration of control and Rap1GAP-depleted cells required ROCK activity (Fig. 6F).

Depletion of Rap1GAP Confers Rac1-dependent, Mesenchymal Cell Migration—Cells that migrate via a mesenchymal mechanism exhibit shared features including a spindle-shaped morphology, the elaboration of actin-rich membrane protrusions, and dependence on Rac1 activity (for review, see Ref. 40). Unlike control and HT29 cells, which were round, Rap1GAP-depleted cells were elongated and exhibited actin-rich protrusions (examples indicated by arrows in Fig. 6, A and B). Focal adhesions were uniformly distributed around the circumference of control and HT29 cells (Fig. 7, A and B). In Rap1GAP-depleted cells, focal adhesions were less uniform in distribution. To assess whether migration was Rac1-dependent, cells were treated with the Rac1 inhibitor, NSC23766 (42). Strikingly, the migration of Rap1GAP-depleted, but not of control cells, required Rac1 activity (Fig. 7, C–E). Moreover, in the presence of the Rac1 inhibitor, the tracks made by Rap1GAP-depleted cells reverted to a circuitous pattern more similar to that
in control cells (Fig. 7C). The selective requirement for Rac1 activity in the migration of Rap1GAP-depleted cells was confirmed using β2-chimaerin, a highly specific Rac1GAP (43). Expression of β2-chimaerin impaired wound closure in Rap1GAP-depleted, but not control or parental HT29 cells (Fig. 7, F and G). Inhibition of Rac1 also restored a rounded morphology and a more uniform distribution of focal adhesions to Rap1GAP-depleted cells (Fig. 7, A and B). Together, these find-
ings demonstrate that depletion of Rap1GAP endows cells with an altered mechanism of cell motility that requires Rap and Rac1 activity.

**Rac1 Suppresses ROCK-dependent Contractility in Rap1GAP-depleted Cells**—Overexpression of activated Rac1 in human tumor cells induced a switch from an amoeboid/rounded mechanism of cell migration to a mesenchymal mechanism (44). To explore the contribution of Rac1 to the suppression of cell contractility in Rap1GAP-depleted cells, the effects of Rac1 inhibition on membrane blebbing were examined. Treatment with NSC (Fig. 8A) or expression of β2-chimaerin (Fig. 8B) induced membrane blebbing in Rap1GAP-depleted cells. Because membrane blebbing was ROCK-dependent, we assessed whether Rac1 impaired ROCK activity. The phosphorylation of two ROCK substrates, the myosin-binding subunit of myosin phosphatase (MYPT1) and myosin 2 light chain (MLC), were examined. Western blotting with two different antibodies revealed that MYPT1 was basally phosphorylated in Rap1GAP-depleted and control cells and that phosphorylation was unaffected by serum stimulation or ROCK inhibition (data not shown). Serum stimulated MLC phosphorylation on serine 18/threonine 19 (Fig. 8, C–F). Although this site is phosphorylated by ROCK and myosin light chain kinase, only inhibition of ROCK blocked MLC phosphorylation (Fig. 8, C–E). Expression of β2-chimaerin increased basal and serum-stimulated MLC phosphorylation in Rap1GAP-depleted cells (Fig. 8, F and G). Similar effects were observed using NSC (Fig. 8H). We conclude that the suppression of contractility in Rap1GAP-depleted cells is not due to global inhibition of ROCK activity, but that the increase in Rac1 signaling suppresses contractility at least in part through inhibition of ROCK activity.

**Down-regulation of Rap1GAP Enhances Migration and Invasion in Other Colon Cancer Cell Lines**—To confirm that the effects of Rap1GAP depletion on migratory and invasive behavior were not restricted to a single cell line, the expression of Rap1GAP was acutely silenced in the human colon cancer cell line, LoVo. LoVo cells were selected for analysis as these cells express Rap1GAP (Fig. 9, B and C). Silencing Rap1GAP increased the migration of individual LoVo cells in phagocytic tracking assays (Fig. 9, A and B). Furthermore, acute silencing of Rap1GAP promoted outgrowth in Matrigel (Fig. 9C). Thus, the consequences of Rap1GAP depletion on tumor cell behavior are likely to be general.

**DISCUSSION**

The significance of the widespread down-regulation of Rap1GAP expression in human tumors is unknown. Our pre-
Manipulating the balance in Rac1 and Rho activity in tumor cells induced a switch from a contractility-based, rounded mechanism of cell motility to Rac-1-dependent mesenchymal cell migration (44). Intriguingly, control and parental HT29 cells exhibited high levels of serum-stimulated contractility, which was suppressed by acute or chronic silencing of Rap1GAP. Moreover, the migration of Rap1GAP-depleted, but not control or parental cells, was Rac1-dependent. Activation of Rap has been shown to stimulate Rac1 activity (12, 45) and to inhibit Rho activity (16, 46, 47). Rap1 and Rap2 activity are stably increased in Rap1GAP-depleted cells (32). Nonetheless, total Rac1 activity was not increased, and global levels of ROCK activity were not decreased by Rap1GAP depletion. Inhibition of Rac1 restored contractility, as evidenced by membrane blebbing, and enhanced MLC phosphorylation in Rap1GAP-depleted cells. Additionally, the tracks made by Rap1GAP-depleted cells reverted to a more circuitous pattern similar to that observed in control cells in the face of Rac1 inhibition. These results suggest that increases in endogenous Rap activity harbor the potential to alter migratory mechanisms, likely through the regulation of discrete pools of Rac1.

The alterations conferred by depleting Rap1GAP, impaired cell/cell adhesion, enhanced adhesion and spreading on collagen (32), and enhanced mesenchymal motility are similar to those observed in invasive cancer cells. Depletion of Rap1GAP enhanced invasion through Matrigel-coated filters in response to serum. The differences in invasion were more striking in cells embedded in Matrigel. HT29 and LoVo cells formed highly compact colonies that retained cell/cell adhesion. By contrast, cells acutely or stably silenced for Rap1GAP invaded the matrix in all directions. Collectively, these data suggest that, at least in vitro, loss of Rap1GAP endows human tumor cells with more aggressive migratory and invasive properties. Consistent with this notion, overexpression of Rap1GAP in tumor cell lines impaired migration in vitro (28, 29, 31, 33–35) and metastasis in vivo (36, 37). Whether depletion of Rap1GAP is sufficient to enhance metastasis remains to be determined.

The precise mechanism through which silencing Rap1GAP confers Rac1-dependent cell motility remains to be elucidated. Activated Rap recruits RacGEFs to the membrane (12). A role for Rap1 in the membrane translocation of Rac1 has been reported (48). It is conceivable that Rap activates specific pools of Rac1 that suppress ROCK-mediated contractility.

In conclusion, alterations in the level of Rap1GAP expression and of endogenous Rap activity appear to be important determinants of migratory and invasive behavior in human tumor cells. Although it is well documented that the migration of tumor cells is highly plastic, the endogenous signals that dictate the selection of migratory mechanisms are poorly understood. Our data suggest a previously unknown role for Rap1GAP in the regulation of migratory mechanisms.

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