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Guanine nucleotide exchange factor Dock7 mediates HGF-induced glioblastoma cell invasion via Rac activation

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Background: Glioblastoma multiforme (GBM), a highly invasive primary brain tumour, remains an incurable disease. Rho GTPases and their activators, guanine nucleotide exchange factors (GEFs), have central roles in GBM invasion. Anti-angiogenic therapies may stimulate GBM invasion via HGF/c-Met signalling. We aim to identify mediators of HGF-induced GBM invasion that may represent targets in a combination anti-angiogenic/anti-invasion therapeutic paradigm.

Methods: Guanine nucleotide exchange factor expression was measured by microarray analysis and western blotting. Specific depletion of proteins was accomplished using siRNA. Cell invasion was determined using matrigel and brain slice assays. Cell proliferation and survival were monitored using sulforhodamine B and colony formation assays. Guanine nucleotide exchange factor and GTPase activities were determined using specific affinity precipitation assays.

Results: We found that expression of Dock7, a GEF, is elevated in human GBM tissue in comparison with non-neoplastic brain. We showed that Dock7 mediates serum- and HGF-induced glioblastoma cell invasion. We also showed that Dock7 co-immunoprecipitates with c-Met and that this interaction is enhanced upon HGF stimulation in a manner that is dependent on the adaptor protein Gab1. Dock7 and Gab1 also co-immunoprecipitate in an HGF-dependent manner. Furthermore, Gab1 is required for HGF-induced Dock7 and Rac1 activation and glioblastoma cell invasion.

Conclusions: Dock7 mediates HGF-induced GBM invasion. Targeting Dock7 in GBM may inhibit c-MET-mediated invasion in tumours treated with anti-angiogenic regimens.

Glioblastoma multiforme (GBM) is the most common and deadly adult brain malignancy (Kohler *et al*, 2011). It is an incurable disease with an average life expectancy of 14 months following diagnosis (Van Meir *et al*, 2010). The morbidity of GBM is primarily due to its rapid growth, angiogenesis and invasion throughout the brain (Furnari *et al*, 2007; Ricard *et al*, 2012).

This invasive behaviour precludes total resection of the tumour, leading to rapid recurrence and currently, no anti-invasion therapeutic strategies exist. Moreover, the recently approved anti-VEGF therapy using bevacizumab has paradoxically been shown to stimulate the invasive behaviour of GBM (Lucio-Eterovic *et al*, 2009; Keunen *et al*, 2011). It is now believed that this occurs,

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at least in part, *via* activation of signalling downstream of the c-Met receptor tyrosine kinase (Lu *et al*, 2012).

Rho GTPases are key mediators of glioblastoma cell invasion (Fortin *et al*, 2012; Kwiatkowska *et al*, 2012; Kwiatkowska and Symons, 2013). These proteins function as molecular switches, cycling between the inactive guanine diphosphate (GDP)-bound and active guanine triphosphate (GTP)-bound states. The exchange of GDP for GTP and thus the activation of Rho GTPases are mediated by guanine nucleotide exchange factors (GEFs) (Bustelo *et al*, 2007). The activation of GEFs, in turn, often occurs downstream of receptor signalling (Schmidt and Hall, 2002). Once activated, Rho GTPases control cellular processes such as vesicular trafficking, cell polarity, cytoskeletal regulation, and gene transcription (Heasman and Ridley, 2008). Rho GEFs and Rho GTPases have a central role in cell migration and invasion, largely by regulating the actin cytoskeleton and the expression of matrix-degrading proteinases (Sahai and Marshall, 2002; Burridge and Wennerberg, 2004). The Rho GTPase signalling pathway is emerging as a promising target for anti-invasion therapy (van der Meel *et al*, 2011).

HGF, also known as a scatter factor, signals through receptor activation of c-Met that has been shown to mediate cell migration and growth in embryogenesis; and growth, invasion and metastasis in cancer cells (Birchmeier *et al*, 2003). HGF is the most potent chemotactic factor known for glioblastoma cells (Brockmann *et al*, 2003) and interestingly, c-Met, the receptor for HGF localises to invasive pseudopodia of motile glioma cells (Beckner *et al*, 2005). Moreover, the expression levels of HGF and its receptor c-Met correlate with astrocytoma grade (Rosen *et al*, 1996; Koochekpour *et al*, 1997; Lamszus *et al*, 1999) and importantly, blocking c-Met signalling, either using ribozymes that target HGF or c-Met, or a c-Met-targeting small molecule, inhibits glioblastoma tumour growth *in vivo* (Abounader *et al*, 1999), highlighting the therapeutic relevance of c-Met signalling for glioblastoma. Interestingly, c-Met-mediated signalling has also been shown recently to be important for the maintenance of tumour-initiating stem-like cells in glioblastoma tumours (Rath *et al*, 2013).

Following c-Met activation, an adaptor protein, Gab1, binds to cytoplasmic tyrosine docking sites on c-Met (Watanabe *et al*, 2006). A requirement for Gab1 has been shown recently in promitility signalling downstream of c-Met (Chaudhuri *et al*, 2011). HGF also activates the Rho family GTPases Rac1 and Cdc42 (Royal *et al*, 2000; Bosse *et al*, 2007; Palamidessi *et al*, 2008). The signalling mechanisms that mediate the activation of these GTPases by HGF/c-Met remain to be fully elucidated however (Gherardi *et al*, 2012).

In this study, we aimed to identify GEFs with enhanced expression in glioblastoma and importance in the context of c-Met-mediated GBM cell invasion. We envisage that identification of the mechanistic role of the key players in GBM cell invasion will lead to a targeted therapeutic approach for the clinical management of GBM especially in combination with an anti-angiogenic (bevacizumab) regimen.

MATERIALS AND METHODS

Dock7 expression profiling in non-neoplastic brain and human gliomas. Dock7 gene expression was mined in the publicly available NCBI Gene Expression Omnibus data set GSE4290 containing 195 clinically annotated brain tumour specimens. Expression values were filtered and principal component (PC) analysis to investigate the relationship between samples was performed as previously described (Tran *et al*, 2006). Box plots for Dock7 expression in the respective samples and survival clusters derived from PC analysis were graphed and the

significance between the respective populations was tested with a two-sample *t*-test assuming unequal variances as previously described (Tran *et al*, 2006).

Human tissue specimens for protein analysis. Fresh human astrocytoma (Grades I–IV) and non-malignant brain tissue specimens obtained from epilepsy reduction surgery were collected under a North Shore University Hospital's Institutional Review Board-approved protocol. Tissue specimens were obtained and immediately frozen in liquid nitrogen. Samples were stored at -80°C until further use. Brain tumour tissues were lysed in a buffer containing 50 mM Tris-HCl (pH 7.5), 2 mM DTT, 1 mM EDTA, 150 mM NaCl, 1% NP-40, supplemented with benzonase (Invitrogen, Grand Island, NY, USA), HALT protease inhibitor cocktail (Thermo, Waltham, MA, USA) and PhosSTOP phosphatase inhibitors (Roche, Madison, WI, USA), on ice with 10–15 strokes aided by a rotor homogeniser. Lysates were decanted by centrifugation at 12 000 g for 15 min at 4°C . Fifty micrograms of proteins was tested by western blotting.

Cell culture. The human glioblastoma cell line U87R-GFP (gift from Peter Forsyth, Moffitt Cancer Center, FL, USA) was cultured in Dulbecco's Modified Eagle's Medium (DMEM) F12 supplemented with 10% fetal bovine serum and $400\ \mu\text{g}\ \text{ml}^{-1}$ G418 as previously described (Johnston *et al*, 2007). SNB19 human glioblastoma cell line was obtained from the Cold Spring Harbor Laboratories Tissue Culture Facility (Cold Spring Harbor, NY, USA) and were maintained in DMEM supplemented with 10% fetal bovine serum. Cells were maintained at 37°C and 5% CO_2 .

siRNA transfections. siRNA duplexes for target genes were purchased from IDT (Coralville, IA, USA). In all, 3×10^5 cells were seeded in each well of a 6-well cell-culture plate in standard growth medium. After 5 h, $2\ \mu\text{l}$ dharmafect 1 (Dharmacon, Inc., Lafayette, CO, USA) in $200\ \mu\text{l}$ Opti-MEM (Gibco-Life Technologies Corp., Norwalk, CT, USA) was gently mixed with siRNA duplex (at desired nM concentration—see below) in $200\ \mu\text{l}$ Opti-MEM. After 20 min, media were removed from cells and this $400\ \mu\text{l}$ mixture was added dropwise to cells. In all, 1.6 ml antibiotic-free medium was added to each well to a final volume of 2 ml per well. The following day, medium was removed and fresh antibiotic-containing medium was added. On day 3, the cells were passaged from 2×6 wells to one 10 cm dish. The following siRNAs (sense strand provided) were used at 5 nM: Dock7 duplex #1 5'-GAUUAG GUUUGCUUCUGUAGCCAGGUU-3'; Dock7 duplex #2 5'-GGC AUGAAAUGGGUAGAUAAUCACA-3'; and Gab1 5'-CAGAUG UCUUGGAAUACUA-3'. A scrambled non-coding negative control was also used (Catalogue # DS NC1). (All siRNA primers were obtained from IDT).

Ex vivo brain slice invasion assay. The brain slice invasion assay was performed as described previously (Valster *et al*, 2005). Animals were maintained and brain slices obtained under a research protocol approved by the Institutional Animal Care and Use Committee of St Joseph's Hospital and Medical Center, Barrow Neurological Institute (Phoenix, AZ, USA). Briefly, 24 h after transfection, 1×10^5 GFP-labelled SNB19 or U87 cells were deposited in $0.5\ \mu\text{l}$ DMEM onto the bilateral putamen of $400\ \mu\text{m}$ thick slices of freshly isolated 4- to 6-week-old mouse brain. Serial Z-sections were collected using a LSM 5 Pascal laser-scanning confocal microscope (Zeiss, Thornwood, NY, USA) and the extent of invasion determined as the maximum depth of invasion of the glioblastoma cells.

3D cell invasion assay. Freshly plated cells were incubated in serum-free medium overnight. Basement membrane extract (Trevigen Inc., Gaithersburg, MD, USA) was thawed on ice at 4°C overnight. The bottom side of 24-well transwell inserts was coated with $1\ \text{mg}\ \text{ml}^{-1}$ fibronectin in PBS at 37°C for 1 h and

washed in sterile PBS and allowed to dry. Cells were trypsinised and counted. In all, 1×10^5 cells were resuspended in $50 \mu\text{l}$ of a mixture of BME diluted to $10 \mu\text{g ml}^{-1}$ with serum-free media and added carefully to the transwell insert. The BME, containing cells, was allowed to polymerise by incubation at 37°C for 30 min. In all, $200 \mu\text{l}$ of serum free medium was added to the upper chamber and $700 \mu\text{l}$ of media supplemented with 20% serum, or 20 ng ml^{-1} HGF (Sigma-Aldrich Corp., St Louis, MO, USA) was added to the lower well. Cells were allowed to invade for 24 h at 37°C and 5% CO_2 . Following this, chambers were placed in 4% (v/v) paraformaldehyde in PBS for 30 min at room temperature and crystal violet for 40 min at room temperature. Chambers were washed in distilled water and non-invading cells were removed from the insert with a cotton swab. Inserts were allowed to dry and were imaged using a Zeiss Axiovert 200M microscope (Zeiss) equipped with a $\times 2.5$ objective. Invading cells were counted using the ImageJ software (National Institutes of Health, Bethesda, MD, USA).

Sulforhodamine B cell proliferation assay. Five thousand cells were added per well of a 96-well plate in serum free, 10% serum or 20 ng ml^{-1} HGF medium in triplicate. Cells were cultured under standard conditions. The medium was changed every 2 days. Cells were fixed 4 h, 3 days and 5 days after seeding as follows: medium was removed, $100 \mu\text{l}$ 10% TCA in PBS was added to each well and left for 60 min at 4°C . Each well was washed five times with deionised water and allowed to dry overnight. In all, $100 \mu\text{l}$ of 0.2% sulforhodamine B (SRB) in 1% acetic acid was added to each well and the plate put on a plate shaker set to 300 r.p.m. at room temperature for 30 min. Each well was washed five times in 1% acetic acid and allowed to dry overnight. In all, $200 \mu\text{l}$ of 10 mM Tris was added to each well and the plate put on a shaker set to 300 r.p.m. at room temperature for 30 min. Absorbance was read at 490 nm.

Colony formation survival assay. Three hundred cells were plated in 6 cm dishes in triplicate in regular medium. One day after plating, ionising radiation was applied to the dishes using a Gammacell 1000 irradiator (^{137}Cs ; Atomic Energy of Canada, Ltd, Mississauga, ON, Canada). Medium was refreshed every 3 days. Ten days after seeding, cells were fixed by removing medium, washing with PBS and adding 4% paraformaldehyde in PBS at room temperature for 30 min. Cells were washed with PBS and stained with 0.2% SRB in 1% acetic acid with shaking for 30 min. Sulforhodamine B was removed and plates were washed in 1% acetic acid and allowed to dry. Plates were scanned and colonies were counted using the ImageJ software.

Immunoprecipitation. Following lysis, $400 \mu\text{g}$ total protein was incubated with $2 \mu\text{g}$ c-Met antibody (H-190) (Santa Cruz Biotechnology, Santa Cruz, CA, USA) or $5 \mu\text{g}$ Dock7 antibody (Abcam, Cambridge, MA, USA) adjusted to a final volume of 1 ml with lysis buffer overnight at 4°C with gentle rotation. In all, $40 \mu\text{l}$ 50% slurry of protein A beads was washed five times by centrifugation at 2500 g for 5 min and resuspending in $100 \mu\text{l}$ lysis buffer. Lysates were added to the beads and incubated by rotating for 4 h at 4°C . Beads were collected by centrifugation at 2500 g for 5 min, supernatant was discarded and beads were washed as above. Protein was eluted by incubation with $30 \mu\text{l}$ $2 \times$ BME-containing gel loading buffer at 90°C for 10 min and analysed by western immunoblotting as described below.

Western immunoblot analysis. Following treatment, cells were placed on ice, media were removed and washed with ice-cold PBS. PBS was removed and cells were lysed with RIPA buffer (Cell Signaling Technology, Inc., Danvers, MA, USA) containing HALT and PhosSTOP protease and phosphatase inhibitor cocktails. Total protein concentration was determined *via* Bradford assay and equal amounts of protein were separated by SDS-PAGE. Following transfer onto PVDF membrane, specific proteins were detected

using the antibodies below. Dock7, as previously described (Watabe-Uchida *et al*, 2006), c-Met (c-12; Santa Cruz), Gab1 (Cell Signaling Technology, Inc.), GST (GE Healthcare, Piscataway, NJ, USA). Membranes were developed by ECL. Films were scanned and bands quantified by densitometry using the ImageJ software.

Analysis of Dock7 activity. Following treatment, cells were lysed and activated Dock7 was immediately precipitated by incubation with freshly prepared GST-Rac1-G15A fusion protein, as recently described (Guilluy *et al*, 2011). Briefly, following incubation, and therefore binding of the fusion protein to active GEFs, the GST protein was pulled down using glutathione sepharose beads and proteins were eluted by boiling in gel loading buffer for 10 min at 90°C . Levels of activated Dock7 were then determined by western blotting.

GTPase activation assay. Following treatment, cells were lysed on ice and active GTP-bound Rac was measured in total cell lysates using the G-Lisa assay (Cytoskeleton Inc., Denver, CO, USA) following the manufacturer's instructions. Active Cdc42 levels were determined using the Active Cdc42 Pull-Down and Detection Kit (Pierce-Thermo Fisher Scientific, Waltham, MA, USA).

Quantification of lamellipodia. Cells plated on laminin-covered coverslips (Becton Dickinson, Franklin Lakes, NJ, USA) were serum starved for 16 h and were then incubated with serum-free DMEM (control) or stimulated for 4 min with serum-free DMEM containing 20 ng ml^{-1} HGF. Cells were fixed with 4% formaldehyde in PBS, permeabilised with 0.1% Triton X-100 in PBS and incubated with Rhodamine-conjugated phalloidin (Molecular Probes-Life Technologies Corp., Norwalk, CT, USA) to stain F-actin. Nuclei were stained with DAPI. Images were acquired using a Zeiss Axiovert 200M microscope. For each experimental condition, images were taken in a random manner. Lamellipodia were traced using the ImageJ software. For each cell, the fraction of the cell perimeter that displays lamellipodia was calculated.

Statistics. All analyses were performed in triplicate. Data are expressed as means with error bars representing standard error of the mean (s.e.m.). Data were analysed using Microsoft Excel. A Student's *t*-test was performed to assess statistical significance with $P < 0.05$ deemed to be significant.

RESULTS

Dock7 expression is increased in high-grade glioma. To discover Rho GEFs of clinical and functional importance in GBM, similar to previous studies (Salhia *et al*, 2008; Seol *et al*, 2009), we focused on GEFs that display enhanced expression in GBM tumour tissue *vs* non-neoplastic brain tissue and with expression that correlates with disease severity. Using a publicly available data set (NCBI, Gene Expression Omnibus GSE4290), we found that Dock7 mRNA levels are increased about two-fold in high-grade astrocytoma in comparison with non-neoplastic brain tissue (Figure 1A). We also examined Dock7 protein expression levels in lysates of tumour and non-neoplastic tissue from independent samples and confirmed increased expression in high-grade glioma tissue *vs* non-neoplastic brain tissue, with a trend towards intermediate expression levels in low-grade tissue (Figure 1B).

We also compared mRNA expression levels of Dock7 in GBM tumours from patients with short-term survival (median = 401 days) and long-term survival (median = 952 days), stratified from the GSE4290 data set as previously described (Tran *et al*, 2006), and found significantly higher Dock7 expression levels in tumours from patients with shorter survival (Figure 1C).

Dock7 expression is required for human glioblastoma tumour cell invasion. To examine the role of Dock7 in the malignant

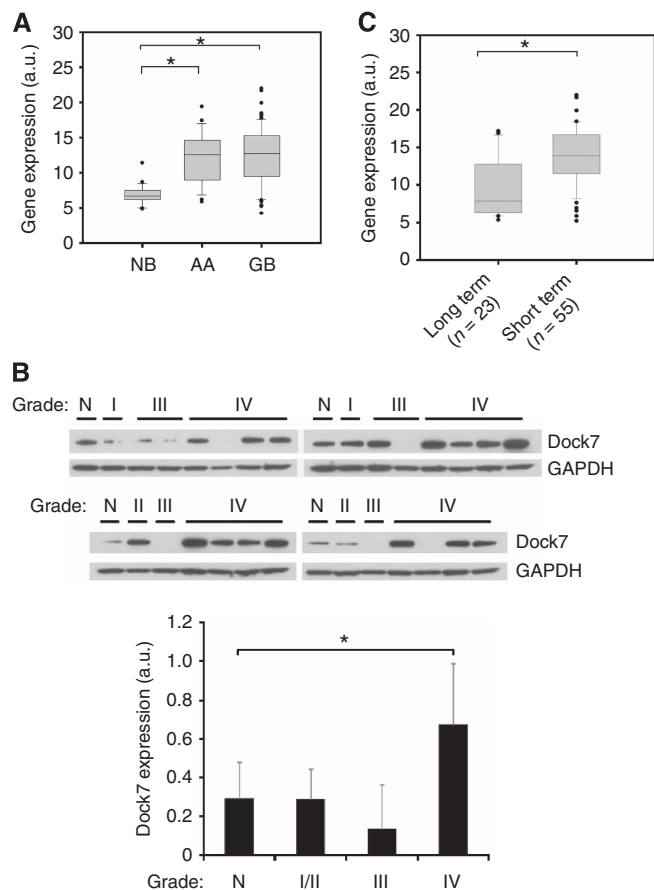


Figure 1. Dock7 expression is elevated in glioblastoma tumours. (A) Box and whisker plots of Dock7 mRNA expression levels from NCBI Gene Expression Omnibus GSE4290 for non-neoplastic brain (NB), anaplastic astrocytoma (AA) and glioblastoma (GB). (B) Western blot analysis showing Dock7 protein expression in non-neoplastic brain (N) and astrocytoma tissue (WHO Grades I-IV). Dock7 expression was normalised by GAPDH expression and was plotted (lower panel). (C) Principal component analysis of brain tumours from NCBI Gene Expression Omnibus GSE4290 relating Dock7 expression level in long-term (median = 952 days) and short-term (median = 401 days) survival groups. Significance between groups was tested with a two-sample t-test assuming unequal variances, * $P < 0.01$.

behaviour of glioblastoma, we first investigated the effect of Dock7 depletion on cell invasion. Using an organotypic mouse brain slice *ex vivo* invasion assay, we found that both SNB19 and U87 GBM cells, using two independent siRNA oligos to minimise the risk of RNA off-target effects, displayed strongly reduced cell invasion (Figure 2A and B).

As mentioned in the Introduction, HGF is the most potent chemoattractant known for glioblastoma cells (Brockmann *et al*, 2003) and we therefore also examined the effect of Dock7 depletion on HGF-stimulated glioblastoma cell invasion using a three-dimensional Matrigel invasion assay. Also in these conditions, depletion of Dock7 potentially inhibited glioblastoma cell invasion, both in U87R cells that have been selected for high invasiveness *in vivo* (Johnston *et al*, 2007) (Figure 2C) and in SNB19 cells (Figure 2D). Thus, these data support an important role for Dock7 in HGF-stimulated glioblastoma cell invasion.

Dock7 does not mediate GBM tumour cell proliferation, survival or radio-resistance. We next assessed whether Dock7 also has a role in other aspects of the malignant behaviour of glioblastoma cells. To determine the role of Dock7 in glioblastoma

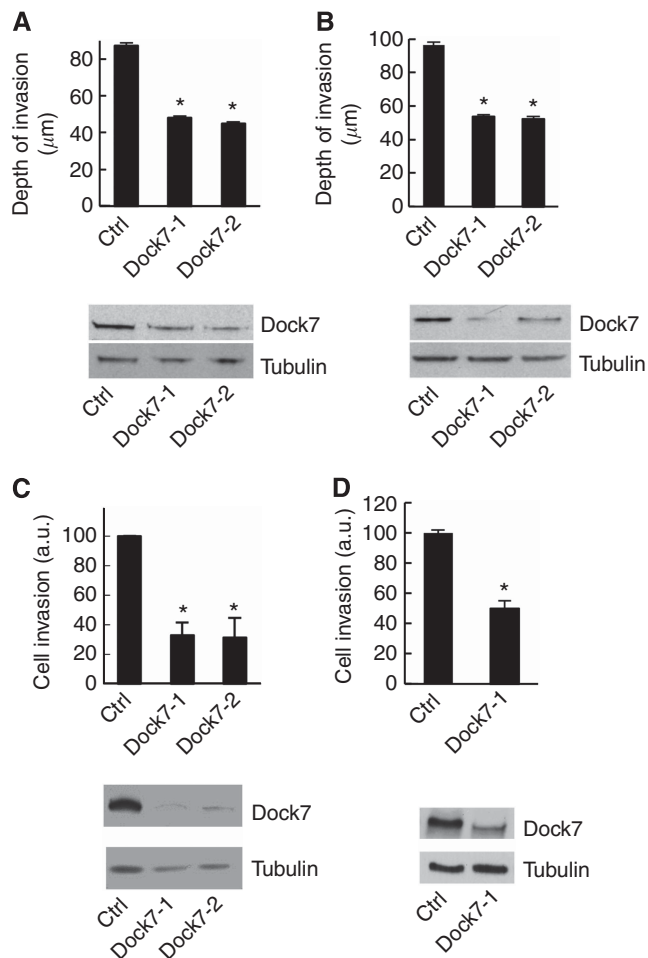


Figure 2. Dock7 depletion inhibits glioblastoma cell invasion. The effect of Dock7 knockdown using two siRNA duplexes (Dock7-1 and Dock7-2) on glioblastoma cell invasion into an *ex vivo* brain tissue slice using (A) U87 and (B) SNB19 cells. A scrambled non-coding negative control (Cat# DS NC1; IDT) was used as a control (ctrl). (C) The effect of Dock7 knockdown on U87 cell invasion towards 20 ng ml^{-1} HGF. (D) The effect of Dock7 knockdown (Dock7-1 siRNA) on SNB19 cell invasion towards HGF. Dock7 knockdown is displayed for each separate cell line under study. * $P < 0.05$, two-tailed t-test.

cell growth, we examined the effect of Dock7 depletion using the SRB colorimetric assay, both in serum-containing and serum-free medium and in serum-free medium supplemented with 20 ng ml^{-1} HGF. We found that depletion of Dock7 did not significantly alter U87R cell proliferation in any of these conditions (Figure 3A-C).

To test for a potential role of Dock7 in glioblastoma cell survival, we used a clonogenicity assay. Knockdown of Dock7 did not cause a significant decrease in colony formation in comparison with control cells however (Figure 3D). We also used this assay to determine whether Dock7 contributes to the resistance of glioblastoma cells to ionising radiation (IR), but we did not observe a significant increase in the sensitivity of the cells to radiation. Thus, taken together, these data indicate that Dock7 does not contribute to the proliferative and survival properties of glioblastoma cells.

Dock7 is required for HGF-induced Rac1 activation. Dock7 has been shown to possess GEF activity towards Rac1 and Cdc42 (Watabe-Uchida *et al*, 2006; Yamauchi *et al*, 2008; Zhou *et al*, 2013). As we found that Dock7 is required for HGF-stimulated

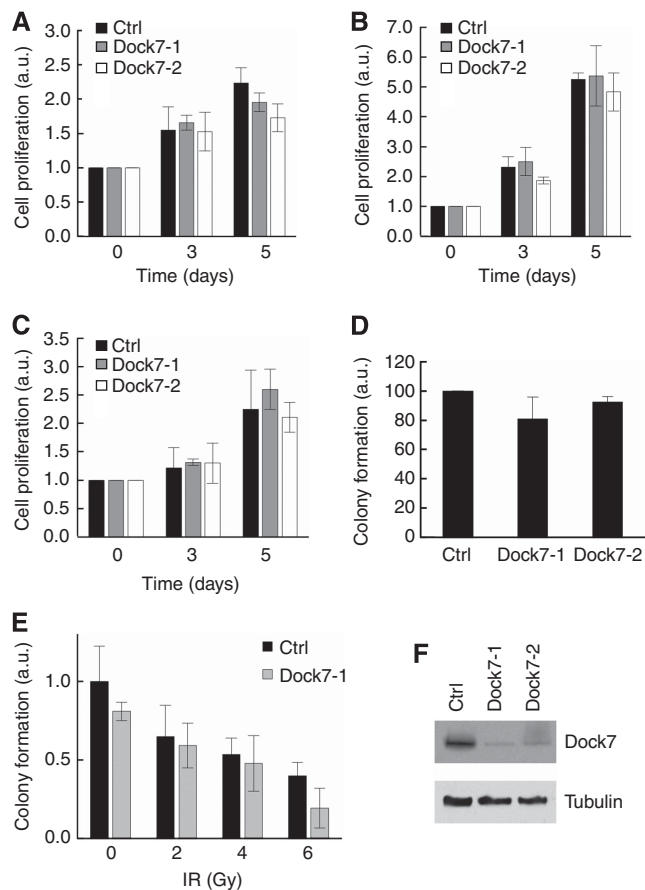


Figure 3. Dock7 is not required for glioblastoma cell proliferation, survival or radio-resistance. The effect of Dock7 knockdown using two siRNA duplexes (Dock7-1 and Dock7-2) on U87R cell proliferation as measured by sulforhodamine B assay in (A) serum-free conditions or (B) 10% serum or (C) 20 ng ml⁻¹ HGF over a 5-day period. Normalisation was performed by assigning a value of 1 to day zero readings and expressing all other readings relative to it. (D) The effect of Dock7 knockdown on SNB19 cell survival using a colony formation assay. Normalisation was performed by assigning the reading for control cells a value of 100% and expressing all other readings relative to it. (E) The effect of Dock7 knockdown on radiation-induced cell death. Normalisation was performed by assigning the reading for non-irradiated control cells a value of 1 and expressing all other readings relative to it. (F) Confirmation of Dock7 knockdown in U87R cells.

glioblastoma cell invasion, we next investigated whether Dock7 is necessary for the activation of these GTPases by HGF. Rac1 activation was quantified using an ELISA that measures the amount of Rac protein in cell lysates through binding to an immobilised Rac effector (Azim *et al*, 2000). Rac1 is the major Rac isoform expressed in glioma cells and therefore this assay essentially reports on the activation state of Rac1 (Chan *et al*, 2005). We found that Dock7 depletion strongly inhibits HGF-stimulated Rac1 activation in U87R cells, at all time points measured (Figure 4A).

To examine whether Dock7 acts on Cdc42 in U87R cells, we used a commercially available pull-down assay that measures activated Cdc42 (see Materials and Methods). However, we failed to detect any HGF-induced Cdc42 activation, even though in the same cells, both EGF and bradykinin resulted in robust Cdc42 activation (Figure 4B) (Kozma *et al*, 1995).

Rac1 is a key signalling element that controls the formation of lamellipodia, actin-rich protrusions at the leading edge of migrating cells (Ridley *et al*, 1992). We therefore also determined

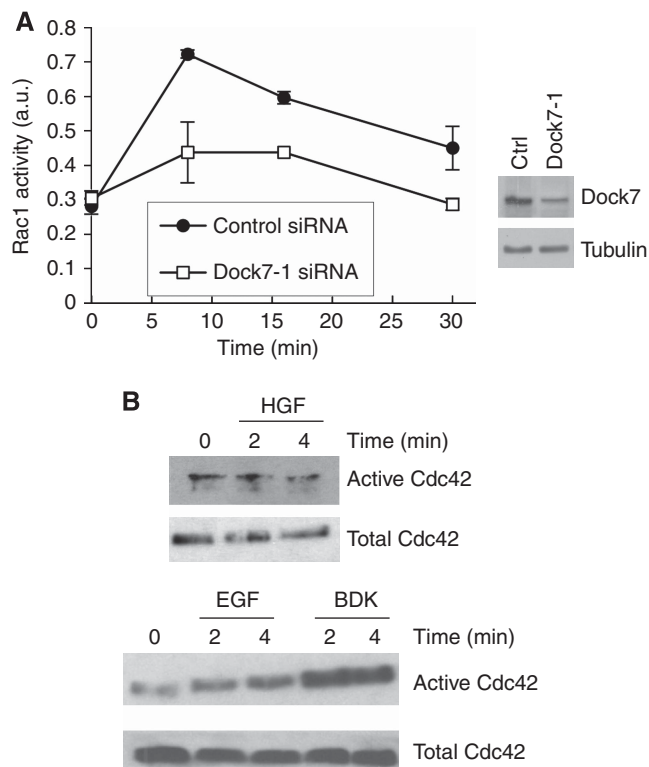


Figure 4. Dock7 is required for HGF-induced activation of Rac1 in human U87R cells. (A) Kinetics of the effect of Dock7 knockdown using the duplex Dock7-1 (right panel; confirmation of knockdown) on HGF-induced Rac1 activation. (B) HGF (20 ng ml⁻¹) does not activate Cdc42 in U87R cells. Both bradykinin (BDK, 1 μM) and EGF (50 ng ml⁻¹) activate Cdc42 in U87R cells.

whether Dock7 mediates HGF-induced lamellipodia formation. We observed that depletion of Dock7 significantly, albeit not completely, decreased HGF-induced lamellipodia formation (Figure 5A and B). HGF is also known to stimulate cell spreading (Matsumoto *et al*, 1994). We found that, in contrast to the partial inhibition of lamellipodia formation caused by Dock7 knockdown, depletion of Dock7 abolished HGF-induced cell spreading, measured 60 min after HGF treatment (Figure 5A and C), indicating a critical role for Dock7 in this function.

Dock7 binds HGF-activated c-Met via the Gab1 scaffold protein.

To further dissect the Dock7-mediated signalling mechanisms that control HGF-induced invasion, we first asked whether Dock7 interacts with c-Met. We readily detected Dock7 in c-Met, but not in control immunoprecipitates (Figure 6A). In addition, we found increased interaction of Dock7 with c-Met upon stimulation of the receptor with HGF (Figure 6A). We also examined whether Dock7 binds to c-Met via the Gab1 adaptor protein. Interestingly, although binding of Dock7 to c-Met was not affected by Gab1 depletion, the HGF-induced increase in binding of Dock7 to c-Met was abolished in Gab1-depleted cells, indicating that this adaptor protein mediates the increased interaction between c-Met and Dock7 upon stimulation with HGF (Figure 6B and C). To further investigate this, we examined the presence of Gab1 in Dock7 immunoprecipitates and found that Dock7 indeed interacts with Gab1 and that this interaction is increased upon HGF stimulation (Figure 6D). As a control, we performed the same analysis in Dock7 knockdown cells and observed decreased Gab1 in the corresponding Dock7 immunoprecipitate (Figure 6E). Thus, taken together, these data indicate that Dock7 interacts with c-Met and that this interaction is mediated, at least in part, by Gab1.

Gab1 adaptor protein is required for HGF-dependent GBM cell invasion, Rac1 activation and Dock7 activation. To further investigate the role of Gab1 as a mediator of c-Met in the control

of glioblastoma cell invasion, we examined the effect of Gab1 depletion on both HGF-induced invasion and Rac1 activation.

Depletion of Gab1 strongly inhibited HGF-induced U87R cell invasion (Figure 7A) and abolished HGF-induced Rac1 activation (Figure 7B). Interestingly, depletion of Gab1 caused an increase in basal Rac1 activity, suggesting the existence of a compensatory mechanism that regulates Rac1 activation.

As Dock7 mediates HGF-stimulated glioblastoma cell invasion, we also investigated whether Dock7 is activated by HGF and whether Gab1 mediates this activation. Using the GST-Rac1-G15A nucleotide binding-deficient mutant to pull down active Dock7 out of cell lysates (Guilluy *et al*, 2011), we observed robust stimulation of Dock7 activity by HGF that is abolished by depletion of Gab1 (Figure 7C and D). In addition, in line with our observation that depletion of Gab1 causes an increase in basal Rac1 activity, we also observed that depleting Gab1 enhances Dock7 activation.

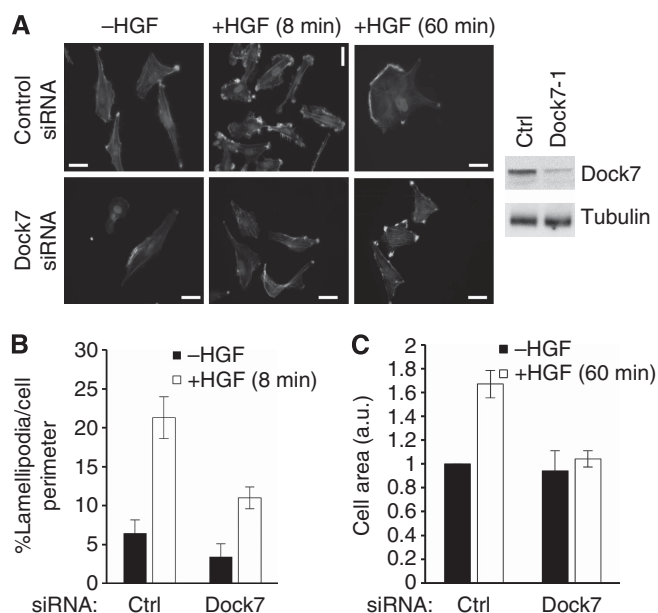


Figure 5. Dock7 is required for HGF-induced lamellipodia formation in SNB19 cells. (A) Micrographs of control and Dock7 knockdown cells (using the duplex Dock7-1) stimulated with HGF for 8 or 60 min and stained with fluorescently labelled phalloidin (right panel; confirmation of knockdown). (B) Quantification of lamellipodia formation. (C) Quantification of cell area. Bars, 10 μ m.

DISCUSSION

In this study, we showed that Dock7 expression levels are increased in glioblastoma vs non-neoplastic brain. In addition, we identified an important role for Dock7 in the invasive behaviour of glioblastoma cells, by mediating HGF-induced Rac activation.

In a previous study, we have provided evidence that Rac1 is constitutively activated in a significant fraction of human GBM tumours (Salhia *et al*, 2008). Our observations add Dock7 to a growing list of GEFs, including Trio, Vav3, SWAP-70, Ect2, and SGEF that have been shown to be overexpressed in GBM tumours and promote Rac1 activation (Salhia *et al*, 2008; Seol *et al*, 2009; Fortin Ensign *et al*, 2013). Dock7, Trio, Vav3 and SWAP-70

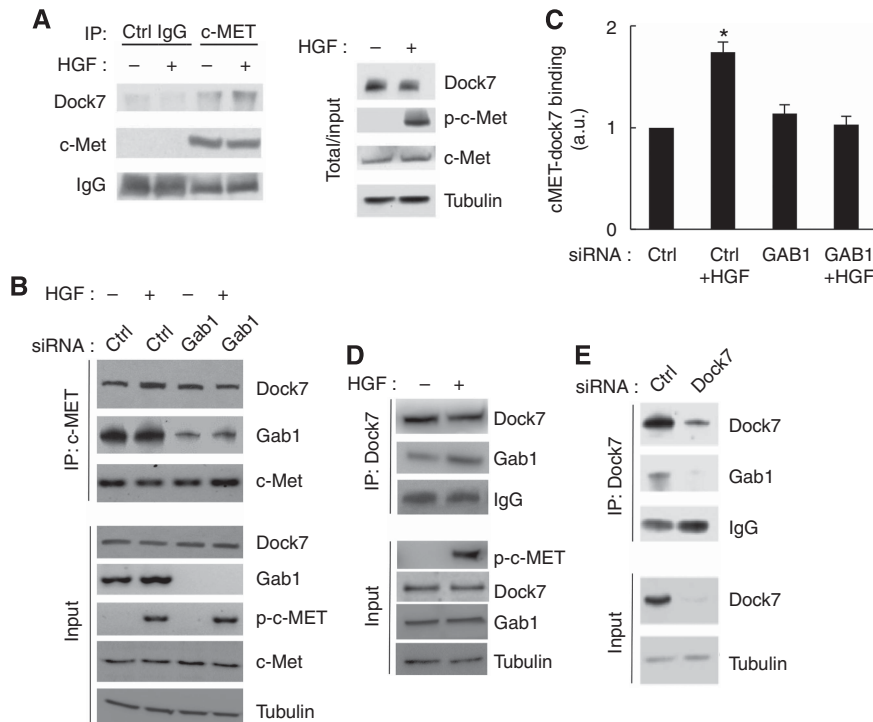


Figure 6. Dock7 binds activated c-Met in a Gab1-dependent manner. (A) Confirmation of c-Met immunoprecipitation specificity using a control rabbit IgG and subsequent immunoblot for Dock7 and c-Met. (B) Western blot demonstrating immunoprecipitation of Dock7 with c-Met from U87R cells. Experiments were performed 3 days after siRNA transfection. (C) Quantification of immunoprecipitation data from five independent experiments, exemplified in (B). Using the ImageJ software, densitometry values for immunoprecipitated Dock7 were normalised to the corresponding densitometry values for c-Met in the immunoprecipitate. (D) Western blot demonstrating immunoprecipitation of Dock7 with Gab1 from U87R cells. (E) Confirmation of Dock7 immunoprecipitation specificity using Dock7 siRNA to show specific pull down of Gab1 with Dock7. * $P < 0.05$, two-tailed t-test.

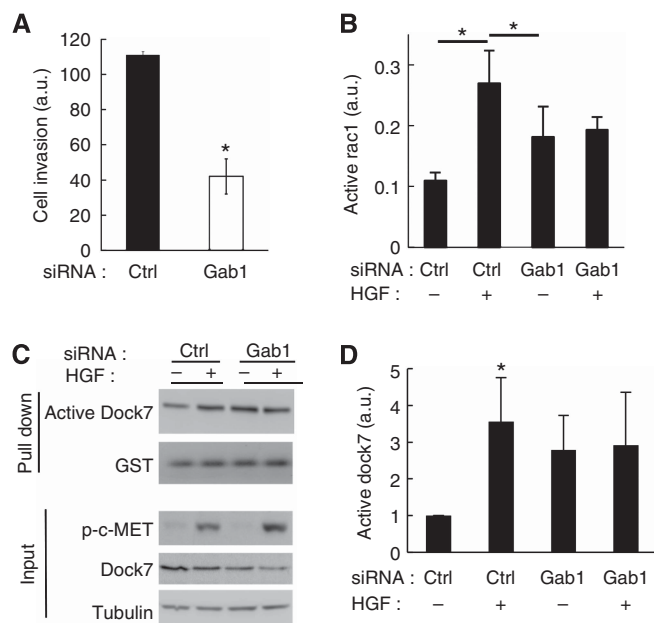


Figure 7. Gab1 mediates HGF-induced cell invasion and activation of Dock7 and Rac1. **(A)** Gab1 depletion reduces HGF-induced U87R cell invasion. Experiments were performed 3 days after siRNA transfection. **(B)** Gab1 depletion inhibits HGF-induced Rac1 activation measured by G-Lisa assay. Experiments were performed 3 days after siRNA transfection. **(C)** The role of Gab1 in HGF-induced Dock7 activation. Experiments were performed 3 days after transfection. Active Dock7 was pulled down using GST-Rac1-15A and subsequently measured by western blotting. **(D)** Quantification of Dock7 activation data from eight independent experiments, exemplified in **(C)**. Activated Dock7 was normalised to the amount of GST-Rac1-15A in the pull down. * $P < 0.05$, two-tailed t-test.

can directly activate Rac1, whereas Ect2 and SGEF are likely to stimulate Rac1 activity *via* Cdc42 and RhoG, respectively. Thus, overexpression of these GEFs, in addition to downregulation of RacGAPs, such as $\beta 2$ -chimaerin (Yuan *et al*, 1995), is likely to contribute to hyperactivation of Rac1 in GBM. Our observations that Dock7 mediates invasion-promoting signalling driven by HGF/*c*-Met, a signalling axis that is hyperactive in GBM, add further clinical significance to our findings.

Interestingly, whereas Rac1 has been shown to be required for glioblastoma cell proliferation and survival (Senger *et al*, 2002; Chan *et al*, 2005; Kwiatkowska *et al*, 2012), we found that depletion of Dock7, only had a marginal, if any, inhibitory effect on these functions. These observations suggest that Dock7, in addition to its catalytic activity, also may direct Rac1-controlled signals towards downstream pathways that are responsible for regulating cell migration and invasion. Such selective activation of downstream signalling could occur *via* interaction of GEFs with scaffold proteins (Marinissen and Gutkind, 2005; Garcia-Mata and Burridge, 2007).

Importantly, our molecular analysis showed that Dock7 interacts with *c*-Met in a manner that partially depends on the Gab1-adaptor protein. We found that Gab1 is necessary for HGF-stimulated binding of Dock7 to *c*-Met, suggesting that Dock7 is recruited to the receptor complex upon binding of Gab1 to the activated receptor. We also found that Dock7 activity is increased by *c*-Met ligation. The mechanistic basis for this increase in Dock7 activity remains to be elucidated. One possibility is that *c*-Met phosphorylates Dock7, thereby enhancing its catalytic activity, as has been shown to be the case for ErbB2 (Yamauchi *et al*, 2008). Another possibility is that binding of Dock7 to Gab1 leads to a

conformational change in Dock7, thereby activating it. Interestingly, in our study depletion of Gab1 causes an increase in the activities of both Dock7 and Rac1, independently of HGF stimulation. This suggests the existence of a compensatory mechanism that modulates the Dock7–Rac1 axis in the absence of Gab1. Future efforts will be directed towards exploring the mechanistic basis of this compensatory mechanism, focusing on the possible roles of other *c*-Met adaptor proteins including Crk and Grb2.

We further showed that Dock7 is necessary for HGF-induced Rac1 activation. HGF has been shown to activate Rac1 in a number of different cell systems (Palamidessi *et al*, 2008; Abella *et al*, 2010). Interestingly, the Rac-specific GEF Tiam1 has been shown to mediate HGF-induced Rac activation in HeLa cells, whereas here we document a role for Dock7 in HGF-induced Rac activation in glioblastoma cells. Whether *c*-Met signalling is relayed by different Rho GEFs in different cell systems is an important question to be addressed in future studies.

There is mounting evidence that HGF/*c*-Met-controlled signalling elements constitute relevant targets for therapeutic intervention in the management of glioblastoma (Steffan *et al*, 2011; Gherardi *et al*, 2012; Rath *et al*, 2013), underscoring the importance of improving our understanding of *c*-Met activated signalling. In principle, GEFs represent good targets for the management of GBM as they convey high signalling specificity, thereby limiting off-target effects of GEF inhibitors (Vigil *et al*, 2010). Moreover, the small molecule GEF inhibitor space is continuing to expand and a number of inhibitors have been successfully developed (Bouquier *et al*, 2009; Evelyn *et al*, 2009; Ferri *et al*, 2009; Shang *et al*, 2013). Emerging technologies, such as tumour-targeted delivery of gene-specific siRNAs, may provide additional therapeutic avenues.

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