

# Rac1 and a GTPase-activating protein, MgcRacGAP, are required for nuclear translocation of STAT transcription factors

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**S**TAT transcription factors are tyrosine phosphorylated upon cytokine stimulation and enter the nucleus to activate target genes. We show that Rac1 and a GTPase-activating protein, MgcRacGAP, bind directly to p-STAT5A and are required to promote its nuclear translocation. Using permeabilized cells, we find that nuclear translocation of purified p-STAT5A is dependent on

the addition of GTP-bound Rac1, MgcRacGAP, importin  $\alpha$ , and importin  $\beta$ . p-STAT3 also enters the nucleus via this transport machinery, and mutant STATs lacking the MgcRacGAP binding site do not enter the nucleus even after phosphorylation. We conclude that GTP-bound Rac1 and MgcRacGAP function as a nuclear transport chaperone for activated STATs.

## Introduction

The signal transducer and activator of transcription (STAT) family consists of seven members (STAT1–4, -5A, -5B, and -6). STATs are phosphorylated by cytokine stimulation, form homo- or heterodimers, and enter the nucleus, where they regulate expression of their target genes (Darnell, 1996; Ihle, 1996). Although STATs have a variety of functions under physiological conditions, the pathological importance of STAT functions has also been reported in many studies. STAT3 and -5 were activated in a broad spectrum of human hematological malignancies as well as in solid tumors (Darnell, 2002). A constitutively active form of STAT5 and -3 transformed IL-3–dependent Ba/F3 cells and fibroblasts, respectively (Onishi et al., 1998; Bromberg et al., 1999; Nosaka et al., 1999). An internal tandem duplication (ITD) mutant of receptor tyrosine kinase Flt3 (ITD-Flt3), a causative mutation of acute myeloid leukemia (Yokota et al., 1997; Hayakawa et al., 2000), induced phosphorylation of STAT5 on its tyrosine residues, thereby playing critical roles in

cell transformation (Mizuki et al., 2000; Zhang et al., 2000; Murata et al., 2003).

The mechanisms by which STATs are phosphorylated by cytokines and the activated STATs regulate the expression of the target genes have been well characterized. How activated STATs are transported to the nucleus has also been investigated; activated STAT1 and -3 were reported to bind importin  $\alpha$ 5 and several importin  $\alpha$ s, respectively, which mediated the nuclear transport of STATs (Sekimoto et al., 1997; McBride et al., 2002; Liu et al., 2005; Ushijima et al., 2005; Ma and Cao, 2006). However, molecules other than importins could also participate in the regulation of the nuclear translocation of STATs.

We have recently described the interactions among STAT3, Rac1, and a Rac/Cdc42 GTPase-activating protein (GAP), MgcRacGAP (male germ cell Rac-GAP), and have shown that MgcRacGAP is required for transcriptional activation of STAT3 (Tonozuka et al., 2004). However, the mechanisms by which Rac and MgcRacGAP regulate transcriptional activation of STAT3 remained unclear. In the present work, we investigated the molecular mechanisms of nuclear transport of a tyrosine-phosphorylated form of STAT5A, a close relative of STAT3, and found that GTP-bound Rac1 and MgcRacGAP were required for transport of activated STATs to the nucleus, indicating a novel function of Rac1 GTPase.

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Abbreviations used in this paper: DBD, DNA binding domain; EMSA, electrophoretic mobility shift analysis; GAP, GTPase-activating protein; ITD, internal tandem duplication; MBP, maltose binding protein; MgcRacGAP, male germ cell Rac-GAP; STAT, signal transducer and activator of transcription; TB, transport buffer.

The online version of this article contains supplemental material.

## Results

### STAT5A, Rac, and MgcRacGAP form a complex in hematopoietic cells

To test whether Rac1 and MgcRacGAP bind STAT5A, as was the case for STAT3 (Tonozuka et al., 2004), we did coimmunoprecipitation. STAT5A and MgcRacGAP were coimmunoprecipitated with Rac1 (Fig. 1 A) and Rac2 in Ba/F3 cells (unpublished data). In addition, STAT5A was coimmunoprecipitated with MgcRacGAP in Ba/F3 cells and in several other human and mouse cell lines, as well as in human primary T cells (unpublished data). These data show that Rac, STAT5A, and MgcRacGAP form a complex in vivo.

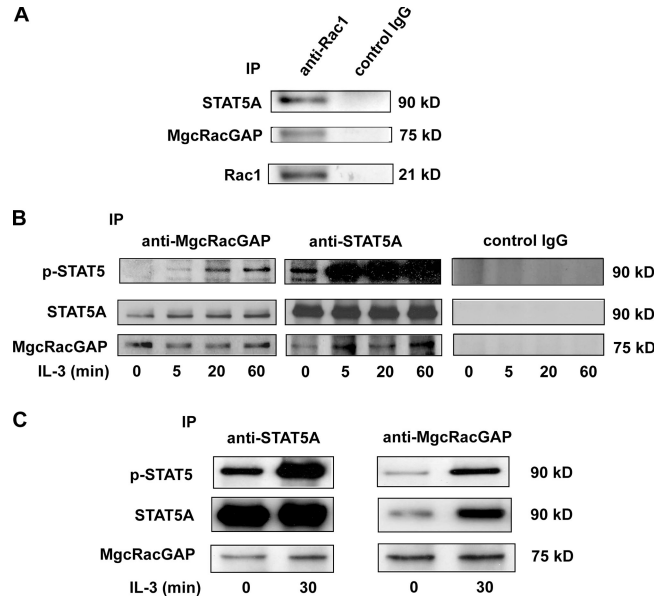
### Augmentation of MgcRacGAP association with STAT5A by IL-3 stimulation

A considerable amount of STAT5A protein was coimmunoprecipitated with MgcRacGAP in IL-3-starved Ba/F3 cells, and this association was enhanced by IL-3 stimulation (Fig. 1 B, left). Vice versa, a small amount of MgcRacGAP protein was coimmunoprecipitated with STAT5A in the starved cells, and this association was enhanced by IL-3 (Fig. 1 B, middle). In Ba/F3 cells expressing a constitutively active form of STAT5A (CA-STAT5A), which is more stable in the phosphorylated form than the wild-type STAT5A (Onishi et al., 1998), a considerable amount of STAT5A protein bound MgcRacGAP, even in unstimulated cells. This binding was also enhanced by IL-3 (Fig. 1 C). Thus, the association between MgcRacGAP and STAT5A does not require phosphorylation of STAT5A, but is enhanced by phosphorylation.

To map the interacting domains between MgcRacGAP and STAT5A, we prepared a series of truncated mutants of MgcRacGAP and STAT5A fused with maltose binding protein (MBP; Fig. S1, a, b, d, and e, available at <http://www.jcb.org/cgi/content/full/jcb.200604073/DC1>). It was found that STAT5A and Rac1 interacted with the Cys-rich and GAP domains of MgcRacGAP, whereas MgcRacGAP interacted with the DNA-binding domain (DBD) of STAT5A (Fig. S1, c and f). The binding domains between STAT5A and MgcRacGAP were similar to those between STAT3 and MgcRacGAP (Tonozuka et al., 2004).

### Simultaneous translocation of STAT5A and MgcRacGAP to the nucleus upon IL-3 stimulation

We next investigated the stoichiometry of STAT5A/MgcRacGAP binding in the cytoplasm or nucleus. IL-3-starved Ba/F3 cells were stimulated with IL-3 for 0, 15, or 90 min, and the cell lysates were fractionated. The cytosol and nuclear fractions were then immunodepleted with the anti-MgcRacGAP or anti-STAT5A antibody. The amounts of total STAT5A and tyrosine-phosphorylated STAT5A (p-STAT5A) in the nuclear fraction increased 15 min after IL-3 stimulation and decreased 90 min after IL-3 stimulation (Fig. 2 A, a–d, lanes for the control antibody). Notably, most of p-STAT5A in the cytosolic fractions was immunodepleted with the anti-MgcRacGAP antibody as well as with the anti-STAT5A antibody (Fig. 2 A, c). On the other hand, a considerable part of p-STAT5A was left in the nuclear extracts of IL-3-stimulated cells after the immunodepletion with the anti-MgcRacGAP antibody

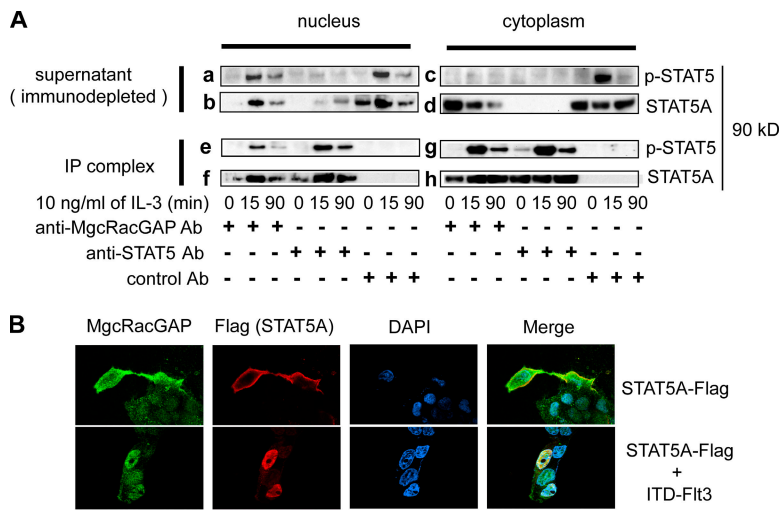


**Figure 1. MgcRacGAP, Rac1, and STAT5 formed a protein complex in IL-3-dependent Ba/F3 cells.** (A) STAT5A and MgcRacGAP were coprecipitated with Rac1. The cell lysates of IL-3-dependent Ba/F3 cells were subjected to immunoprecipitation with an anti-Rac1 or control antibody, followed by the immunoblotting with the anti-MgcRacGAP, anti-STAT5A, or anti-Rac1 antibody. (B) IL-3 enhanced association between STAT5A and MgcRacGAP. Ba/F3 cells were incubated in the presence or absence of 5 ng/ml IL-3 for the times indicated, and the cell lysates were subjected to immunoprecipitation with the anti-MgcRacGAP, anti-STAT5A, or control antibody, followed by the immunoblotting with the anti-p-STAT5 (top), anti-STAT5A (middle), or anti-MgcRacGAP antibody (bottom). Each row of images of the immunoprecipitation using the anti-MgcRacGAP and anti-STAT5A antibodies is derived from the same exposure of one gel, and each using the control antibody is derived from a similar exposure of the different gel. (C) The association of STAT5A and MgcRacGAP was enhanced in Ba/F3 cells expressing CA-STAT5A. Ba/F3 cells expressing CA-STAT5A were incubated in the presence or absence of 5 ng/ml IL-3 for 30 min, and cell lysates were subjected to immunoprecipitation with the anti-STAT5A (left) or anti-MgcRacGAP antibody (right), followed by the immunoblotting with the anti-p-STAT5 (top), anti-STAT5A (middle), or anti-MgcRacGAP antibody (bottom). Each row of images is derived from the same exposure of one gel.

(Fig. 2 A, a). These results suggested that most of p-STAT5A was bound by MgcRacGAP in the cytoplasm of IL-3-stimulated cells and was released from MgcRacGAP in the nucleus.

The amount of cytoplasmic STAT5A immunoprecipitated with the anti-MgcRacGAP antibody gradually increased after IL-3 stimulation (Fig. 2 A, h), and concomitantly the amount of cytoplasmic STAT5A immunodepleted with the anti-MgcRacGAP antibody gradually decreased (Fig. 2 A, d), implicating that MgcRacGAP maintained interaction with STAT5A in the cytoplasm of IL-3-stimulated cells even after the dephosphorylation of STAT5A. The fractionation was confirmed by Western blotting with the anti-HDAC (for nuclear fraction) or RhoA (for cytosol fraction) antibody (unpublished data).

Next, we visualized STAT5A and MgcRacGAP by immunostaining using adherent 293T cells. To enhance phosphorylation and nuclear translocation of STAT5, we used a constitutively active tyrosine kinase receptor, ITD-Flt3 (Yokota et al., 1997). In the absence of ITD-Flt3, ectopically expressed STAT5A-Flag localized to the cytoplasm and colocalized in part with the endogenous MgcRacGAP. Expression of ITD-Flt3 led to translocation



**Figure 2. STAT5A and MgcRacGAP simultaneously entered the nucleus.** (A) Stoichiometry of the association between MgcRacGAP and p-STAT5A or total STAT5A in the cytoplasm and nucleus. IL-3-starved Ba/F3 cells were stimulated with IL-3 for the times indicated, and cytosol and nuclear extracts were prepared as described previously (Nakamura et al., 2002). 10  $\mu$ g of protein for each of the extracts was immunodepleted with the anti-MgcRacGAP, anti-STAT5A, or control antibody, followed by immunoblotting with the anti-p-STAT5 and anti-STAT5A antibodies (a–d). The immunoprecipitates were also examined by Western blotting with the anti-p-STAT5 and anti-STAT5A antibodies (e–h). (B) STAT5A and MgcRacGAP translocated into the nucleus upon ITD-Flt3 stimulation in 293T cells. Cells were transfected with pME/STAT5A-Flag together with pMKIT (MOCK; top) or pMKIT/ITD-Flt3 (bottom). After 36 h, cells were immunostained with the anti-MgcRacGAP and anti-Flag antibodies and viewed using a Fluoview FV300 confocal microscope. Bar, 10  $\mu$ m.

and colocalization of STAT5A and MgcRacGAP in the nucleus (Fig. 2 B). These results indicated that MgcRacGAP translocated to the nucleus concurrently with STAT5A in response to IL-3 and ITD-Flt3 stimulation. Intriguingly, a dominant-negative form of Rac1, N17Rac1, completely inhibited the ITD-Flt3-induced nuclear translocation of STAT5A (Fig. S2, available at <http://www.jcb.org/cgi/content/full/jcb.200604073/DC1>). This result suggested that the GTP-bound form of Rac1 was required for the nuclear accumulation of activated STAT5A. However, N17Rac1 was recently reported to inhibit not only Rac1 but also other Rho-GTPases (Debrececi et al., 2004). To confirm that the N17Rac1 inhibition of nuclear translocation of p-STAT5A was indeed due to the inhibition of Rac1, we used mouse embryonic fibroblasts derived from gene-targeted conditional Rac1-flox mice in the Rac2-null background (Gu et al., 2003).

#### Rac1 is required for the nuclear translocation of p-STAT5A in mouse embryonic fibroblast cells

ITD-Flt3 induced the nuclear localization of STAT5A-Flag in the presence of Rac1 (Fig. 3 A, a and b). However, when Rac1 was depleted by Cre recombinase in Rac2<sup>-/-</sup>Rac1<sup>flox/flox</sup> fibroblasts (Fig. 3 B), the nuclear translocation of STAT5A was severely impaired (Fig. 3 A, c and d), and even p-STAT5 mostly remained in the cytoplasm (Fig. 3 A, e and f). In addition, CA-STAT5A did not enter the nucleus in the absence of Rac1 (unpublished data). We performed a similar analysis using Rac1<sup>flox/flox</sup>Rac2<sup>wt/wt</sup> fibroblasts and obtained identical results. These results demonstrate that Rac1 plays an essential role in the nuclear translocation of p-STAT5A.

#### Rac1 and MgcRacGAP were required for the nuclear accumulation and transcriptional activation of STAT5A

We next used siRNA to knock down Rac1 or MgcRacGAP expression in Ba/F3 cells where STAT5 activation is required for cell growth. The siRNA treatment for Rac1 or MgcRacGAP resulted in severe growth retardation of Ba/F3 cells and caused apoptosis in some cells. The total cell number was only one tenth or one fifth 48 h after siRNA treatment for Rac1 or MgcRac-

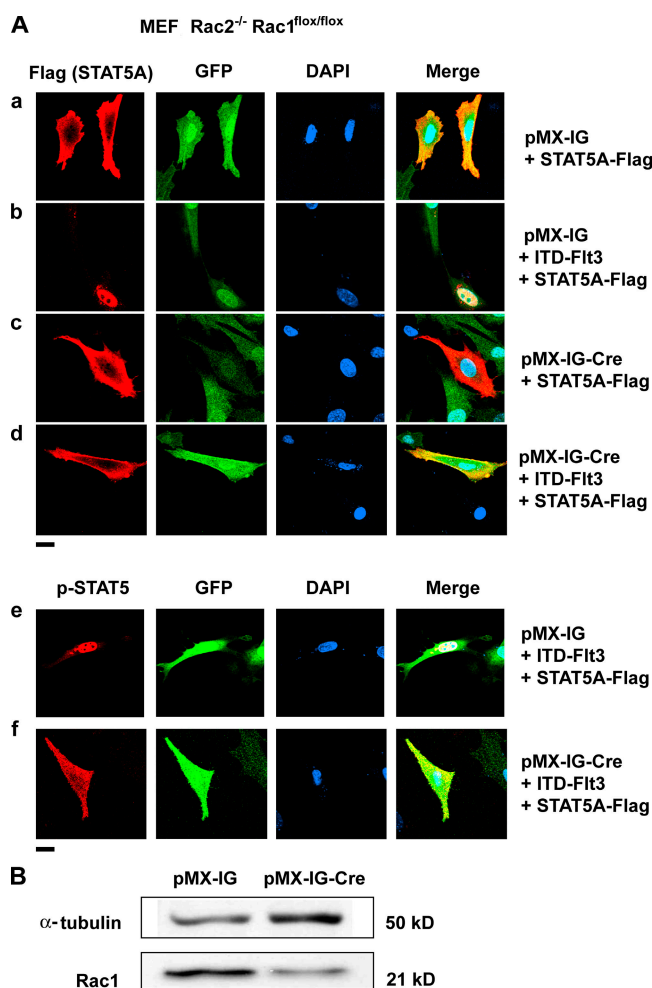
GAP, respectively (unpublished data). The siRNA treatment for MgcRacGAP led to the formation of multinucleated cells, as reported previously (Mishima et al., 2002), but no more than 20% of the cells, indicating the failure of cytokinesis by MgcRacGAP depletion is not the major cause of the growth inhibition.

We then did semiquantitative RT-PCR analysis to test whether transcriptional activation of STAT5 is affected by the knock down of Rac1 or MgcRacGAP and found that expression of bcl-xL, one of the STAT5 target genes, was severely impaired by the siRNA treatment (Fig. 4 A). We also confirmed that siRNA treatments specifically decreased the expression levels of Rac1 or MgcRacGAP protein but not those of RhoA and HDAC, similar to the results shown in Fig. 4 B (not depicted).

We also investigated whether knock down of Rac1 or MgcRacGAP affects the subcellular distribution of STAT5A and p-STAT5A in Ba/F3 cells before and after IL-3 stimulation. The siRNA-treated Ba/F3 cells were starved for 6 h after the isolation of live cells using Ficoll and stimulated with IL-3 (15 min), and the cell lysates were fractionated. The siRNA treatments specifically decreased expression levels of Rac1 or MgcRacGAP protein (Fig. 4 B, c and d) but not those of RhoA and HDAC (Fig. 4 B, e and f). The IL-3-induced nuclear accumulation of STAT5A and p-STAT5A was almost completely blocked in Ba/F3 cells treated with either Rac1 or MgcRacGAP siRNA when compared with those treated with the control siRNA (Fig. 4 B, a and b). The same treatment moderately decreased the amounts of p-STAT5A and total STAT5A in the cytoplasmic fraction (Fig. 4 B, a and b), suggesting that Rac1 and MgcRacGAP enhance the IL-3-induced phosphorylation of STAT5A and somehow stabilize STAT5A in the cytoplasm.

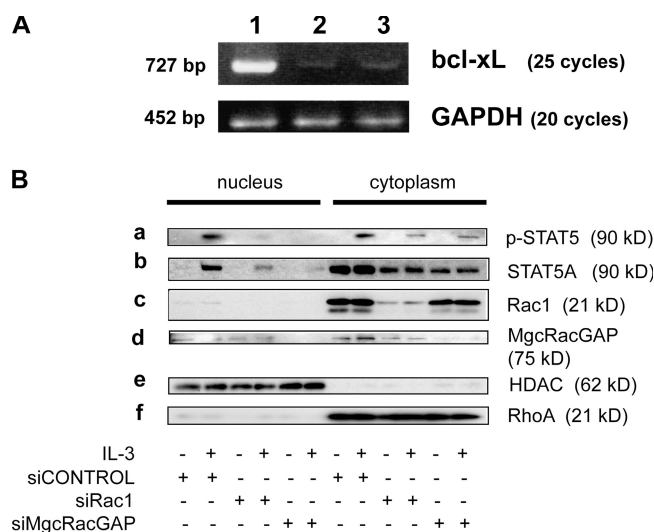
#### Direct interaction between STATs and MgcRacGAP/Rac1 regulates the activation of STATs through facilitating their tyrosine phosphorylation and nuclear translocation

We previously found that STAT3 bound MgcRacGAP through its DBD (Tonozuka et al., 2004). To examine whether MgcRacGAP regulated transcriptional activity of STAT3 and -5A through direct interaction, we attempted to produce mutant STATs lacking



**Figure 3. The nuclear translocation of p-STAT5 was not observed in the Rac1-knockout mouse embryonic fibroblasts.** (A) The nuclear translocation of ITD-Flt3-induced p-STAT5 was impaired in the Rac2<sup>-/-</sup>Rac1<sup>flox/flox</sup> fibroblasts. The Rac2<sup>-/-</sup>Rac1<sup>flox/flox</sup> fibroblasts were transfected with a control pMX-IG (a and b) or pMX-IG-Cre (c and d) retrovirus vector. After 3 d, cells were transiently cotransfected with pME/STAT5A-Flag and MOCK (a and c) or pMKIT/ITD-Flt3 (b and d). After 36 h, the cells were fixed and immunostained with the anti-Flag antibody (a–d, left) or anti-p-STAT5 antibody (e and f, left). Bars, 10  $\mu$ m. (B) Expression of Rac1 was depleted in the Rac2<sup>-/-</sup>Rac1<sup>flox/flox</sup> fibroblasts by Cre recombinase. Expression of  $\alpha$ -tubulin and Rac1 was examined in the Rac2<sup>-/-</sup>Rac1<sup>flox/flox</sup> fibroblasts retrovirally transfected with a control pMX-IG or pMX-IG-Cre.

the MgcRacGAP binding site. To this end, we narrowed down the binding site in DBD-STAT3 to a 25-amino-acid stretch, using MBP-fused DBD-STAT3 truncations (DB1-DB6; Fig. S3 A, available at <http://www.jcb.org/cgi/content/full/jcb.200604073/DC1>). We found that only DB2 (aa 338–362) of DBD-STAT3 interacted with MgcRacGAP (Fig. S3 B). Conversely, the mutant of DBD-STAT3 lacking DB2 (DBD-STAT3-dDB2) did not bind MgcRacGAP (Fig. S3 C). These results clearly demonstrated that the DB2 region (25 amino acid) of STAT3 bound MgcRacGAP. This region is well conserved among STAT family proteins and harbors a  $\beta$ -sheet structure, which is thought to mediate protein–protein interaction. Purified MgcRacGAP was pulled down by the MBP-DB2 of STAT3, and the corresponding region of STAT5 (aa 341–365) fused with MBP but not by MBP alone, demonstrating that MgcRacGAP directly bound DB2 of

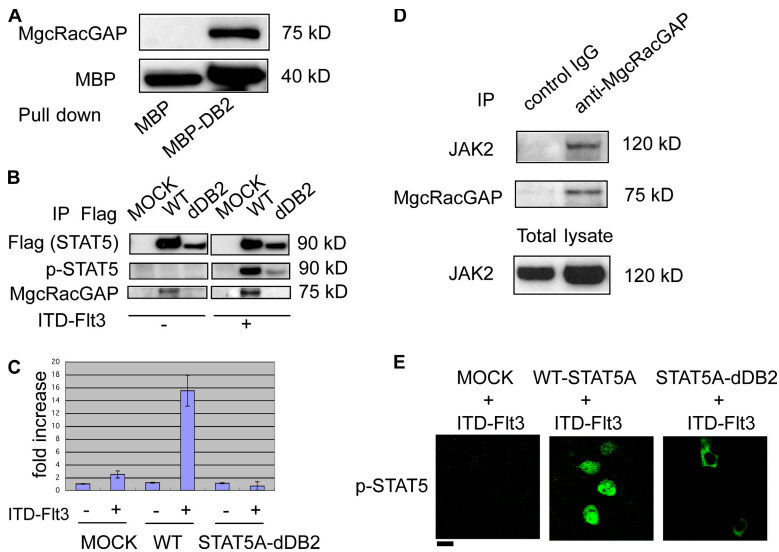


**Figure 4. Rac1 and MgcRacGAP were required for IL-3-induced nuclear accumulation and transcriptional activation of p-STAT5A.** (A) IL-3-induced transcriptional activation of STAT5A was suppressed by knock down of Rac1 or MgcRacGAP. Expression of bcl-xL or GAPDH mRNA was examined in Ba/F3 cells treated with the control siRNA (lane 1), Rac1 siRNA (lane 2), or MgcRacGAP siRNA (lane 3). 24 h after the siRNA treatment, the live cells were collected using Ficoll and subjected to semiquantitative RT-PCR. (B) IL-3-induced nuclear accumulation of p-STAT5A was impaired by knock down of Rac1 or MgcRacGAP. The intracellular distribution of p-STAT5A or total STAT5A in the IL-3-stimulated or unstimulated Ba/F3 cells pretreated with the control, Rac1, or MgcRacGAP siRNA (a and b). Note that Rac1 or MgcRacGAP expression was specifically suppressed by siRNA (c and d). Cytosol and nuclear extracts were prepared as described previously (Nakamura et al., 2002) and validated by Western blot using an anti-HDAC antibody or anti-RhoA antibody (e and f).

STAT3 and -5 (Fig. S3 D and Fig. 5 A). Both of the STAT3 and -5A mutants lacking DB2 (STAT3- and STAT5A-dDB2) did not bind MgcRacGAP and the extent of tyrosine phosphorylation of these mutants was less prominent after IL-6 or ITD-Flt3 stimulation (Fig. 5 B and not depicted). In addition, STAT3- and STAT5A-dDB2 lacked their transcriptional activities (Fig. S3 E and Fig. 5 C). These results indicated that the interaction of MgcRacGAP/Rac1 with STAT3 and -5A facilitates cytokine receptor-induced tyrosine phosphorylation of both STAT3 and -5A. Considerable decrease in the tyrosine phosphorylation of STAT5A was also observed when Rac1 or MgcRacGAP was knocked down (Fig. 4 B, a). Intriguingly, MgcRacGAP also interacted with JAK2 (Fig. 5 D), suggesting that MgcRacGAP/Rac1 also mediated the tyrosine phosphorylation of STATs through the interaction with JAK2. Importantly, STAT3- and STAT5A-dDB2 that do not bind MgcRacGAP did not enter the nucleus even after tyrosine phosphorylation by IL-6 or ITD-Flt3 (Fig. 5 E and not depicted), suggesting that MgcRacGAP/Rac1 is required not only for nuclear translocation of p-STATs but also for efficient tyrosine phosphorylation of STATs.

**MgcRacGAP and GTP-bound Rac1 were required for the nuclear translocation of p-STAT5A in cytosol-free digitonin-permeabilized cells**

We established a nuclear transport assay using semi-intact permeabilized cells (Adam et al., 1990), which enables us to



**Figure 5. The mutant of STAT5A, which lacks MgcRacGAP binding site, was not efficiently tyrosine phosphorylated by ITD-Fit3 stimulation and did not enter the nucleus even after tyrosine phosphorylation.** (A) The DB2 region of STAT5 directly interacted with MgcRacGAP in vitro. Full-length MgcRacGAP was expressed in Sf-9 cells using the baculovirus vector and was purified from infected Sf-9 cells. The recombinant MgcRacGAP was pulled down by MBP-DB2 or MBP-bound beads and subjected to Western blot analysis with the anti-MgcRacGAP (top) or anti-MBP antibody for the loading control (bottom). (B) The deletion mutant of DB2 did not bind MgcRacGAP, and the STAT5 phosphorylation was considerably impaired by the deletion of DB2. Expression and tyrosine phosphorylation of Flag-tagged STAT5A-dDB2 (top and middle, respectively) were examined in the MOCK or ITD-Fit3-transfected 293T cells. The interactions of MgcRacGAP with the WT-STAT5A or STAT5A-dDB2 were also examined in the MOCK or ITD-Fit3-transfected 293T cells (bottom). Images of the immunoblots using the MOCK or ITD-Fit3-transfected cells are derived from the same exposure of one gel that was cut to remove intervening lanes. (C) The transcriptional activity of STAT5-dDB2 was impaired. Luciferase activities were examined in the lysates of ITD-Fit3-stimulated 293T cells cotransfected with the STAT5-reporter plasmid together with internal control reporter plasmids

and the MOCK vector (pME), the expression vector for the Flag-tagged WT-STAT5, or STAT5-dDB2 mutant. The results shown are the mean  $\pm$  SD of three independent experiments. (D) MgcRacGAP was coprecipitated with JAK2. The cell lysates of 293T cells transfected with the expression vector (pRK5) for JAK2 were subjected to immunoprecipitation with the anti-MgcRacGAP or control antibody, followed by the immunoblotting with the anti-JAK2 (top) or anti-MgcRacGAP antibody (middle). Levels of transfected JAK2 were assayed by blotting with the anti-JAK2 antibody (bottom). (E) STAT5A-dDB2 did not enter the nucleus even after the phosphorylation. The 293T cells were cotransfected with pMKIT/ITD-Fit3 together with the MOCK (left), the expression vector for the Flag-tagged WT-STAT5A (middle), or STAT5A-dDB2 (right). After 24 h, the cells were fixed and immunostained with the anti-p-STAT5 antibody. Bar, 10  $\mu$ m.

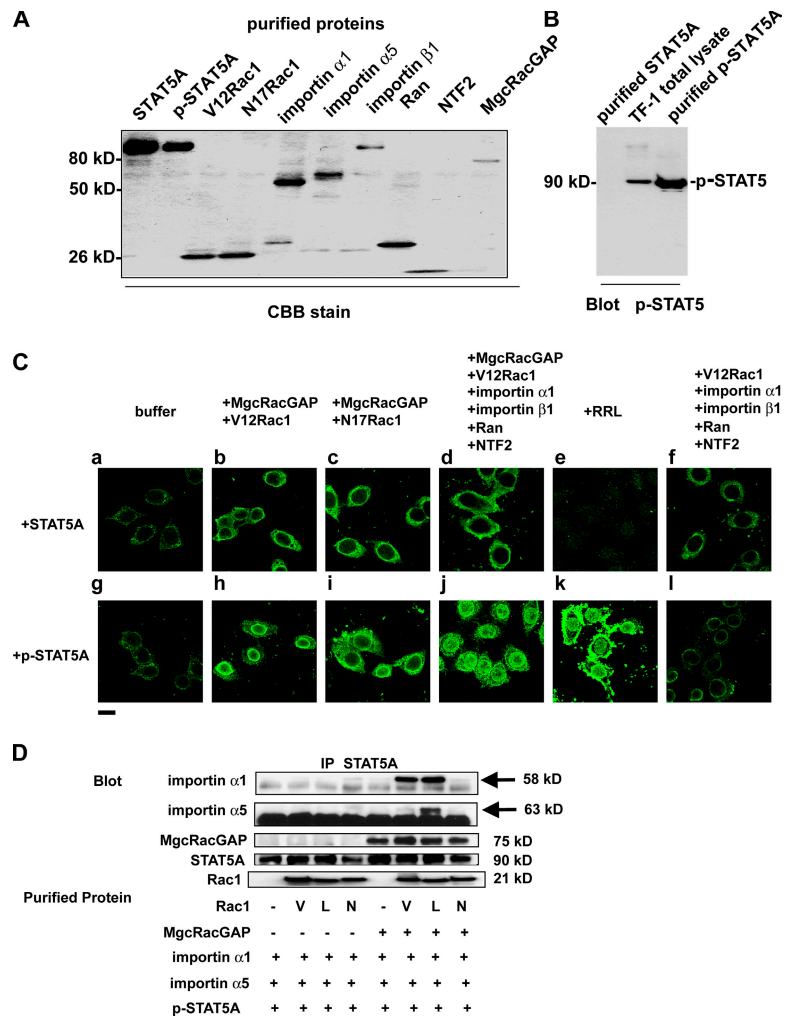
biochemically analyze the roles of Rac1 and MgcRacGAP in the nuclear import of p-STAT5A. We confirmed the purities of STAT5A, MgcRacGAP, V12Rac1, N17Rac1, importin  $\alpha$ 1, importin  $\alpha$ 5, importin  $\beta$ 1, Ran, and NTF2 produced by Sf-9 cells, and the tyrosine phosphorylation of STAT5A induced by coexpression with the kinase domain of JAK2 in Sf-9 cells (Fig. 6, A and B). It was confirmed that the purified p-STAT5A bound DNA in electrophoretic mobility shift analysis (EMSA) in a similar fashion with GM-CSF-activated STAT5 in TF-1 cells (Fig. S4 A, available at <http://www.jcb.org/cgi/content/full/jcb.200604073/DC1>), indicating that the recombinant in vivo phosphorylated STAT5A formed a proper dimer. Permeabilized HeLa cells were incubated with the indicated combinations of purified proteins in transport buffer (TB) plus an energy regenerating system. After the import reaction in the cells incubated with purified unphosphorylated STAT5A, a considerable amount of unphosphorylated STAT5A was detected at the cytoplasm in most cells (Fig. 6 C, a). The addition of purified MgcRacGAP, V12Rac1, importin  $\alpha$ 1, and importin  $\beta$ 1 did not affect localization of unphosphorylated STAT5A (Fig. 6 C, b–d and f). Although rabbit reticulocyte lysate reduced cytoplasmic localization of unphosphorylated STAT5A (Fig. 6 C, e), it induced both the nuclear and plasma membrane localization of p-STAT5A (Fig. 6 C, k). These results suggested that rabbit reticulocyte lysate contained cofactors that are required for the nuclear translocation of p-STAT5A in this transport assay. Interestingly, p-STAT5A accumulated at the nuclear membrane, with some migrating into the nucleus in the presence of purified MgcRacGAP and V12Rac1, but the nuclear translocation of p-STAT5A was inhibited in the presence of purified MgcRacGAP and N17Rac1 (Fig. 6 C, h and i). These results indicate that the GTP-bound form of Rac1 and MgcRacGAP facilitate

the nuclear translocation of p-STAT5A. Given that purified importin  $\beta$ 1 also accumulated mostly in the nuclear envelope and only partially migrated to the nucleus in our assay system (unpublished data) as reported previously (Kutay et al., 1997), the accumulation of p-STAT5 and importin  $\beta$ 1 in the nucleus might have been caused by residual amounts of nuclear transporters left in the assay system. Thus, it is likely that the GTP-bound form of Rac1 and MgcRacGAP play critical roles in targeting p-STAT5A to the nuclear envelope and that cofactors are required for the efficient nuclear import of p-STAT5A from the nuclear envelope. In fact, nuclear translocation of p-STAT5A was enhanced by further addition of the purified nuclear transporters, including importin  $\alpha$ 1, importin  $\beta$ 1, Ran, and NTF2 to the assay (Fig. 6 C, j). This nuclear translocation of p-STAT5A was not observed in the absence of MgcRacGAP even in the presence of cofactors (Fig. 6 C, l).

To confirm whether the unphosphorylated recombinant STAT5A conserved a native folded state, we did nuclear transport assay using the in vitro phosphorylated STAT5A. The recombinant full-length JAK2 efficiently phosphorylated the recombinant STAT5A in the kinase reaction buffer (Fig. S4 B). This in vitro phosphorylated STAT5A behaved in the nuclear transport assay like the in vivo phosphorylated STAT5A (Fig. S5, a–i, available at <http://www.jcb.org/cgi/content/full/jcb.200604073/DC1>). The nuclear transport of p-STAT5A requires both MgcRacGAP and V12Rac1. The nuclear import of the in vitro phosphorylated recombinant STAT5A was also achieved by the presence of the cytosol fraction of HeLa cells (HeLa-CS), which had been prepared as described previously (Adam et al., 1990). In addition, immunodepletion of MgcRacGAP or Rac1 considerably inhibited the nuclear import of the in vitro phosphorylated recombinant STAT5A (Fig. S5, j–m). This inhibition

Figure 6. Purified p-STAT5A accumulated to the nuclear envelope in the presence of V12Rac1 and MgcRacGAP in the nuclear transport assay.

(A) Coomassie blue (CBB) staining of purified STAT5A, p-STAT5A, V12Rac1, N17Rac1, importin  $\alpha$ 1, importin  $\alpha$ 5, importin  $\beta$ 1, Ran, NTF2, or MgcRacGAP. (B) Western blot analysis of the STAT5A-Flag protein purified from Sf-9 cells with or without coexpression with the kinase domain of JAK using the anti-p-STAT5 antibody. Total cell lysate of GM-CSF-stimulated TF-1 was used as a control. (C) The nuclear transport assay. HeLa cells were permeabilized with 40  $\mu$ g/ml digitonin. Incubation with 50  $\mu$ l import mix was done at 37°C for 30 min. Import mix contained TB, an energy regenerating system, and a single or combinations of the following purified proteins as indicated: 1  $\mu$ M STAT5A, p-STAT5A, V12Rac1, N17Rac1, MgcRacGAP, importin  $\alpha$ 1, importin  $\beta$ 1, Ran, or NTF2. After the import reaction, the cells were fixed. STAT5A protein was detected using the anti-STAT5A antibody. Cells were examined using a Fluoview FV300 confocal microscope. A representative result of three independent experiments is shown. Bar, 10  $\mu$ m. (D) The direct bindings of both GTP-bound Rac1 and MgcRacGAP facilitated the interaction of p-STAT5A with importin  $\alpha$ s. Purified p-STAT5A was incubated with importin  $\alpha$ s in the absence or presence of the indicated combinations of V12Rac1, L61Rac1, N17Rac1, or MgcRacGAP in TB containing 5% BSA to block nonspecific bindings. 1  $\mu$ g of each purified protein was used for each sample. After the incubation for 30 min at RT, STAT5A was immunoprecipitated with anti-STAT5A antibody and washed three times with TB. The immunoprecipitates were subjected to Western blot analysis with the anti-importin  $\alpha$ 1, anti-importin  $\alpha$ 5, anti-Rac1, anti-MgcRacGAP, or anti-STAT5A antibody.



was restored by add-back of the purified recombinant MgcRacGAP or Rac1 (Fig. S5, n and o).

To determine whether the Rac1 activation or the presence of MgcRacGAP is required for the interaction of p-STAT5A with importin  $\alpha$ s, an in vitro binding assay was done using purified proteins. Intriguingly, p-STAT5A formed complexes with importin  $\alpha$ 1 and  $\alpha$ 5 only in the presence of both MgcRacGAP and V12Rac1 or another constitutively active mutant L61Rac1, but not N17Rac1 (Fig. 6 D). These results demonstrated that GTP-bound Rac1 and MgcRacGAP functions as p-STAT5A nuclear chaperone, facilitating p-STAT5A to form protein complexes with importin  $\alpha$ s.

## Discussion

In the present work, we demonstrate that Rac1 and MgcRacGAP are essential for the nuclear translocation of STAT5A, based on the following observations. First, Rac1 and MgcRacGAP directly bound STAT5A, and the interaction between MgcRacGAP and STAT5A was enhanced by IL-3 stimulation. Second, STAT5A and MgcRacGAP simultaneously entered the nucleus upon IL-3 and ITD-Flt3 stimulation. Third, knock down of Rac1 or MgcRacGAP profoundly inhibited both the IL-3–

induced transcriptional activation of STAT5A and the nuclear accumulation of p-STAT5A in IL-3-dependent Ba/F3 cells. Fourth, depletion of Rac1 in fibroblasts, as well as expression of N17Rac1 in 293T cells, prevented p-STAT5A from entering the nucleus. Fifth, p-STAT5A lacking the MgcRacGAP binding site (p-STAT5A-dDB2) did not accumulate in the nucleus. Last, in a nuclear transport assay, purified V12Rac1 and MgcRacGAP induced accumulation of purified p-STAT5A on the nuclear envelope, with some p-STAT5A migrating into the nucleus, and the further addition of nuclear transporters, including importin  $\alpha$ 1, importin  $\beta$ 1, Ran, and NTF2, achieved the efficient nuclear translocation of p-STAT5A. Moreover, either the absence of MgcRacGAP or the presence of N17Rac1 inhibited this nuclear translocation of p-STAT5A.

Simon et al. (2000) suggested that an active form but not an inactive form of Rac1 bound STAT3 and played important roles in EGF-induced STAT3 activation. These authors did not, however, specifically examine the nuclear transport of STAT3. Interestingly, EGF receptor-mediated endocytosis is required for cytoplasmic transport of STAT3 (Bild et al., 2002), and MgcRacGAP is recruited to the EGF receptor complex after EGF stimulation (Blagoev et al., 2003). We also found that STAT3 bound Rac1 and Rac2, which was enhanced by IL-6 stimulation.

In addition, STAT3 bound MgcRacGAP, which was required for the transcriptional activation of STAT3, and some population of MgcRacGAP entered the nucleus together with STAT3 (Tonozuka et al., 2004). Although these results suggested a role of Rac1/MgcRacGAP in STAT3 activation, the underlying molecular mechanisms remained elusive. We studied the functional interactions using a nuclear transport assay and found that the nuclear translocation of p-STAT3 as well as p-STAT5A was induced in the presence of a combination of purified proteins, including V12Rac1, MgcRacGAP, importin  $\alpha$ 1, importin  $\beta$ 1, Ran, and NTF2 (Fig. 6, Fig. S5, and not depicted). These results demonstrate a novel Rac1 function in the nuclear transport of p-STAT3 as well as p-STAT5A.

Although we showed the results for STAT5A, we obtained identical results in experiments so far performed for closely related STAT5B (unpublished data). In addition, the phenotypes of STAT3- and STAT5A-dDB2 were nearly identical (Fig. 5 and Fig. S3), and the region of STAT3 that binds to MgcRacGAP (STAT3-DBD-DB2) is well conserved among STAT family proteins, suggesting a general role for MgcRacGAP and Rac1 in the nuclear transport of p-STAT proteins.

#### Involvement of Rac1 in the nuclear transport of STATs

The Rho family small GTPases play key roles in a variety of cellular functions, including regulation of cell cycle, transcription, and transformation (Bishop and Hall, 2000). Among them, the Rac subfamily consists of three known members: Rac1, Rac2, and Rac3. Although Rac1 and Rac3 are ubiquitously expressed, Rac2 expression is specific in hematopoietic cells. Rac1 and Rac2 were implicated in both distinct and overlapping functions, including cell migration, membrane ruffling, production of superoxide, and phagocytosis (Ridley, 1995; Roberts et al., 1999; Bishop and Hall, 2000; Williams et al., 2000; Gu et al., 2003; Cancelas et al., 2005). Interestingly, the C-terminal region of Rac1 but not Rac2 or Rac3 contained a functional NLS, suggesting a role for Rac1 in the nucleus. Consistent with this, Rac1 was reported to play a role in the nuclear import of SmgGDS and p120 catenin (Lanning et al., 2003), members of the importin  $\alpha$ -like armadillo family of proteins (Peifer et al., 1994; Chook and Blobel, 2001). In the present paper, using Rac1-deficient mouse embryonic fibroblasts, we demonstrate that Rac1 was critically required for the nuclear transport of p-STAT5A (Fig. 3).

#### Requirement of cofactors involved in importin $\alpha/\beta$ pathway for nuclear import of STATs

It was reported that unphosphorylated STATs shuttled between the cytoplasm and nucleus (Zeng et al., 2002; Marg et al., 2004). Activated STAT1 was reported to bind importin  $\alpha$ 5, leading to its nuclear translocation (Sekimoto et al., 1997; McBride et al., 2002). How activated STAT3 is imported to the nucleus has remained controversial. Ushijima et al. (2005) showed that activated STAT3 binds importin  $\alpha$ 1,  $\alpha$ 3, and  $\alpha$ 5, and Ma and Cao (2006) demonstrated that activated STAT3 binds importin  $\alpha$ 5 and  $\alpha$ 7 but not  $\alpha$ 1,  $\alpha$ 3, or  $\alpha$ 4, whereas Liu et al. (2005) reported that STAT3 nuclear import is independent of tyrosine phosphor-

ylation and mediated by importin  $\alpha$ 3. On the other hand, how activated STAT5A is imported to the nucleus remained largely elusive. It was reported that the ERBB4/HER4 receptor tyrosine kinase, which harbors the NLS sequence, functions as a STAT5A nuclear chaperone, implicating the NLS of STAT5A-associated molecules in the nuclear translocation of STAT5A (Williams et al., 2004). However, unlike ERBB4/HER4, ITD-Flt3 does not harbor an NLS and did not enter the nucleus (unpublished data). In the nuclear transport assay, most p-STAT5A accumulated to the nuclear envelope in the presence of V12Rac1 and MgcRacGAP, and further addition of the purified nuclear transporters, including importin  $\alpha$ 1, importin  $\beta$ 1, Ran, and NTF2, facilitated the nuclear translocation of p-STAT5A (Fig. 6 C, j). Together, it is likely that the complex of p-STAT5, GTP-bound Rac1, and MgcRacGAP translocates to the nuclear envelope, where it recruits other factors such as importin  $\alpha/\beta$  to pass through the nuclear pore complex into the nucleus. Indeed, direct interaction of both GTP-bound Rac1 and MgcRacGAP facilitated the interaction of p-STAT5A with importin  $\alpha$  (Fig. 6 D). In agreement with this, Rac1 harbors an NLS (Lanning et al., 2003) and MgcRacGAP harbors a bipartite NLS and binds importin  $\alpha$  (unpublished data). Interestingly, a mutant of MgcRacGAP lacking NLS strongly blocked the nuclear translocation of p-STATs in the nuclear transport assay even with V12Rac1, importin  $\alpha$ 1, importin  $\beta$ 1, Ran, and NTF2 (unpublished data), suggesting a role of MgcRacGAP as an NLS-containing nuclear chaperone of p-STATs. Establishment of the nuclear transport assay for p-STATs has enabled us to clearly demonstrate the requirement of Rac1 and MgcRacGAP for the nuclear translocation of p-STATs.

The activities of small GTPases are regulated by two classes of proteins, GAPs and GEFs (guanine nucleotide exchange factors). In this paper, we did not address GEFs, but some, such as smgGDS or ECT-2, may also participate in the nuclear transport of STAT proteins. Based on the results that p-STAT5A binds importin  $\alpha$ s only in the presence of MgcRacGAP and active forms of Rac1 but not inactive form of Rac1, we speculate that Rac1 inactivation by MgcRacGAP release p-STATs from the importin complex in the nucleus. To prove this hypothesis and clarify its molecular mechanisms, further work will be required.

#### Coordinate control of cell division and transcription

Another interesting question raised by our work concerns the coordinate control of cell division and transcription. We originally identified MgcRacGAP as a GAP protein that regulates IL-6-induced macrophage differentiation of leukemic M1 cells (Kawashima et al., 2000). Later, we and others found that MgcRacGAP or Cyk-4, an orthologue in *Caenorhabditis elegans*, played essential roles in cytokinesis (Jantsch-Plunger et al., 2000; Hirose et al., 2001; Van de Putte et al., 2001; Mishima et al., 2002). We further demonstrated that MgcRacGAP was phosphorylated at Serine 387 by Aurora-B at the midbody, functionally converted from Rac/Cdc42-GAP to Rho-GAP, and played essential roles to complete cell division in cytokinesis (Minoshima et al., 2003). In interphase, MgcRacGAP formed a

complex with Rac1 and STAT3 and was required for the full transcriptional activation of STAT3, thereby enhancing the differentiation of IL-6-stimulated M1 cells (Tonozuka et al., 2004). On the other hand, when STAT5 was activated by IL-3 or ITD-Flt3 in conjunction with Rac1 and MgcRacGAP, the cells proliferate. Thus, MgcRacGAP functions as a Rac-GAP to activate transcription of STAT in the nucleus of interphase cells, probably leading to cell proliferation or differentiation. At cytokinesis, it functions as a Rho-GAP to complete cytokinesis, indicating that the distinct roles of the Rho family small GTPases depend on the cell cycle.

### Does Rac1 play a general role in nuclear transport of transcription-related proteins?

The experiments using N17Rac1 showed that Rac1 contributes to maximal activation of STAT1 and -3 in response to IFN- $\gamma$  (Park et al., 2004). The molecular mechanisms of this phenomenon can be explained by our current results. Esufali and Bapat (2004) suggested that Rac1 plays some role in redistribution of  $\beta$ -catenin and that a mutant Rac1 lacking its NLS hampers nuclear localization of  $\beta$ -catenin, leading to attenuation of the  $\beta$ -catenin-dependent transcriptional activity of T cell factor/lymphoid enhancing factor. The authors stated that it was not yet clear whether the Rac1/ $\beta$ -catenin association facilitated nuclear import or retention of  $\beta$ -catenin or, alternatively, Rac1 augments the function of  $\beta$ -catenin as a coactivator. Given the results of the present study, however, it is likely that Rac1 also plays a critical role in the nuclear transport of  $\beta$ -catenin, suggesting a general role of Rac1 GTPase for the nuclear transport of transcription factors. It is tempting to speculate that Rac1 is a molecular link between changes in cytoskeletal organization and alterations in transcription.

## Materials and methods

### Culture, cytokines, and antibodies

Ba/F3 cells were maintained in RPMI1640 medium (Invitrogen) containing 10% FCS and 1 ng/ml mIL-3 (R&D Systems). An ecotropic retrovirus packaging cell line PLATE was maintained as described previously (Hirose et al., 2001). An anti-STAT5A antibody and anti-STAT5B antibody were obtained from R&D Systems. Affinity-purified anti-MgcRacGAP antibody was produced as described previously (Hirose et al., 2001). An anti-Rac1 mAb and anti-importin  $\alpha$ 1 mAb were purchased from BD Biosciences. The rabbit polyclonal anti-Rac1, anti-RhoA, anti-JAK2, and goat polyclonal anti-HDAC or anti-importin  $\alpha$ 5 antibodies were obtained from Santa Cruz Biotechnology, Inc.

### Immunoprecipitation and Western blotting

Immunoprecipitation, gel electrophoresis, and immunoblotting were done as described previously (Kawashima et al., 2001), with minor modifications. Cell lysates ( $2 \times 10^7$  cells/ml) were incubated at 4°C for 2 h with the indicated antibodies and protein A-Sepharose. The immunoprecipitates were subjected to Western blot analysis with an anti-p-STAT5 mAb (Upstate Biotechnology), anti-MgcRacGAP, or anti-STAT5A antibody. The loading amounts were verified with the anti-STAT5A or anti-MgcRacGAP antibody after stripping the filters. The filter-bound antibody was detected using the ECL system (GE Healthcare). Cytosol and nuclear fractions were prepared as described previously (Nakamura et al., 2002).

### MBP pull-down assays

MBP fusion proteins (0.5  $\mu$ g) bound to amylose resin beads were incubated with cell lysates (10  $\mu$ g) from IL-3-stimulated Ba/F3 cells as described previously (Tonozuka et al., 2004).

### Transfection and immunostaining

The 293T cells were transfected with 1.0  $\mu$ g pME/STAT5A-Flag together with 0.5  $\mu$ g pMKIT (MOCK) or pMKIT/ITD-Flt3, and in some experiments cells were transfected with 0.5  $\mu$ g pME/STAT5A-HA and 0.5  $\mu$ g pMKIT (MOCK) or pMKIT/ITD-Flt3 together with 1.0  $\mu$ g pCMV5/N17Rac1-Flag, using Lipofectamine Plus reagents (Life Technologies). After 24 h, cells were plated on glass coverslips, and the next day the cells were immunostained as described previously (Hirose et al., 2001).

### Microscopy

Fluorescence images were analyzed on a confocal microscope (Fluoview FV300 Scanning Laser Biological Microscope IX 70 system; Olympus) equipped with two lasers (Ar 488 and HeNe 543) using a 60 $\times$  oil objective (PlanApo; Olympus). Fluoview version 4.3 software (Olympus) was used for image acquisition from confocal microscopy. Photoshop 7.0 or Photoshop Elements 2.0 software (Adobe) was used for processing of images.

### RNA interference and semiquantitative RT-PCR

For the silencing of Rac1 or MgcRacGAP, SMARTpool Rac1 or MgcRacGAP siRNA (L041170 or L040081; Dharmacon) was used. A control siRNA was used as a nonsilencing control (Tonozuka et al., 2004). 5  $\mu$ l of 40  $\mu$ M double-stranded siRNA were introduced into  $2 \times 10^6$  cells of Ba/3F cells with Nucleofector II (Amaxa) set at program T-16 using a Cell Line Nucleofector kit V (Amaxa) according to the manufacturer's instruction. A control vector carrying GFP was introduced to >80% of Ba/3F cells under this condition. 24 h after transfection, live cells were isolated using Ficol-Paque PLUS (GE Healthcare), and gene expression was examined by semiquantitative RT-PCR analysis as described previously (Nosaka et al., 1999). The primers used are as follows: 5' bcl-x, 5'-GAAAGAATTCACCAT-GTCTCAGAGCAACCGG-3'; 3' bcl-x, 5'-GAAAGCGCCGCTCACTCC-GACTGAAGAGTG-3'; 5' GAPDH, 5'-ACCACAGTCCATGCCATCAC-3'; 3' GAPDH, 5'-TCCACCACCTGTGTGTA-3'.

### Production of retroviruses

High-titer retroviruses harboring Cre recombinase were produced in a transient retrovirus packaging cell line PLATE (Morita et al., 2000) and were used to deplete Rac1 in Rac2<sup>-/-</sup>Rac1<sup>lox/lox</sup> fibroblasts (Fig. 3 B).

### Generation, expression, and purification of recombinant proteins in Sf-9 cells

To construct baculovirus vectors, the cDNAs encoding STAT5A, MgcRacGAP, V12Rac1, L61Rac1, N17Rac1, importin  $\alpha$ s, importin  $\beta$ 1, Ran, and NTF2 with the C-terminal Flag epitope tag, and a kinase domain of JAK2 (JH1; Saharinen et al., 2000) were subcloned into pBacPAK (BD Biosciences). The resulting constructs were used to obtain recombinant baculoviruses by cotransfection with Bsu36 I-digested BacPAK viral DNA (BD Biosciences) into Sf-9 cells according to the manufacturer's protocol. For protein expression, Sf-9 cells were infected with high-titer viral stocks for 96 h and lysed. The lysate was clarified by centrifugation, and the supernatant was immunoprecipitated with the anti-Flag M2-agarose affinity gel (Sigma-Aldrich) for 2 h at 4°C. The recombinant Flag-tagged proteins were eluted with 3 $\times$  Flag peptide (Sigma-Aldrich).

### EMSA using purified p-STAT5A

To determine whether purified p-STAT5A formed a proper dimer, EMSA was performed using consensus sequence of STAT5A as a probe, as described previously (Kawashima et al., 2001).

### In vitro kinase reaction

An in vitro kinase reaction of purified STAT5A was performed as described previously with minor modifications (Quelle et al., 1995). In vitro phosphorylated STAT5A was immunoprecipitated with the anti-Flag M2-agarose affinity gel and reeluted with a 3 $\times$  Flag peptide. The purified in vitro phosphorylated STAT5A was dialyzed against TB, and the final concentrations of STAT5A protein were determined for use in SDS-PAGE and in the nuclear transport assay.

### Preparation of fluorescent conjugates

FITC-labeled BSA (Sigma-Aldrich) conjugated with a synthetic peptide containing the SV40 large T antigen (CGGGPKKKRKVED; NLS-conjugated FITC-BSA) was prepared as described previously (Adam et al., 1990), as a control protein harboring an NLS. We confirmed that NLS-conjugated FITC-BSA was imported to the nucleus in our experimental conditions as reported previously (Kutay et al., 1997), which was not inhibited by immunodepletion of MgcRacGAP or Rac1 (Fig. S5, p-i).



### Import assays with permeabilized cells

HeLa cells were grown on poly-L-lysine-coated coverslips and permeabilized with 40  $\mu\text{g/ml}$  digitonin (Roche) in TB [20 mM Hepes, pH 7.3, 110 mM KOAC, 2 mM Mg(OAC)<sub>2</sub>, 1 mM EGTA, 2 mM DTT, 0.4 mM PMSF, 3  $\mu\text{g/ml}$  aprotinin, 2  $\mu\text{g/ml}$  pepstatin A, 1  $\mu\text{g/ml}$  leupeptin, and 20 mg/ml BSA] for 10 min at RT. Subsequently, the cells were washed twice in TB. Incubation with 50  $\mu\text{l}$  import mix was performed at 37°C for 30 min. The import mix contained TB, an energy regenerating system (0.5 mM ATP, 0.5 mM GTP, 10 mM creatine phosphate, and 30 U/ml creatine phosphokinase), and 1  $\mu\text{M}$  of purified unphosphorylated or phosphorylated STAT5A alone, or STAT5A plus the 1  $\mu\text{M}$  of other purified cofactor proteins as indicated in Fig. 6 C. After the import reaction, the cells were washed with ice-cold TB and immunostained with the anti-STAT5A antibody and anti-p-STAT5 mAb as described previously (Hirose et al., 2001). Fixed cells were examined using a Fluoview FV300 confocal microscope (Olympus).

### Online supplemental material

Fig. S1 depicts the binding domains of MgcRacGAP with STAT5A and that of STAT5A with MgcRacGAP. Fig. S2 shows that N17Rac1 expression inhibits the nuclear translocation of p-STAT5A. Fig. S3 shows that the DB2 region is required for the interaction of STAT3 with MgcRacGAP and the transcriptional activation of STAT3. Fig. S4 shows that purified p-STAT5A forms a dimer and binds DNA containing the STAT5 consensus sequence and that the purified STAT5A can be phosphorylated *in vitro*. Fig. S5 shows that the *in vitro* phosphorylated recombinant STAT5A can be imported to the nucleus in the nuclear transport assay and that immunodepletion of Rac1 or MgcRacGAP specifically inhibits the nuclear import of p-STAT5A using HeLa cytosol extract. Online supplemental material is available at <http://www.jcb.org/cgi/content/full/jcb.200604073/DC1>.

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