



The M-Ras-RA-GEF-2-Rap1 Pathway Mediates Tumor Necrosis Factor- α -dependent Regulation of Integrin Activation in Splenocytes

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(Degree)

博士 (医学)

(Date of Degree)

2007-09-25

(Date of Publication)

2008-02-04

(Resource Type)

doctoral thesis

(Report Number)

甲4061

(URL)

<https://hdl.handle.net/20.500.14094/D1004061>

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The M-Ras-RA-GEF-2-Rap1 Pathway Mediates Tumor Necrosis Factor- α -dependent Regulation of Integrin Activation in Splenocytes

**M-Ras-RA-GEF-2-Rap1 経路は、脾臓細胞において
Tumor Necrosis Factor- α 依存性のインテグリン活性化を制御する**

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饗場 篤、片桐 晃子、木梨 達雄、中尾 和貴、片岡 徹

Ras ファミリー 低分子量 G 蛋白質 グアニンヌクレオチド交換因子 **M-Ras**
RA-GEF-2 Rap1 インテグリン TNF- α 細胞接着 脾臓 B 細胞

Title: The M-Ras-RA-GEF-2-Rap1 Pathway Mediates Tumor Necrosis Factor- α -dependent Regulation of Integrin Activation in Splenocytes

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Running head: Role of RA-GEF-2 in integrin activation

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Abbreviations used: ES, embryonic stem; GAPDH, glyceraldehyde-3-phosphate dehydrogenase; GEF, guanine nucleotide exchange factor; HA, hemagglutinin; HBSS, Hank's balanced salt solution; ICAM-1, intercellular adhesion molecule 1; IL, interleukin; LFA-1, lymphocyte function-associated antigen 1; PBS, phosphate-buffered saline; PCR, Polymerase Chain Reaction; PLC, phospholipase C; PMA, phorbol myristate acetate; RA, Ras/Rap1-associating; RID, RalGDS-Ras-interacting domain; RNAi, RNA Interference; RT, reverse transcription; SDF-1, stromal cell-derived factor-1; siRNA, small interfering RNA; TNF- α , tumor necrosis factor- α ; TRAF, TNF receptor-associated factors.

Abstract

The Rap1 small GTPase has been implicated in regulation of integrin-mediated leukocyte adhesion downstream of various chemokines and cytokines in many aspects of inflammatory and immune responses. However, the mechanism for Rap1 regulation in the adhesion signaling remains unclear. RA-GEF-2 is a member of the multiple-member-family of guanine nucleotide exchange factors (GEFs) for Rap1 and characterized by the possession of a Ras/Rap1-associating domain, interacting with M-Ras-GTP as an effector, in addition to the GEF catalytic domain. Here, we show that RA-GEF-2 is specifically responsible for the activation of Rap1 that mediates tumor necrosis factor- α (TNF- α)-triggered integrin activation. In BAF3 hematopoietic cells, activated M-Ras potently induced lymphocyte function-associated antigen 1 (LFA-1)-mediated cell aggregation. This activation was totally abrogated by knockdown of RA-GEF-2 or Rap1. TNF- α treatment activated LFA-1 in a manner dependent on M-Ras, RA-GEF-2 and Rap1, and induced activation of M-Ras and Rap1 in the plasma membrane, which was accompanied by recruitment of RA-GEF-2. Finally, we demonstrated that M-Ras and RA-GEF-2 were indeed involved in TNF- α -stimulated and Rap1-mediated LFA-1 activation in splenocytes by using mice deficient in RA-GEF-2. These findings proved a crucial role of the crosstalk between two Ras-family GTPases M-Ras and Rap1, mediated by RA-GEF-2, in adhesion signaling.

1 INTRODUCTION

2
3 The integrin family member lymphocyte function-associated antigen 1 (LFA-1), a
4 heterodimer consisting of α_L and β_2 subunits, is involved in diverse aspects of leukocyte
5 function, including extravasation, migration and immunological synapse formation with
6 antigen-presenting cells (Dustin *et al.*, 2004; Kinashi, 2005). Intercellular adhesion molecule
7 1 (ICAM-1) interacts with LFA-1 with the highest affinity to mediate adhesion. LFA-1 on
8 circulating lymphocytes has low avidity, which is required to be upregulated by extracellular
9 stimuli for ligand binding. Intracellular signals that increase LFA-1 avidity, which are called
10 "inside-out" signals, are triggered by various cytokines, chemokines and antigen stimulation
11 of the T cell receptor. Although diverse signaling molecules have been implicated in the
12 regulation of LFA-1 avidity, mechanisms underlying this signaling remain largely unknown.

13 Rap1 is a member of the Ras family of small GTPases, and is highly expressed in
14 hematopoietic cells including lymphocytes (Bos *et al.*, 2001; Stork and Dillon, 2005). In its
15 GTP-bound active form, Rap1 interacts with various effector molecules to initiate
16 downstream signaling pathways. The first identified Rap1 function is the antagonism to
17 Ras-dependent activation of the Raf-1/extracellular signal-regulated kinase cascade
18 (Kitayama *et al.*, 1989; Cook *et al.*, 1993), which may also underlie the role of Rap1 in T cell
19 anergy (Boussiotis *et al.*, 1997). Another Raf family member B-Raf, in contrast to Raf-1, is
20 directly activated by Rap1, leading to the activation of the extracellular signal-regulated
21 kinase pathway (Ohtsuka *et al.*, 1996; Vossler *et al.*, 1997). The different Rap1 action on
22 these two Raf kinases is attributable to the difference in affinity of Rap1 to the cysteine-rich
23 domain (Okada *et al.*, 1999).

24 The involvement of Rap1 in integrin-mediated cell adhesion was proposed from the
25 effect of the overexpression of activated Rap1 and its negative regulator SPA-1 (Tsukamoto
26 *et al.*, 1999; Caron *et al.*, 2000; Katagiri *et al.*, 2000; Reedquist *et al.*, 2000). Two Rap1
27 effectors, RAPL and RIAM, have been shown to act as a link between activated Rap1 and
28 integrin (Katagiri *et al.*, 2003; Lafuente *et al.*, 2004). RAPL was isolated as a Rap1 effector
29 enriched in lymphoid tissues, containing a Ras/Rap1-associating (RA) domain, and was

shown to induce lymphocyte polarization and the redistribution of LFA-1, leading to enhanced adhesion (Katagiri *et al.*, 2003). In support of a pivotal role for RAPL, *RAPL* gene knockout, in fact, caused an impairment in lymphocyte adhesion and migration (Katagiri *et al.*, 2004). Recently, the serine/threonine kinase Mst1 was identified as a binding protein of RAPL, and its involvement in chemokine-induced cell polarization and LFA-1-mediated adhesion was demonstrated (Katagiri *et al.*, 2006).

The activation of Rap1 in response to various upstream signals is mediated by guanine nucleotide exchange factors (GEFs) (Bos *et al.*, 2001). The first identified Rap1 GEF termed C3G associates with the adaptor protein Crk, and forms a complex with receptor and non-receptor protein tyrosine kinases in response to extracellular stimuli (Gotoh *et al.*, 1995). Epac1 (also called cAMP-GEFI) and Epac2 (also called cAMP-GEFII) are activated by direct binding of cAMP, being responsible for cAMP-dependent Rap1 activation (de Rooij *et al.*, 1998; Kawasaki *et al.*, 1998; de Rooij *et al.*, 2000). Another Epac subfamily member called Repac (also called GFR/MR-GEF) binds to the activated form of M-Ras, which downregulates the activity of Repac (Ichiba *et al.*, 1999; de Rooij *et al.*, 2000; Rebhun *et al.*, 2000). The third subfamily is constituted of two calcium- and diacylglycerol-regulated Rap1 GEFs termed CalDAG-GEFI and CalDAG-GEFIII, which contain calcium-binding EF and diacylglycerol-binding C1 domains (Kawasaki *et al.*, 1998; Yamashita *et al.*, 2000).

Two related GEFs called RA-GEF-1 (also called PDZ-GEF1/nRapGEP/CNrasGEF) (de Rooij *et al.*, 1999; Liao *et al.*, 1999; Ohtsuka *et al.*, 1999; Pham *et al.*, 2000) and RA-GEF-2 (Gao *et al.*, 2001) constitute another Rap1 GEF subfamily. These two GEFs have both GEF and RA domains, serving not only as an upstream regulator, but also as a downstream target, of Ras family small GTPases. In fact, RA-GEF-1 acts both downstream and upstream of Rap1, amplifying Rap1-dependent B-Raf activation in the Golgi apparatus (Liao *et al.*, 1999; Liao *et al.*, 2001). On the other hand, RA-GEF-2 mediates M-Ras-dependent Rap1 activation in the plasma membrane (Gao *et al.*, 2001). Although RA-GEF-1 and RA-GEF-2 have been biochemically characterized as a link between Ras family small GTPases, specific signaling that involves these GEFs and Rap1 remains elusive.

1 Furthermore, the physiological function of these GEFs in developing and adult mice remains
2 totally unknown.

3 In this paper, we examine the role of RA-GEF-2 in the regulation of Rap1 that is
4 responsible for integrin-mediated lymphocyte adhesion by using both cultured cell lines and
5 RA-GEF-2-deficient mice.

6

MATERIALS AND METHODS

Cells and cDNA Transfection

The mouse hematopoietic cell line BAF3 expressing human LFA-1 (BAF/hLFA-1) was suspended with RPMI 1640 containing 2 mM L-glutamine, 50 μ M β -mercaptoethanol, 10% fetal bovine serum and 10% WEHI-3B conditioned medium as a source of interleukin (IL)-3 (Katagiri *et al.*, 2000). The cDNA for constitutively active mutant human M-Ras (M-Ras[Q71L]) was subcloned into a mammalian expression vector, pcDNA3.1HisC (Invitrogen, Carlsbad, CA), generating pcDNA3.1HisC-M-Ras[Q71L]. For the stable expression of M-Ras[Q71L], BAF/hLFA-1 cells were transfected with pcDNA3.1HisC-M-Ras[Q71L] by electroporation at 300 V and 900 μ F and incubated for 24 h at 37°C. After incubation, cells were selected with G418 (SIGMA, St. Louis, MO) at 1 mg/ml, and several BAF/hLFA-1/M-Ras[Q71L] clones were isolated by a limiting dilution. Construction of pFLAG-CMV2-M-Ras[Q71L], pFLAG-CMV2-M-Ras, pFLAG-CMV2-H-Ras, pFLAG-CMV2-Rap1, and pEF-BOS-HA-Rap1 was previously described (Liao *et al.*, 1999; Gao *et al.*, 2001). These plasmids were introduced into BAF/hLFA-1 cells by electroporation as described above.

Antibodies

Anti-RA-GEF-2 antiserum against a synthetic peptide (CLEPRDITDPVYKTVTSSTD), which corresponds to the C-terminal amino acid sequence of RA-GEF-2, was raised in rabbits (Operon Biotechnologies, Tokyo, Japan). The antibody was affinity-purified from the antiserum using Sepharose 4B resin conjugated with the antigenic peptide. Mouse monoclonal anti-actin (Santa Cruz Biotechnology, Santa Cruz, CA), goat polyclonal anti-M-Ras (Santa Cruz Biotechnology), rabbit polyclonal anti-Rap1 (Santa Cruz Biotechnology), mouse monoclonal anti-FLAG (SIGMA), mouse monoclonal anti-6 \times His (Clontech, Palo Alto, CA), and mouse monoclonal anti-hemagglutinin (HA) (Invivogen, San Diego, CA), mouse monoclonal anti-human LFA-1 (MEM-25) (MONOSAN®, Uden, NED), and rat monoclonal anti-mouse LFA-1 (M17/4) (Santa Cruz Biotechnology) antibodies were commercially

obtained.

Immunoblotting

Proteins were extracted from BAF3-derived cells or mouse tissues by lysis buffer A (50 mM Tris-HCl [pH 7.4], 200 mM NaCl, 2.5 mM MgCl₂, 10% glycerol, 1% Nonidet P-40, 1 mM phenylmethylsulfonyl fluoride, 1 mM leupeptin) and subjected to sodium dodecyl sulfate-polyacrylamide gel electrophoresis. Proteins were transferred to a nitrocellulose membrane (Whatman, Brentford, UK) and incubated with primary antibodies, followed by horseradish peroxidase-conjugated secondary antibodies. The enhanced chemiluminescence system (GE Healthcare Bio-Sciences, Piscataway, NJ) was applied for detection.

Gene Silencing by RNA Interference (RNAi)

BAF3-derived cells were transfected with StealthTM small interfering RNAs (siRNAs) against mouse RA-GEF-2 (sense sequence; 5'-ggacuccugaggacuuaauuuau-3'), mouse Rap1A (sense sequence; 5'-aaggacuacuagcuuguacucacgc-3'), and control (negative control Med GC) by electroporation at 300 V and 300 μ F. After electroporation, cells were incubated for 48 h at 37°C and subjected to further experiments. Spleen cells were transfected with siRNAs against mouse M-Ras (sense sequence; 5'-ucagguaggagucucaaugguggg-3') and control followed by incubation for 24 h at 37°C. StealthTM siRNAs were purchased from Invitrogen.

Immunostaining

BAF/hLFA-1 cells were transfected with pFLAG-CMV2-M-Ras[Q71L], incubated for 24-32 h at 37°C, and then serum-starved for another 16 h. Following fixation with 4% paraformaldehyde for 15 min, cells were mounted on aminosilane-coated slide glasses (MATSUNAMI GLASS, Osaka, Japan) until dried-up and permeabilized by cold methanol for 1 min on ice. Cells were stained with mouse monoclonal anti-FLAG antibody and Alexa Fluor® 546-labeled goat anti-mouse IgG (H+L) (Molecular Probes, Eugene, OR) for M-Ras[Q71L], and anti-RA-GEF-2 antibody and Alexa Fluor® 488-labeled goat anti-rabbit IgG (H+L) (Molecular Probes) for endogenous RA-GEF-2. After staining, cells were

observed by confocal laser scanning microscopy (LSM510 META; Carl Zeiss, Jena, Germany).

Cell Preparation from the Thymus and Spleen

Thymocytes and splenocytes were prepared essentially as described (Sebzda *et al.*, 2002; Duchniewicz *et al.*, 2006; Kawasaki *et al.*, 2006). The thymus and spleen were disaggregated with fine forceps, passed through a fine mesh filter to obtain a single cell suspension and washed twice with Hank's balanced salt solution (HBSS; Invitrogen). After washing, cells were incubated for 5 min in erythrocyte lysis buffer (155 mM NH₄Cl, 10 mM KHCO₃, 0.1 mM EDTA) to remove erythrocytes and washed with HBSS three times.

Adhesion Assays

BAF/hLFA-1 cells were transfected with pFLAG-CMV2-M-Ras, pFLAG-CMV2-H-Ras, pFLAG-CMV2-Rap1 or a combination of pFLAG-CMV2-M-Ras and pEF-BOS-HA-Rap1 and incubated for 48 h at 37°C. Maxisorb 96-well plates (Nunc, Roskilde, DK) were coated with 100 µl of 2 µg/ml ICAM-1/Fc (R&D Systems, Minneapolis, MN) in phosphate-buffered saline (PBS) (Invitrogen) at 4°C over night, washed three times with PBS, and blocked with 2% bovine serum albumin in HBSS for 1 hr at 37°C. The blocked 96-well plates were washed three times with 0.5% bovine serum albumin in HBSS. Cells were collected and washed with HBSS three times and stained with 2.5 µM biscarboxyethyl-carboxyfluorescein acetoxymethylether (BCECF/AM; Calbiochem, San Diego, CA) in HBSS for 30 min at 37°C. After staining, cells were washed with HBSS three times and suspended in adhesion assay medium (RPMI 1640 containing 10 mM Hepes [pH 7.4] and 5% fetal bovine serum). Stained BAF3-derived cells (1×10^5 /well) or thymus and spleen cells (1×10^6 /well) suspended in 100 µl of adhesion assay medium were stimulated with phorbol myristate acetate (PMA) (10 ng/ml) or tumor necrosis factor- α (TNF- α) (10 ng/ml) for 30 min at 37°C. For inhibition, stained cells were incubated with 20 µg/ml anti-human LFA-1 (MEM-25) or anti-mouse LFA-1 (M17/4) antibodies for 30 min before loading (Garnotel *et al.*, 1995; Mueller *et al.*, 2004). Non-adherent cells were removed by washing three times with adhesion assay medium

prewarmed at 37°C. Cells bound to ICAM-1 were detected by fluorescence analyzer (FLA3000G; Fuji-film, Tokyo, Japan) at an excitation of 473 nm and an emission of 520 nm. The adhesion was represented as a percentage of total input detected before incubation.

Production of Recombinant Retrovirus and Infection

The SPA-1 cDNA was a generous gift from Drs. Masakazu Hattori and Nagahiro Minato (Kyoto University, Kyoto, Japan). Retroviral expression vectors pLPCX (Clontech) and pLPCX carrying the FLAG-tagged mouse SPA-1 cDNA were introduced into Phoenix ecotropic retroviral packaging cells (ATCC inventory #SD3444) with the Superfect transfection reagent (Qiagen, Valencia, CA) according to the manufacturer's protocol to produce the viral supernatant. After incubation for 48 h at 37°C, the viral supernatant was collected, filtrated, and used for infection. BAF/hLFA-1/M-Ras[Q71L] cells (5×10^5) were suspended in 1 ml of the viral supernatant with 10 µg/ml polybrene for 2 h at 32°C. After infection, cells were incubated in medium for 24 h at 37°C and then selected for 10 days in the presence of 2 µg/ml puromycin.

Preparation of the Plasma Membrane Fraction

pEF-BOS-HA-Rap1 was transfected into BAF/hLFA-1 cells with or without pFLAG-CMV2-M-Ras[Q71L]. After transfection, cells were incubated for 24-32 h at 37°C and then serum-starved for another 16 h. To see TNF-α-dependent activation of M-Ras and Rap1, pFLAG-CMV2-M-Ras and pEF-BOS-HA-Rap1 were co-transfected into BAF/hLFA-1 cells and incubated for 48 h at 37°C. After incubation, cells were treated with or without 10 ng/ml TNF-α for 30 min at 37°C. The plasma membrane fraction was prepared by sucrose density gradient centrifugation as described (Gao *et al.*, 2001). In this fraction, the plasma membrane is highly enriched as estimated by the specific activity of the marker enzyme alkaline phosphatase (Gao *et al.*, 2001). The plasma membrane fraction resuspended in lysis buffer A was subjected to pull-down assays. For *in vivo* study, the plasma membrane-enriched fraction was prepared from spleen cells of wild-type mice as described (Peirce *et al.*, 2004) with minor modifications. Briefly, spleen cells were transfected with siRNAs against mouse

M-Ras and control for 24 h at 37°C, followed by TNF- α treatment for 30 min at 37°C. After treatment, cells were resuspended in hypotonic lysis buffer B (20 mM Tris-HCl [pH 7.4], 1 mM MgCl₂, 1 μ g/ml leupeptin, 1 mM phenylmethylsulfonyl fluoride), homogenized using a Potter-Elvehjem homogenizer and centrifuged at 4,000 \times g for 5 min. The supernatant was further centrifuged at 20,000 \times g for 30 min, and the pellet containing the plasma membrane, but not microsomes, was resuspended in lysis buffer A, and subjected to pull-down assays.

Pull-down Assays

Extracts from the plasma membrane fraction or plasma membrane-enriched fraction prepared by lysis buffer A were added to the RalGDS-Ras-interacting domain (RID) (for Rap1-GTP) or the c-Raf-Ras-binding domain (for M-Ras-GTP), which had been immobilized on glutathione agarose resins, and mixed with slow agitation for 1 h at 4°C. Resins were washed three times with lysis buffer A, and bound proteins were eluted from resins by sodium dodecyl sulfate-polyacrylamide gel electrophoresis sample buffer followed by immunoblotting.

Flow Cytometric Analysis

Spleen cells were incubated with fluorescein isothiocyanate-conjugated anti-mouse LFA-1 (M17/4) antibody (1 μ g per 1 \times 10⁶ cells) in HBSS on ice for 30 min, washed with HBSS three times, and subjected to flow cytometric analysis by FACScaliber (Beckton Dickinson, San Jose, CA).

Separation of Splenic B and non-B Cells

Polystyrene flasks (25 cm², CORNING, Corning, NY) were coated with 2 ml of 1 mg/ml rabbit anti-mouse immunoglobulins antibody (DAKO, Glostrup, Denmark) in PBS at 4°C over night and washed three times with PBS. After preparation of splenocytes, cells were resuspended in adhesion assay medium (1.5 \times 10⁷/ml), and 3 ml of cell suspension was added to the flask. After incubation for 1 hr at 37°C with gentle swirling, non-adherent cells (non-B cell enrichment, 4.5 \times 10⁷ cells/spleen) were carefully resuspended by rocking the flask and

recovered from the cell suspension medium. The flask with adherent cells (B cell enrichment, 1.0×10^8 cells/spleen) was washed 5 times with PBS, and 2 ml of 4 mg/ml lidocaine (SIGMA) solution in PBS was added. After incubation for 15 min, cells were removed from the flask by pipetting, collected by centrifugation ($200 \times g$, 10 min), and washed twice with PBS.

Construction of the Targeting Vector

An *RA-GEF-2* genomic DNA fragment was cloned from a 129/Sv mouse genomic bacterial artificial chromosome library (Invitrogen) and used for the construction of a targeting vector. A 522-bp *MfeI*-*BsgI* fragment, which harbors exon 21 of *RA-GEF-2* coding for the part of the GEF domain, was inserted into a construct in which it is flanked with loxP at its 5' end and with loxP-*TK-neo*-loxP at its 3' end. *TK-neo*, which expresses the neomycin-resistance gene under the control of the thymidine kinase promoter, was inserted in an inverted orientation. Subsequently, a 9.2-kb *KpnI*-*MfeI* fragment and a 4.3-kb *BsgI*-*StyI* fragment of *RA-GEF-2* were inserted as the 5'- and 3'-arms for homologous recombination, respectively (Figure 5A). Finally, the diphtheria toxin A chain cassette for negative selection was inserted into the 3' end of the right arm.

Gene Targeting and Generation of Mutant Mice

To generate the floxed *RA-GEF-2* (*RA-GEF-2^{lox}*) allele, in which exon 21 is flanked with loxP sites (Figure 5A), mouse EB3 embryonic stem (ES) cells (a gift from Dr. Hitoshi Niwa, RIKEN Center for Developmental Biology, Kobe, Japan) derived from the 129/Ola strain were transfected with the targeting vector, linearized by *NotI* cleavage, by electroporation at 0.8 kV, 3 μ F and subjected to selection with G418. G418-resistant 768 clones were isolated and screened by Southern blot analysis of their genomic DNAs. One ES clone, carrying the properly generated *RA-GEF-2^{lox}* allele, was injected into mouse C57BL/6 blastocysts to generate chimeric males, which were subsequently bred with C57BL/6 females to generate *RA-GEF-2^{+ / flox}* mice. Subsequently, *RA-GEF-2^{+ / flox}* mice were bred with *CAG-Cre* transgenic mice (a generous gift from Dr. Jun-ichi Miyazaki, Osaka Univ., Osaka, Japan) (Sakai and Miyazaki, 1997) to yield *RA-GEF-2^{+ / -}* mice carrying an *RA-GEF-2* allele with a deletion of

1 exon 21-loxP-*TK-neo* by Cre-mediated recombination between the terminal loxP sites. Finally,
 2 *RA-GEF-2*^{-/-} mice were generated by interbreeding *RA-GEF-2*^{+/-} mice. All animals were
 3 maintained under standard housing condition with a 12-h~12-h dark-light cycle at the animal
 4 facilities of Kobe University Graduate School of Medicine according to institutional
 5 guidelines.

7 ***Genotyping by Southern Blotting and Polymerase Chain Reaction (PCR)***

8 Genomic DNAs isolated from ES cells or mouse tails were digested by *Bsu36I*, separated by
 9 agarose gel electrophoresis and transferred to a nylon membrane (Hybond N+, GE Healthcare
 10 Bio-Sciences) with alkaline transfer buffer (0.4 N NaOH, 0.6 M NaCl). Membranes were
 11 hybridized with the ³²P-radiolabeled 3' probe (Figure 5A) generated by PCR (the sense and
 12 antisense primers were 5'-ccgtcatattattcgaatgacttctgcc-3' and 5'-gcctcagctataaagtgagtacag-3',
 13 respectively), and the hybridized signals were detected by STORM bioimage analyzer (GE
 14 Healthcare Bio-Sciences). For genotyping by PCR, a trio of primers were used for
 15 amplification of the *RA-GEF-2* alleles; #1 (5'-gagccttgagatacagaaacttg-3') located upstream
 16 of the 5'-terminal loxP site in the *RA-GEF-2*^{lox} allele, #2 (5'-cttgacaacagggaagagtg-3') in
 17 exon 21, and #3 (5'-ctaggaggtgtcagcaaag-3') downstream of the 3'-terminal loxP site. The
 18 amplified DNA fragments were separated by agarose gel electrophoresis.

20 ***Reverse Transcription (RT)-PCR Analysis***

21 Mouse spleen cells were transfected with siRNAs against mouse M-Ras and control,
 22 incubated for 24 h at 37°C, and treated with TNF-α for 30 min at 37°C. Transcripts were
 23 prepared by using the Sepasol(R)-RNA I Super kit (Nacalai tesque, Kyoto, Japan) according
 24 to the manufacturer's protocol. The first-strand oligo(dT)-primed cDNA was synthesized as
 25 previously described (Wu *et al.*, 2003). Primers used for amplification of M-Ras were
 26 5'-cgctgttccaagtgaaaacc-3' and 5'-ggctgtcacaagatgacac-3'. Primers used for amplification of
 27 *RA-GEF-2* were 5'-gaggcacttgggaaaagtg -3' and 5'-cttgacaacagggaagagtg -3'. Primers for
 28 amplification of mouse glyceraldehyde-3-phosphate dehydrogenase (GAPDH)
 29 (5'-gtgaaggctcggtgtgaacggattt-3', and 5'-cacagtcttctgagtggcagtgat-3') were used as an internal

1 control.
2

1 RESULTS

3 *RA-GEF-2 Mediates M-Ras-induced Rap1 Activation and LFA-1-ICAM-1-dependent Cell* 4 *Aggregation*

5 To evaluate the involvement of M-Ras and RA-GEF-2 in cell adhesion through the interaction
6 between LFA-1 and ICAM-1, we employed the BAF/hLFA-1 cell line, which was derived
7 from nonadherent IL-3-dependent mouse BAF3 cells (Katagiri *et al.*, 2000). BAF/hLFA-1
8 cells stably express human LFA-1, and show PMA-dependent adhesion to ICAM-1 (Katagiri
9 *et al.*, 2000). In addition, overexpression of constitutively activated Rap1 potently enhanced
10 the binding of BAF/hLFA-1 cells to ICAM-1, suggesting that Rap1 is involved in
11 LFA-1-ICAM-1-dependent cell adhesion (Katagiri *et al.*, 2000). We isolated a
12 BAF/hLFA-1-derived clone (designated BAF/hLFA-1/M-Ras[Q71L]), which stably expresses
13 constitutively activated M-Ras mutant (M-Ras[Q71L]) (Figure 1A). Endogenous RA-GEF-2
14 may exist as two splice variants as suggested by doublet bands in immunoblot analysis
15 (Figure 1A) and the occurrence of two mRNA species previously reported (Gao *et al.*, 2001).
16 The expression of endogenous M-Ras in BAF/hLFA-1 cells was undetectable by anti-M-Ras
17 antibody (Figure 1A). BAF/hLFA-1/M-Ras[Q71L] cells have an increased tendency to
18 aggregate, which was completely suppressed in the presence of anti-human LFA-1 antibody,
19 indicating that M-Ras-dependent cell aggregation was indeed mediated by LFA-1-ICAM-1
20 interaction (Figure 1B). We hypothesized that RA-GEF-2 may activate endogenous Rap1
21 upon overexpression of activated M-Ras, leading to the formation of cell aggregates. In fact,
22 when the expression of RA-GEF-2 was abrogated by RNAi, LFA-1-dependent cell
23 aggregation and adhesion to ICAM-1 were significantly reduced (Figure 2, A-C). Similarly,
24 the decrease in the expression level of Rap1 by its specific siRNA inhibited the
25 M-Ras[Q71L]-dependent cell aggregation and adhesion to ICAM-1, thus placing Rap1
26 downstream of M-Ras (Figure 2, A-C). Reduction in adhesion to ICAM-1 by RA-GEF-2 or
27 Rap1 knockdown was almost complete because adhesion of parental BAF/hLFA-1 cells to
28 ICAM-1 was 8.4 ± 2.2 (%). Furthermore, overexpression of SPA-1, a GTPase-activating
29 protein specific for Rap1, which negatively regulates Rap1 activity, potently inhibited

1 M-Ras-induced cell aggregation (Figure 2, D and E). Rap2, another substrate for RA-GEF-2,
2 was not expressed in BAF/hLFA-1 cells as determined by RT-PCR (data not shown).

3 We have shown that, in COS-7 cells, activated M-Ras was expressed exclusively in
4 the plasma membrane, and RA-GEF-2 was recruited from the cytoplasm to the plasma
5 membrane through the binding of its RA domain to M-Ras, when they were co-expressed
6 (Gao *et al.*, 2001). Likewise, activated M-Ras and RA-GEF-2 seemed to be co-localized, in
7 part, in the plasma membrane in BAF/hLFA-1/M-Ras[Q71L] cells, whereas RA-GEF-2 was
8 localized throughout the cytoplasm when expressed alone in BAF/hLFA-1 cells (Figure 3A).
9 In the majority of cells investigated, these proteins showed equivalent subcellular localization.
10 Recruitment of RA-GEF-2 to the plasma membrane was also detected by immunoblot
11 analysis when constitutively activated M-Ras was expressed (Figure 3B). The GTP-bound
12 active form of plasma membrane-localized Rap1 was consequently increased as shown by
13 pull-down assay in activated M-Ras-expressing cells (Figure 3B). In addition, downregulation
14 of RA-GEF-2 by RNAi inhibited Rap1 activation in the plasma membrane, indicating that
15 RA-GEF-2 mediated M-Ras-induced Rap1 activation (Figure 3B).

17 ***TNF- α Induces LFA-1-ICAM-1-dependent Adhesion Specifically through M-Ras***

18 A diverse array of cytokines modulates LFA-1-ICAM-1-dependent lymphocyte adhesion. As
19 a step to identify cytokines that stimulate the M-Ras-RA-GEF-2-Rap1 pathway to induce cell
20 adhesion, we examined the involvement of M-Ras in integrin activation signaling downstream
21 of the TNF- α receptor because Rap1 was reported to mediate TNF- α -triggered activation of
22 integrins (Caron *et al.*, 2000). For this purpose, we isolated BAF/hLFA-1-derived clones that
23 express wild-type M-Ras, Ha-Ras or Rap1, and then compared adhesion to ICAM-1 upon
24 TNF- α treatment. TNF- α selectively induced the adhesion of BAF/hLFA-1 cells harboring
25 M-Ras, but not those harboring Ha-Ras or Rap1 (Figure 4A). PMA, as a positive control,
26 enhanced the adhesion of all of the above cell lines (data not shown). Co-expression of Rap1
27 did not enhance M-Ras-mediated cell adhesion in response to TNF- α , suggesting that the
28 expression level of endogenous Rap1 is sufficiently high in BAF/hLFA-1 cells (Figure 4A).
29 The effect of TNF- α in M-Ras-expressing cells was dose-dependent with maximal induction

at the concentration of 10 to 100 ng/ml (Figure 4B). TNF- α -induced adhesion of these cells was indeed blocked by anti-LFA-1 antibody (Figure 4C). In addition, downregulation of RA-GEF-2 or Rap1 expression by respective siRNAs rendered cells insensitive to TNF- α (Figure 4D). TNF- α also induced the recruitment of RA-GEF-2 to the plasma membrane in cells expressing wild-type M-Ras (Figure 4E). This recruitment occurred presumably through the specific interaction of RA-GEF-2 with the GTP-bound form of M-Ras in the plasma membrane generated upon stimulation with TNF- α (Figure 4E). The Rap1-GTP level in the plasma membrane was also increased following TNF- α stimulation, probably due to the action of RA-GEF-2 recruited to this region (Figure 4E). Amounts of plasma membrane-localized M-Ras and Rap1 were unaffected by TNF- α treatment (Figure 4E). Taken together, in BAF3-derived cells, TNF- α specifically activated the signaling pathway consisting of M-Ras, RA-GEF-2 and Rap1, leading to LFA-1-dependent cell adhesion.

Generation of RA-GEF-2-deficient Mice

To evaluate the physiological relevance of the M-Ras-RA-GEF-2-Rap1 pathway in terms of lymphocyte adhesion, we generated mice with functional disruption of the *RA-GEF-2* gene. We first created mice carrying an *RA-GEF-2* allele with exon 21 (encoding the GEF domain) and the *TK-neo* cassette flanked by *loxP* sites (floxed allele) (Figure 5A). Hybridization of *Bsu36I*-digested genomic DNAs with a 3' external probe identified 7.7-kb and 8.8-kb bands of the wild-type and floxed alleles, respectively, in *RA-GEF-2*^{+/*lox*} mice (Figure 5B). Insertion of the 5' *loxP* site was also confirmed by PCR. Primers #1 and #2 (described in Figure 5A) amplified 540-bp and 586-bp products from the wild-type and floxed alleles, respectively (Figure 5C). *RA-GEF-2*^{+/*lox*} mice were then bred with mice carrying a transgene encoding the Cre recombinase under the control of the CAG promoter (Sakai *et al.*, 1997). Out-of-frame deletion of exon 21 resulted in functional disruption of the *RA-GEF-2* allele (knockout allele). *RA-GEF-2*^{+/-} mice were then intercrossed to produce *RA-GEF-2* homozygous null (*RA-GEF-2*^{-/-}) offsprings, in which the *RA-GEF-2* knockout allele was confirmed by PCR. Primers #1 and #3 (described in Figure 5A) amplified a 792-bp and 316-bp products from the wild-type and knockout alleles, respectively (Figure 5D). *RA-GEF-2*^{-/-} pups were born at the

expected Mendelian ratios ($RA-GEF-2^{+/+} : RA-GEF-2^{+/-} : RA-GEF-2^{-/-} = 26.7\% : 47.8\% : 25.6\%$; 90 pups) without any intrauterine loss or early death, and were fertile and indistinguishable from their $RA-GEF-2^{+/+}$ or $RA-GEF-2^{+/-}$ littermates in appearance and growth. $RA-GEF-2^{-/-}$ mice grew normally for at least 37 weeks. Anatomical examination revealed that their spleens were enlarged compared with those of wild-type mice, whereas the other tissues appeared grossly normal (Figure 5E). We observed no obvious histological abnormalities in the $RA-GEF-2$ knockout spleen.

The expression of the $RA-GEF-2$ protein in mouse tissues was examined by immunoblotting. $RA-GEF-2$ was highly expressed in the thymus and spleen, and weakly in the lung, brain and pancreas in a wild-type mouse (Figure 5F). $RA-GEF-2$ was also weakly expressed in other hematopoietic tissues such as lymph nodes and bone marrow (data not shown). As expected, no $RA-GEF-2$ protein was detected in $RA-GEF-2^{-/-}$ mouse tissues (Figure 5F).

LFA-1-ICAM-1-mediated Adhesion of Mouse Splenocytes

Thymocytes and splenocytes were isolated from wild-type and $RA-GEF-2^{-/-}$ mice, and their adhesion to ICAM-1 was examined. PMA and $TNF-\alpha$ enhanced adhesion of thymocytes and splenocytes from the wild-type mouse (Figure 6A). This adhesion was mediated by LFA-1 as indicated by the inhibitory effect of anti-LFA-1 antibody (Figure 6A). $RA-GEF-2$ knockout exerted virtually no effect on PMA- or $TNF-\alpha$ -stimulated thymocyte adhesion. In marked contrast, $TNF-\alpha$ -stimulated adhesion of $RA-GEF-2^{-/-}$ splenocytes was significantly reduced compared with wild-type ones, whereas PMA-stimulated adhesion remained unaffected (Figure 6A). Furthermore, downregulation of the expression of endogenous M-Ras or $RA-GEF-2$ in splenocytes by RNAi completely blocked $TNF-\alpha$ -stimulated adhesion (Figure 6B). The expression level of LFA-1 on the cell surface of splenocytes did not show any detectable difference between wild-type and $RA-GEF-2^{-/-}$ mice as determined by FACScan analysis, indicating that the impaired $TNF-\alpha$ -induced adhesion in $RA-GEF-2^{-/-}$ splenocytes was not due to the loss of LFA-1 expression (Figure 6C). To identify the cell population that actually responds to $TNF-\alpha$, we separated B and non-B cells from wild-type and $RA-GEF-2^{-/-}$

spleens, and their adhesion to ICAM-1 was examined. TNF- α stimulated the adhesion of splenic B cells, but not non-B cells, isolated from wild-type mice (Figure 6D). In contrast, B cells from *RA-GEF-2*^{-/-} splenocytes did not show any TNF- α -induced adhesion (Figure 6D). The expression level of M-Ras was almost the same in splenic B cells, splenic non-B cells and thymocytes isolated from wild-type and *RA-GEF-2*^{-/-} knockout mice (Figure 6D). *RA-GEF-2* expression levels in these cell types were also similar in wild-type mice (Figure 6D). In addition, we observed similar nuclear translocation of NF- κ B following TNF- α stimulation for 30 min in wild-type and *RA-GEF-2*-deficient splenic B cells, suggesting that TNF- α receptor expression did not alter significantly in *RA-GEF-2*-deficient cells (data not known). Collectively, *RA-GEF-2* is indispensable for transduction of TNF- α -induced adhesion signals in splenic B cells, whereas *RA-GEF-2* does not appear to play an essential role in adhesion signaling in thymocytes, even though it is abundantly expressed in the thymus as well.

Additionally, the activation of plasma membrane-localized Rap1 upon TNF- α treatment was detected by pull-down assays, which was abrogated when the expression of M-Ras was downregulated by RNAi (Figure 6E). Downregulation of M-Ras by RNAi also inhibited the recruitment of *RA-GEF-2* to the plasma membrane (Figure 6E). Importantly, TNF- α -induced Rap1 activation was completely inhibited in the plasma membrane of *RA-GEF-2*^{-/-} splenocytes (Figure 6E). The chemokine stromal cell-derived factor-1 (SDF-1), like TNF- α , induces Rap1 activation and integrin-mediated cell adhesion in B cells (McLeod *et al.*, 2002). In marked contrast to those induced by TNF- α , SDF-1-dependent Rap1 activation and adhesion to ICAM-1 were totally unaffected by *RA-GEF-2* knockout in splenocytes (Figure 6, F and G). Therefore, M-Ras and *RA-GEF-2* are indeed involved in Rap1 activation specifically downstream of the TNF- α receptor in mouse splenocytes.

DISCUSSION

Signaling for integrin activation in response to a diverse array of cytokines and chemokines has been shown to involve Rap1 (Bos *et al.*, 2001; Kinashi, 2005). Evidence emerging from recent mouse genetics by the use of gene targeting also supports this notion. Primary hematopoietic cells isolated from the spleen and thymus of Rap1A-null mice showed diminished adhesion through LFA-1-ICAM-1 and VLA-4-fibronectin interactions, although these mice were viable and fertile (Duchniewicz *et al.*, 2006). Rap1B is predominantly expressed in platelets, and has been implicated in agonist-induced activation of the platelet-specific integrin $\alpha_{IIb}\beta_3$. Agonist-dependent aggregation of platelets from Rap1B-deficient mice was indeed reduced, and Rap1B-deficient mice were protected from arterial thrombosis (Chrzanowska-Wodnicka *et al.*, 2005). Considering these literatures, Rap1 certainly exerts a pivotal role in lymphocyte adhesion *in vivo* as suggested by previous *in vitro* studies.

However, regulatory mechanisms for Rap1-mediated cell adhesion *in vivo* are only partly understood. Although diverse GEFs for Rap1 exist in mammalian cells, only CalDAG-GEFI (also known as RasGRP2) has been implicated in Rap1-mediated integrin activation through *in vivo* studies. CalDAG-GEFI contains calcium- and diacylglycerol-binding domains, and thus acts downstream of phospholipase C (PLC) enzymes (Kawasaki *et al.*, 1998). Platelets from *CalDAG-GEFI*^{-/-} mice are severely compromised in integrin-dependent aggregation and thrombus formation in response to multiple extracellular factors, including thrombin, thromboxane A₂ and ADP, as a consequence of their inability to signal through CalDAG-GEFI to Rap1 (Crittenden *et al.*, 2004). Therefore, CalDAG-GEFI is thought to be responsible for Rap1 activation downstream of diverse cell surface receptors that trigger platelet aggregation (Crittenden *et al.*, 2004). Additionally, CalDAG-GEFI activates Rap1 in response to the "outside-in" signal evoked by integrin $\alpha_2\beta_1$ (the collagen receptor), leading to cell adhesion mediated by the integrin $\alpha_{IIb}\beta_3$ (Bernardi *et al.*, 2006).

In macrophages, multiple inflammatory mediators, such as lipopolysaccharide, TNF- α and platelet-activating factor, have been shown to activate the β 2 integrin through the activation of Rap1 (Caron *et al.*, 2000). CD31 (PECAM-1)-induced integrin activation also involves Rap1 in T cells (Reedquist *et al.*, 2000). Here, we have shown that M-Ras and RA-GEF-2 are involved in TNF- α -induced Rap1 activation signaling that leads to LFA-1 activation in splenocytes (Figure 7). In contrast, other tested cytokines, such as IL-4, leukemia inhibitory factor, IL-7, IL-10, and IL-15, which trigger integrin-mediated adhesion did not employ M-Ras and RA-GEF-2 as downstream signaling components (data not shown). Similarly, the chemokine SDF-1 does not require M-Ras for the induction of BAF/hLFA-1 cell adhesion (Katagiri *et al.*, 2006). In the thymus, RA-GEF-2 was not necessary for TNF- α -induced, integrin-mediated adhesion, although expression levels of M-Ras and RA-GEF-2 were comparable to those in the spleen (Figure 5F, 6A and 6D). Therefore, Rap1 activation mediated by M-Ras and RA-GEF-2 is thought to have an indispensable role in TNF- α signaling specifically in splenocytes, particularly splenic B cells. The activation of Rap1 independent of M-Ras and RA-GEF-2 may occur upon TNF- α stimulation in the thymus, although the mechanism remains obscure. If M-Ras acts downstream of the TNF- α receptor also in thymocytes, another RA-GEF family member RA-GEF-1 may possibly be involved in TNF- α stimulation of Rap1, because we have recently found that the RA domain of RA-GEF-1 also binds to the GTP-bound form of M-Ras in addition to that of Rap1 (unpublished results). Alternatively, Rap1-independent signaling may have a major role in TNF- α -induced adhesion in thymocytes. Recently, M-Ras null mice were generated, exhibiting no gross morphological defects at both anatomical and histological levels, further supporting the notion that M-Ras may have a highly specific role in TNF- α signaling in splenic B cells (Nuñez Rodriguez *et al.*, 2006.). In *Drosophila* embryonic macrophages, an ortholog of mammalian RA-GEFs, designated Dizzy, is responsible for Rap1-dependent integrin activation, leading to cell adhesion, cell shape change and cell migration (Huelsmann *et al.*, 2006). Thus, RA-GEF-2 and Rap1-mediated integrin regulatory signaling may be conserved in eukaryotes. However, it remains unclear whether an M-Ras ortholog is involved upstream of Dizzy in *Drosophila*.

Upon the binding of its ligand, the TNF- α receptor undergoes homotrimerization, initiating the formation of a large signaling complex consisting of two classes of cytoplasmic adaptors: TNF receptor-associated factors (TRAFs) and death domain-containing molecules such as TRADD and FADD (Locksley *et al.*, 2001; Aggarwal, 2003; Gilmore, 2006). Downstream of these adaptors, the I κ B kinase complex phosphorylates I κ B proteins, leading to their proteasomal degradation. In turn, NF- κ B dimers are released, and translocated to the nucleus, where transcription of an array of genes for adhesion molecules, chemokines, cytokines and enzymes is induced. In addition to the NF- κ B pathway, c-Jun N-terminal kinase and p38 mitogen-activated protein kinase cascades are activated in response to TNF- α stimulation. In contrast to these well characterized signaling pathways, mechanisms for TNF- α -dependent M-Ras activation remain totally unknown. Indeed, no GEF that acts on M-Ras downstream of the TNF- α receptor has heretofore been identified. Intriguingly, conditional knockout of the *TRAF2* gene in splenic B cells was accompanied by significant splenomegaly similarly to *RA-GEF-2* knockout (Grech *et al.*, 2004). Thus, RA-GEF-2 may function downstream of TRAF2 in TNF- α signaling. The role of TRAF2 in the regulation of Rap1-dependent integrin activation will possibly be elucidated in future studies.

The RA-GEF-2-mediated signaling pathway described herein is interesting in that it involves two Ras family GTPases M-Ras and Rap1 in tandem, which are specifically linked by RA-GEF-2. TNF- α -activated M-Ras, which exists almost exclusively in the plasma membrane, recruited RA-GEF-2, which in turn activated Rap1. By this mechanism, a subset of Rap1 that is localized in the plasma membrane, in fact, became activated upon TNF- α stimulation (Figure 4E). Thus, tandemly arranged M-Ras and Rap1 may be primarily responsible for the subcellular region-specific regulation of downstream signaling. Another reported link between M-Ras and Rap1 is a GEF termed Repac/GFR/MR-GEF, which is most abundantly expressed in the brain (Rebhun *et al.*, 2000). In contrast to RA-GEF-2, the ability of Repac/GFR/MR-GEF to promote Rap1 is inhibited by the activated form of M-Ras, and a physiological role of this signaling remains obscure. PLC ϵ , a PLC isoform that harbors both RA and Rap1 GEF domains, also serves as a link between Ras family GTPases. Both Ras and Rap1 bind to the RA domain of PLC ϵ in a GTP-dependent manner, thereby recruiting PLC ϵ

1 predominantly to the plasma membrane and the Golgi apparatus, respectively (Song *et al.*,
2 2001). Rap1 GEF activity of PLC ϵ contributes to prolonged activation of Rap1 specifically in
3 the Golgi apparatus (Jin *et al.*, 2001; Song *et al.*, 2002). Therefore, PLC ϵ also acts as a
4 subcellular region-specific regulator of Rap1.

5 RAPL is identical to NORE1B (Tommasi *et al.*, 2002), which directly binds to M-Ras
6 in a GTP-dependent manner through its RA domain (Ehrhardt *et al.*, 2001). However,
7 mutationally or TNF- α -activated M-Ras did not transduce the signal directly to RAPL, but
8 instead required RA-GEF-2 and Rap1 for the induction of LFA-1-dependent adhesion of both
9 BAF3 cells and primary splenocytes as illustrated in this study. Although the mechanisms
10 remain largely obscure, there may be some difference in the action of Rap1 and M-Ras on
11 RAPL/NORE1B within the cell. Rap1, but not M-Ras, may exist in a microdomain of the
12 plasma membrane, in which RAPL can regulate LFA-1, and thus the binding of RAPL to
13 M-Ras, as observed *in vitro*, may not result in the recruitment of RAPL to the vicinity of
14 LFA-1. As another possibility, the binding to M-Ras, unlike Rap1, may be insufficient for
15 RAPL to become competent for the regulation of downstream molecules, such as Mst1. These
16 possibilities will be examined in future studies.

17

1 **ACKNOWLEDGMENTS**

2

3 We thank Masakazu Hattori and Nagahiro Minato for the SPA-1 cDNA, Jun-ichi Miyazaki
4 for CAG-Cre mice. This investigation was supported by Grants-in-aid for Scientific Research
5 on Priority Areas "Cell and Tissue Disorganization in Cancer" and "Systems Genomics",
6 Scientific Research (B), and the 21st Century COE Research Program "Signaling Mechanisms
7 by Protein Modification Reactions" from the Ministry of Education, Culture, Sports, Science
8 and Technology of Japan.

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18

1 Figure Legends

2

3 **Figure 1.** Induction of LFA-1-mediated cellular aggregation by M-Ras[Q71L]. (A)

4 Expression of endogenous RA-GEF-2, but not M-Ras. Protein extracts from BAF/hLFA-1
5 and BAF/hLFA-1/M-Ras[Q71L] cells were used for immunoblotting to detect endogenous
6 RA-GEF-2 by anti-RA-GEF-2 antibody. Anti-M-Ras and anti-6×His antibodies were used to
7 detect total M-Ras and M-Ras[Q71L], respectively. Anti-actin antibody was used to detect
8 actin as an internal control. (B) M-Ras[Q71L]-induced cellular aggregation. BAF/hLFA-1
9 cells transfected with or without M-Ras[Q71L] were observed by microscopy. Anti-human
10 LFA-1 antibody (MEM-25) was added to the culture medium to examine the involvement of
11 LFA-1 in the cellular aggregation. The result is representative of more than 10 independent
12 clones, which gave equivalent results. Each bar shows 100 μm.

13

14 **Figure 2.** The involvement of RA-GEF-2 and Rap1 in M-Ras[Q71L]-induced cellular

15 aggregation. (A) Inhibition of M-Ras[Q71L]-induced cellular aggregation by downregulation
16 of RA-GEF-2 and Rap1. BAF/hLFA-1/M-Ras[Q71L] cells transfected with siRNAs against
17 control (Control), RA-GEF-2, and Rap1 were observed by microscopy. Each bar shows 100
18 μm. (B) SiRNA-induced reduction of protein expression. After siRNA treatment, protein
19 extracts were prepared and used for immunoblotting by anti-RA-GEF-2 and Rap1 antibodies.
20 (C) Inhibition of M-Ras[Q71L]-induced adhesion to ICAM-1 by downregulation of
21 RA-GEF-2 and Rap1. BAF/hLFA-1/M-Ras[Q71L] cells transfected with siRNAs against
22 control (Control), RA-GEF-2, and Rap1 were examined for adhesion to ICAM-1. Cells bound
23 to ICAM-1 were detected by fluorescence analyzer. Bars represent the average and standard
24 error of three independent experiments performed in triplicate. **, $P < 0.01$. (D) Inhibition of
25 M-Ras[Q71L]-induced cellular aggregation by SPA-1. SPA-1 was not expressed (pLPCX) or
26 expressed (SPA-1) in BAF/hLFA-1/M-Ras[Q71L] cells, and cells were observed by
27 microscopy. Each bar shows 50 μm. (E) The expression of SPA-1. Protein extracts prepared
28 from cells shown in (D) were used to confirm the expression of SPA-1 by immunoblotting
29 using anti-FLAG antibody.

Figure 3. RA-GEF-2 recruitment and Rap1 activation in the plasma membrane by M-Ras[Q71L]. (A) Co-localization of M-Ras[Q71L] and RA-GEF-2 in the plasma membrane. FLAG-M-Ras[Q71L] and endogenous RA-GEF-2 in FLAG-M-Ras[Q71L]-transfected (upper panels) or mock-transfected (lower panel) BAF/hLFA-1 cells were detected by anti-FLAG and anti-RA-GEF-2 antibodies, respectively. Each Bar shows 5 μ m. (B) Rap1 activation and RA-GEF-2 recruitment in the plasma membrane fraction by M-Ras[Q71L]. HA-Rap1 was expressed with or without FLAG-M-Ras[Q71L] in BAF/hLFA-1 cells. Cells were treated with siRNAs against control (Control) and RA-GEF-2, and extracts of the plasma membrane fraction were prepared after serum starvation. The GTP-bound form of HA-Rap1 in the plasma membrane fraction and the total cell lysate was precipitated by the use of GST-RalGDS-RID and detected by immunoblotting using anti-HA antibody. The expression of endogenous RA-GEF-2, FLAG-M-Ras[Q71L] and HA-Rap1 in the plasma membrane fraction was monitored by immunoblotting using anti-RA-GEF-2, anti-FLAG and anti-HA antibodies, respectively. The expression of endogenous RA-GEF-2 and HA-Rap1 in total cell lysates was also monitored by anti-RA-GEF-2 and anti-HA antibodies, respectively.

Figure 4. Induction of LFA-1-mediated adhesion to ICAM-1 via M-Ras by TNF- α . (A) Induction of adhesion to ICAM-1 in M-Ras-expressing BAF/hLFA-1 cells by TNF- α . BAF/hLFA-1 cells transfected with M-Ras, H-Ras, Rap1 or a combination of M-Ras and Rap1 were examined for adhesion to ICAM-1 in the presence and absence of TNF- α . Cells bound to ICAM-1 were detected by fluorescence analyzer. Each bar represents the average and standard error of six independent experiments performed in triplicate. **, $P < 0.01$. (B) Dose dependency of TNF- α on adhesion to ICAM-1. M-Ras-expressing BAF/hLFA-1 cells were treated with TNF- α (0.1 – 100 ng/ml) and examined for adhesion to ICAM-1. (C) Inhibition of TNF- α -induced adhesion to ICAM-1 by anti-human LFA-1 (MEM-25) antibody. After treatment with anti-LFA-1 antibody, M-Ras-expressing BAF/hLFA-1 cells were treated with or without TNF- α and examined for adhesion to ICAM-1. **, $P < 0.01$. (D) Inhibition of adhesion to ICAM-1 by downregulation of RA-GEF-2 and Rap1. M-Ras-expressing

BAF/hLFA-1 cells transfected with siRNAs against control (Control), RA-GEF-2 or Rap1 were treated with or without TNF- α and examined for adhesion to ICAM-1. Data in (B), (C), and (D) are shown as the average and standard error of three independent experiments performed in triplicate. **, $P < 0.01$. (E) The activation of M-Ras and Rap1 and RA-GEF-2 recruitment in the plasma membrane fraction in response to TNF- α . FLAG-M-Ras and HA-Rap1 were co-expressed in BAF/hLFA-1 cells, and extracts of the plasma membrane fraction were prepared after TNF- α treatment. The GTP-bound form of FLAG-M-Ras was precipitated from the extracts by the use of GST-c-Raf-Ras-binding domain and detected by immunoblotting using anti-FLAG antibody. The GTP-bound form of HA-Rap1 in the plasma membrane fraction and the total cell lysate was precipitated by the use of GST-RalGDS-RID and detected by immunoblotting using anti-HA antibody. The amount of endogenous RA-GEF-2, FLAG-M-Ras and HA-Rap1 in the plasma membrane fraction was monitored by immunoblotting using anti-RA-GEF-2, anti-FLAG and anti-HA antibodies, respectively. The amount of endogenous RA-GEF-2 and HA-Rap1 in total cell lysates was also monitored by anti-RA-GEF-2 and anti-HA antibodies, respectively.

Figure 5. Targeted disruption of the mouse *RA-GEF-2* gene. (A) Schematic representation of the wild-type allele (*RA-GEF-2*⁺), the floxed allele (*RA-GEF-2*^{fllox}), and the knockout allele (*RA-GEF-2*⁻). Exons 19 ~ 22 (white rectangles), loxP sites (black triangles), *TK-neo* (neo) and diphtheria toxin A chain cassette (DT-A) are indicated in the targeting construct. The 3' probe for Southern blot analysis, a trio of primers for genotyping by PCR, and restriction enzyme sites (*K*, *KpnI*; *B*, *Bsu36I*; *M*, *MfeI*; *Bsg*, *BsgI*; *S*, *StyI*; *N*, *NotI*) are also shown. (B) Southern blot analysis of the *RA-GEF-2* gene. Genomic DNA was prepared from mouse tails, digested by *Bsu36I*, and hybridized with the 3' external probe. The *RA-GEF-2*⁺ allele generated a 7.7-kb band, and the *RA-GEF-2*^{fllox} allele generated an 8.8-kb band. (C) PCR analysis of the leftmost loxP. Genomic DNA was prepared from mouse tails and analyzed by PCR. Primers #1 and #2 generated a 540-bp band from *RA-GEF-2*⁺ allele, and a 586-bp band from the *RA-GEF-2*^{fllox} allele. (D) PCR-based genotyping of *RA-GEF-2* knockout. Primers #1 and #3 generated a 792-bp band from the *RA-GEF-2*⁺ allele, and a 316-bp band from the *RA-GEF-2*⁻

allele. (E) Comparison of tissue-to-body weight ratios between *RA-GEF-2^{-/-}* (KO) and wild-type (WT) mice. At least seven mice of each genotype were examined at the ages of 8 wk. ***, $P < 0.001$. (F) Immunoblot analysis for RA-GEF-2. Protein extracts were prepared from tissues and used for immunoblotting by anti-RA-GEF-2 antibody.

Figure 6. Impaired LFA-1-mediated adhesion to ICAM-1 in *RA-GEF-2^{-/-}* cells from the spleen, but not the thymus. (A) Comparison of adhesion to ICAM-1 between cells from wild-type and *RA-GEF-2^{-/-}* mice. Thymocytes and splenocytes isolated from wild-type and *RA-GEF-2^{-/-}* mice were left untreated or treated with PMA and TNF- α . After treatment, cells were examined for adhesion to ICAM-1. Anti-mouse LFA-1 antibody was added 30 min before TNF- α treatment. Fold increase of adhesion to ICAM-1 of cells treated with PMA or TNF- α compared to that of untreated cells was shown. The percentages of adhesion to ICAM-1 of untreated cells are as follows; wild-type thymocytes, 7.6 % ; *RA-GEF-2^{-/-}* thymocytes, 7.3 %; wild-type splenocytes, 9.7 %; *RA-GEF-2^{-/-}* splenocytes, 5.9 %. Each bar represents the average and standard error of six independent experiments performed in triplicate. ■, wild-type mice; □, *RA-GEF-2^{-/-}* mice; **, $P < 0.01$. (B) Inhibition of TNF- α -induced adhesion to ICAM-1 by downregulation of M-Ras and RA-GEF-2 in splenocytes. Splenocytes isolated from wild-type and *RA-GEF-2^{-/-}* mice were transfected with siRNAs against control (Control), M-Ras and RA-GEF-2, and left untreated or treated with TNF- α . After treatment, cells were examined for adhesion to ICAM-1. Fold increase of adhesion to ICAM-1 of cells treated with TNF- α compared to that of untreated cells was shown. The percentages of adhesion to ICAM-1 of untreated cells are as follows; wild-type splenocytes, 9.2 %; *RA-GEF-2^{-/-}* splenocytes, 6.3 %. Each bar represents the average and standard error of three independent experiments performed in triplicate. ■, wild-type mice; □, *RA-GEF-2^{-/-}* mice; *, $P < 0.05$. Reduction of the expression of M-Ras and RA-GEF-2 in wild-type (+/+) and *RA-GEF-2^{-/-}* (-/-) splenocytes by specific siRNAs was confirmed by RT-PCR and immunoblotting, respectively. (C) Equal expression of LFA-1 in wild-type and *RA-GEF-2^{-/-}* mice. LFA-1 on the cell surface was stained with fluorescein isothiocyanate-conjugated anti-mouse LFA-1 (M17/4) antibody and analyzed by FACScaliber.

1 White- and gray-colored histograms represent the isotype control, and cells stained with
 2 anti-LFA-1 antibody, respectively. (D) TNF- α -induced adhesion of splenic B and non-B cells.
 3 Splenic B and non-B cells isolated from wild-type and *RA-GEF-2*^{-/-} mice were left untreated
 4 or treated with TNF- α . After treatment, cells were examined for adhesion to ICAM-1. Fold
 5 increase of adhesion to ICAM-1 of TNF- α -treated cells compared to that of untreated cells
 6 was shown. The percentages of adhesion to ICAM-1 of untreated cells are as follows;
 7 wild-type B cells, 5.8 %; *RA-GEF-2*^{-/-} B cells, 4.6 %; wild-type non-B cells, 11.7 %;
 8 *RA-GEF-2*^{-/-} non-B cells, 11.3 %. Each bar represents the average and standard error of four
 9 independent experiments performed in triplicate. ■, wild-type mice; □, *RA-GEF-2*^{-/-} mice;
 10 **, $P < 0.01$. The expression level of M-Ras and RA-GEF-2 in splenic B cells, splenic non-B
 11 cells and thymocytes from wild-type (+/+) and *RA-GEF-2*^{-/-} (-/-) mice was confirmed by
 12 RT-PCR. (E) The activation of Rap1 and RA-GEF-2 recruitment in the plasma membrane
 13 fraction in response to TNF- α . Splenocytes isolated from wild-type and *RA-GEF-2*^{-/-} mice
 14 were transfected with siRNAs against control (Control) and M-Ras, and extracts from the
 15 plasma membrane-enriched fraction were prepared after TNF- α treatment. The GTP-bound
 16 form of endogenous Rap1 in the plasma membrane fraction and the total cell lysate was
 17 precipitated from the extracts by the use of GST-RalGDS-RID and detected by
 18 immunoblotting using anti-Rap1 antibody. The amount of endogenous RA-GEF-2 and Rap1
 19 in the plasma membrane-enriched fraction was monitored by immunoblotting using
 20 anti-RA-GEF-2 and anti-Rap1 antibodies, respectively. The amount of endogenous
 21 RA-GEF-2 and Rap1 in total cell lysates was also monitored by anti-RA-GEF-2 and
 22 anti-Rap1 antibodies, respectively. Inhibition of M-Ras expression by the siRNA in total cell
 23 lysates was confirmed by RT-PCR using primers specific for M-Ras. (F) Adhesion to
 24 ICAM-1 of splenocytes in response to TNF- α and SDF-1. Splenocytes isolated from
 25 wild-type and *RA-GEF-2*^{-/-} mice were treated with TNF- α (10 ng/ml) and SDF-1 (100 ng/ml)
 26 for 30 min and 5 min, respectively, and examined for adhesion to ICAM-1. Fold increase of
 27 adhesion to ICAM-1 of cells treated with TNF- α or SDF-1 compared to that of untreated cells
 28 was shown. The percentages of adhesion to ICAM-1 of untreated cells are as follows;
 29 wild-type splenocytes, 7.7 %; *RA-GEF-2*^{-/-} splenocytes, 5.3 %. Each bar represents the

1 average and standard error of three independent experiments performed in triplicate. ■,
2 wild-type mice; □, *RA-GEF-2*^{-/-} mice; **, $P < 0.01$. (G) The activation of Rap1 in
3 splenocytes treated with TNF- α and SDF-1. Splenocytes isolated from wild-type and
4 *RA-GEF-2*^{-/-} mice were treated with TNF- α and SDF-1 as in Fig. 6F. After treatment, the
5 GTP-bound form of endogenous Rap1 was precipitated from total cell lysates by the use of
6 GST-RalGDS-RID and detected by immunoblotting using anti-Rap1 antibody. The amount of
7 endogenous Rap1 in total cell lysates was monitored by anti-Rap1 antibody.

8
9 **Figure 7.** Presumptive signaling pathway. Upon binding to the receptor (1), TNF- α activates
10 M-Ras (2). The activated M-Ras recruits RA-GEF-2 from the cytoplasm to the plasma
11 membrane (3), and RA-GEF-2 activates Rap1 in the plasma membrane (4) to induce LFA-1
12 activation (5) and adherence to ICAM-1 (6).

Figure 1

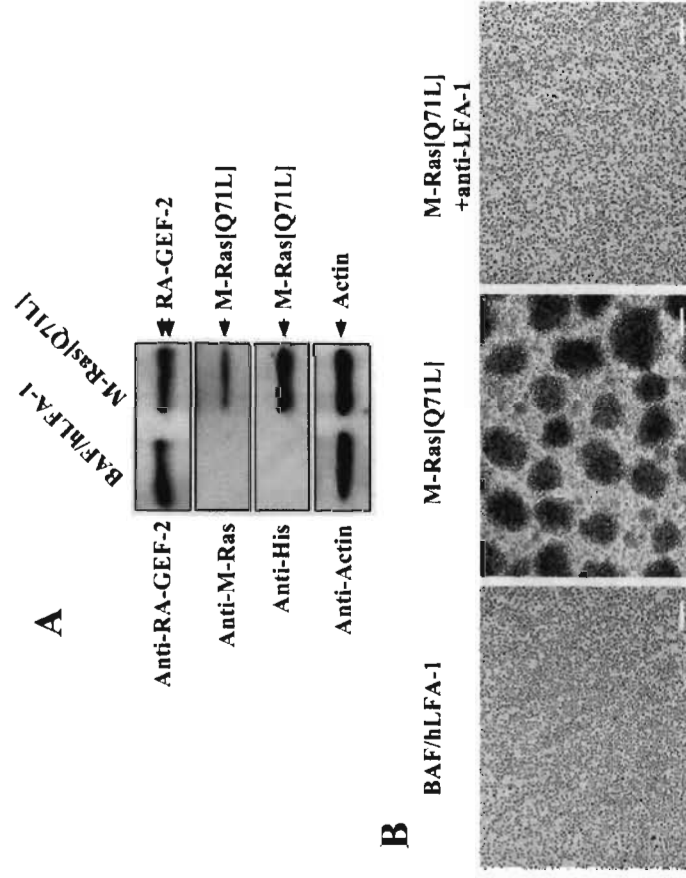


Figure 2

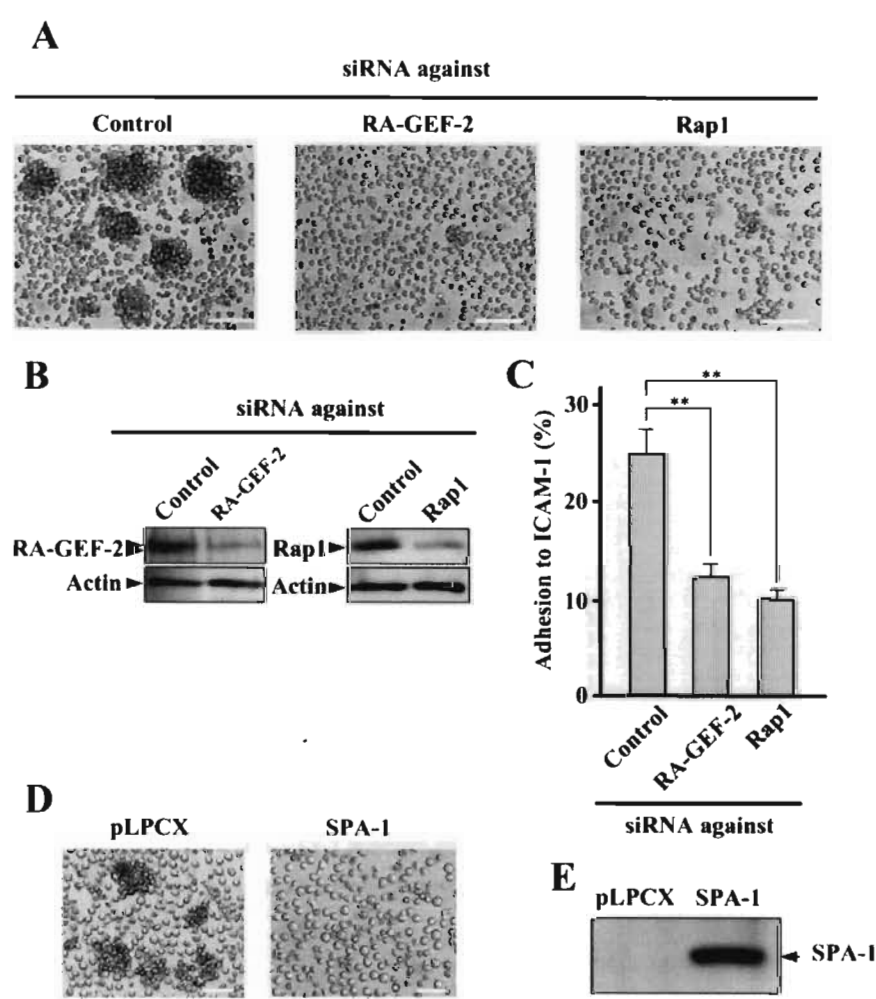


Figure 3

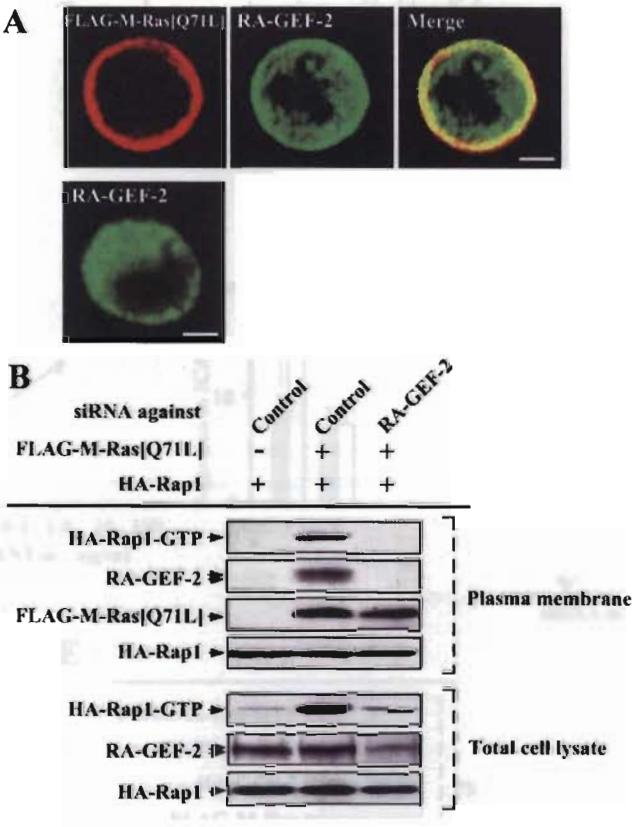


Figure 4

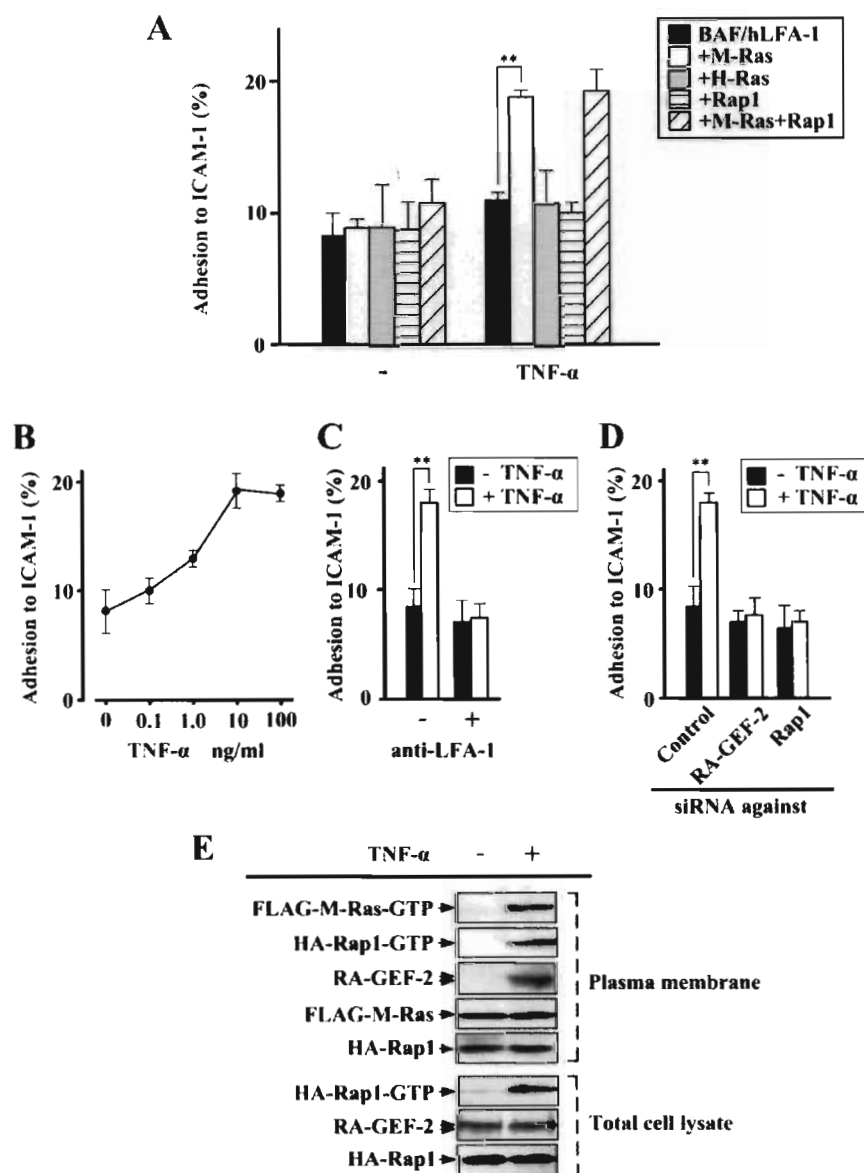


Figure 5

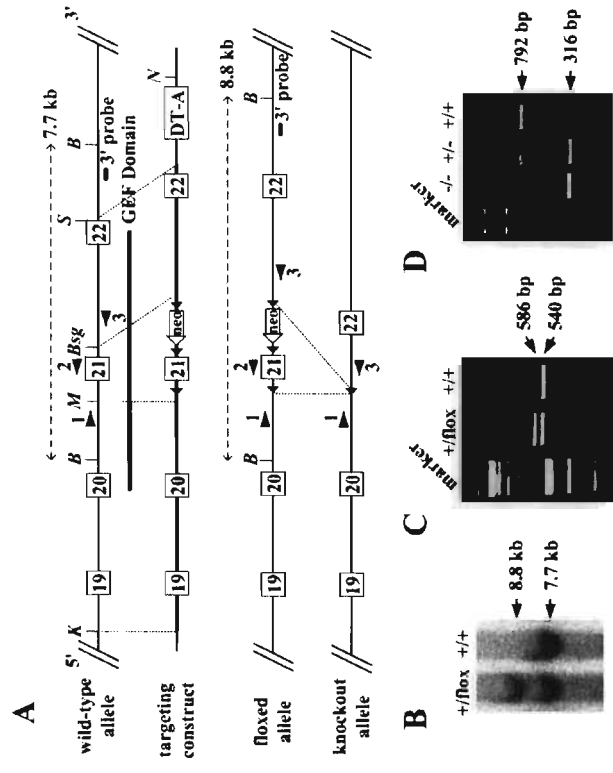
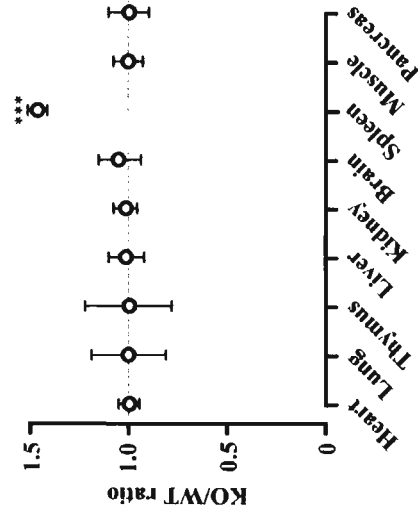


Figure 5

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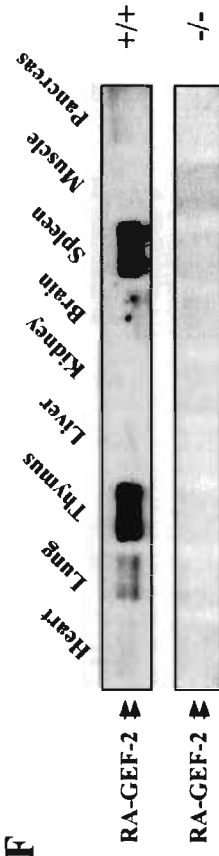


Figure 6

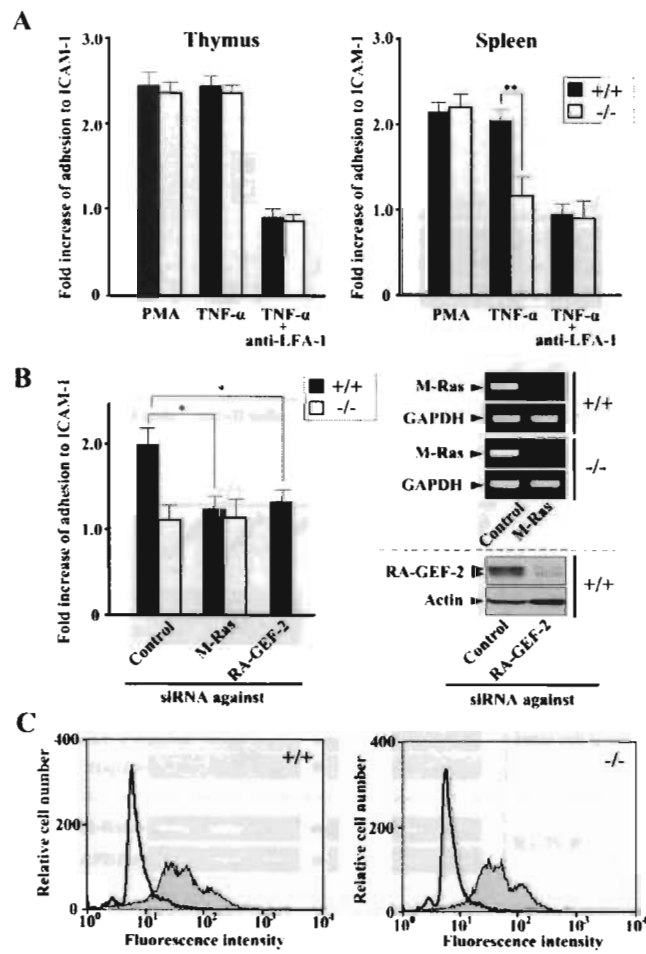


Figure 6

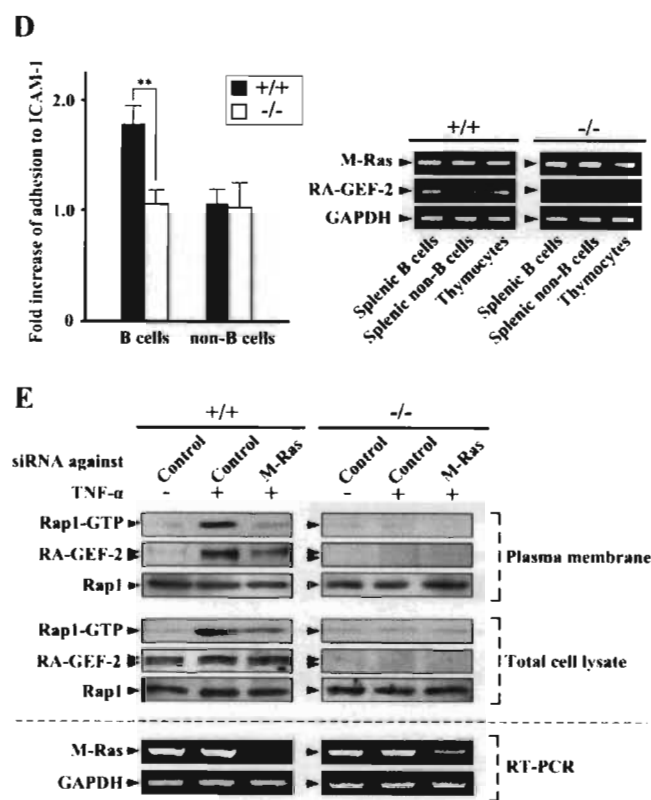


Figure 6

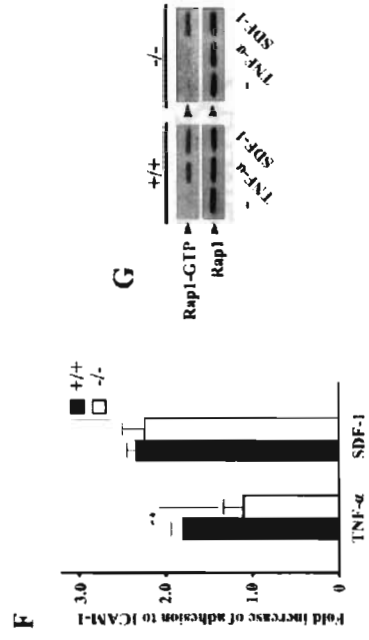


Figure 7

