Abstract. As an atypical member of the Rab family, Rab24 has several attributes distinguishing this protein from the other members. Based on the yeast two-hybrid system, interaction between human RAB24 and two proteins, cyclophilin A (CyP-A) and gamma-aminobutyric acid type A receptor-associated protein (GABARAP), was detected and identified in COS-7 cells. Interestingly, RAB24 is predominantly localized in the nuclei of COS-7 cells, which is different from previous reports using other cell lines. RAB24 (D123I) can trigger the accumulation of intracellular inclusions, with small quantities of intranuclear inclusions in some cells. The GTPase activity of RAB24 and its two mutants was detected.

Introduction

Rab24 is an atypical Rab protein whose function is currently unknown. Subcellular localization experiments showed that Rab24 is distributed in the endoplasmic reticulum/cis-Golgi region and on late endosomal structures, and this subcellular distribution pattern of Rab24 may indicate its involvement in autophagy-related processes (1,2). Rab24 presents unusual characteristics, including low intrinsic GTPase activity, and is inefficiently prenylated when compared with other Rab proteins (3). Another study also suggested that Rab24 might play a unique role in the degradation of misfolded cellular proteins or trafficking of proteins to the nuclear envelope (4). It has been reported that GFP-Rab24 is associated with autophagosomes in amino-acid starved cells; furthermore, Rab24 could be colocalized with LC3 (microtubule-associated protein light chain 3, a mammalian homolog of the yeast protein Apg8/Aut7), which suggested that Rab24 protein may be involved in the autophagic pathway since LC3 is an essential gene for autophagy (2). Although a possible role of Rab24 in autophagy has been suggested, its specific physiological function in mammalian cells remains to be defined. It should be mentioned that all of these study results were based on Rab24 from mice. For human RAB24, which varies by two amino acids (localized at the nonconservative region) and shares 98% identity with Rab24, there is no functional study, although it has been identified and characterized (5). Based on the yeast two-hybrid system, three proteins interacting with RAB24, human GABARAP [γ-aminobutyric-acid-type-A (GABA)-receptor-associated protein], cyclophilin A (CyP-A) and TUBA6 (human tubulin alpha 6), were detected, and the former two were individually cotransfected with RAB24 into COS-7 cells for further identification. The findings on the association between RAB24 and GABARAP and CyP-A in mammalian cells are helpful for further functional study on these three proteins.

It was surprising to find that human RAB24 is localized predominantly in the nuclei of COS-7 cells, which is different from previous reports that Rab24 is only distributed in the cytoplasm of other cell lines. RAB24 has low GTPase activity, and two mutants of RAB24 [RAB24 (Y17A) and RAB24 (S67Q)] produced a substantial increase in RAB24 GTPase activity, which has been suggested previously using Rab24 from mice.

Materials and methods

Strains, media, plasmids and cell line. The E. coli strains DH5α and BL21 (Rosetta) were grown in Luria Broth medium in the presence of antibiotics, as required for different plasmids, pLexA, pEGFP-C1 and pDsRed-N1 plasmids were purchased from Clontech, and pGEX-4T1 was from Amersham Pharmacia. COS-7 (African green monkey kidney cells) was purchased from the Chinese Academy of Science, cultured and stored in our laboratory. Media and reagents for cell culture were purchased from Invitrogen.
Expression constructs. The cDNA encoding RAB24 was obtained by PCR amplification of the recombinant clone constructed by our laboratory and cloned into various plasmids, including pLexA, pEGFP-C1 and pGEX-4T1. The RAB24 cDNA was altered by overlap-extension PCR, using the appropriate mutator oligonucleotides (listed below) to introduce amino acid substitutions at D123 (D123I, pEGFP-C1 and pGEX-4T1), Y17 (Y17A, pGEX-4T1), and S67 (S67Q, pGEX-4T1). ORFs of CyP-A and GABARAP were obtained using PCR amplification for human multiple tissue cDNA, and cloned into plasmid pDsRed-N1, respectively. All primers and restriction enzymes of expression constructs are: pLexA-RAB24 sense (EcoRI) 5'-CGGAATTCGCGAAGACGTACGATTATCTCTTC-3', and antisense (XhoI) 5'-CCGCTCGAGAAGTAGCGAGCAACGAAAGAAACTG-3'; pEGFP-C1-RAB24 sense (EcoRI) 5'-CGGGATCCGTGATGACAACAGCTGTAGAAGTAGG-3'; pGEX4T1-RAB24 sense (BamHI) 5'-CGGGATCCGCG GGCAGCGCGTGGACGTCAA-3', and antisense (XhoI) 5'-CCGCTCGAGAAGTAGCGAGCAACGAAAGAAACTG-3'; pDsRed-N1-CyP A sense (HindIII) 5'-CCAAGCTTACCA TGGAGTTCGTGTACAAAGAAGC-3', and antisense (BamHI) 5'-CGGGATCCCGCAGACCGTAGACACTTTCGTCAC-3'; pDsRed-N1-GABARAP sense (EcoRI) 5'-CGGAATTCTGACCATGGTCAACCCCACCGTGTTCTTC-3', and antisense (BamHI) 5'-CGGGATCCCGTTCGAGTTGTCCACAGTCAATG-3'.

Figure 1. Subcellular distribution of human GABARAP and CyP-A in cultured COS-7 cells. The 2 μg constructs of pDsRed-N1/CyP-A or pDsRed-N1/GABARAP were transfected individually into COS-7 with 3 μl Lipofectamine 2000. The nucleus was counterstained with DAPI and viewed using an R2100 Confocal Microscope with appropriate filters (543 nm). (A) Subcellular localization of DsRed-GABARAP, predominantly distributed in the cytoplasm. (B) Subcellular localization of DsRed-CyP-A, distributed only in the cytoplasm, often encircling the nucleus.

Figure 2. Subcellular distribution of human RAB24 and RAB24 (D123I) in cultured COS-7 cells. The 2 μg constructs of pEGFP-C1/ RAB24 or pEGFP-C1/ RAB24 (D123I) were transfected individually into COS-7. The samples were viewed using an R2100 Confocal Microscope with appropriate filters (488 nm). (A) Subcellular localization of GFP-RAB24, predominantly distributed in the nucleus, with small amounts at the perinuclear position. The distribution pattern would be quite different with other cell lines. (B) Subcellular localization of GFP-RAB24 (D123I) collectively form intracellular inclusions, mostly encircling the nucleus (possibly in the endoplasmic reticulum) and without the ability to gain passage. In some cells, intranuclear inclusions were visible, but the nuclear envelope is disrupted (two lower panels).
Cell culture and liposome-mediated transfection. COS-7 cells were grown in Dulbecco's modified Eagle medium containing 10% FBS and maintained at 37°C in a 5% CO₂ atmosphere. On the day before transfection, cells were plated on coverslips in the 6-well tissue culture plates. Cells were transfected at 60-80% confluence with the indicated expression vectors using Lipofectamine 2000 (Invitrogen) according the manufacturer's instructions. Following 32-48 h transfection, cells were washed 3 times with 1X PBS, fixed with 4% paraformaldehyde for 15 min washed 3 times with 1X PBS, counterstained with 4',6'-diamidino-2-phenylindole for 10 min, and washed 2 times with ddH₂O. The samples were then viewed using an R2100 Confocal Microscope (Bio-Rad) with the appropriate filters, and the photos were captured with Leica DC viewer software. The overlays were processed with Photoshop 6.0.

Figure 3. Colocalization of RAB24 and CYP-A or RABARAP in COS-7 cells. The 2 μg constructs of pDsRed-N1/CyP-A or pDsRed-N1/GABARAP and 2 μg pEGFP-C1/RAB24 were co-transfected individually into COS-7 cells. (A) Colocalization of RAB24 and CyP-A is always distributed around the nucleus in large dots. GFP-RAB24 is also distributed in the nucleus. (B) Colocalization of RAB24 and RABARAP was often distributed in the cytoplasm near the nucleus (two upper panels). Surprisingly, DsRed-GABARAP entered the nucleus and was colocalized with GFP-RAB24 in some cells (low panel).

Yeast two-hybrid system. The MATCHMAKER LexA two-hybrid system and human fetal brain MATCHMAKER LexA cDNA libraries were purchased from Clontech. The yeast two-hybridization was performed according to the manual supplied by the manufacturer, and a sequential transformation method was adopted. All positive clones were classified by yeast clone hybridization and the length of PCR product using the AD fusion site-specific primer (sense, 5'-CCA GCCTCTTGCTGAGTGGAGATG-3'; antisense, 5'-CGT AAATTTCTGGCAAGGTAGACAAG-3'). The independent clones were further verified with a yeast mating test. PCR products amplified from positive clones were sequenced on an ABI PRISM™ 377 DNA sequencer (Perking-Elmer), and the sequencing results were directly submitted to the NCBI website (http://www.ncbi.nlm.nih.gov/blast) to perform BLAST analysis.

Identification for the interacting proteins of RAB24

Yeast mating test. The yeast mating test was used to verify the interaction between RAB24 and CyP-A, GABARAP and TUBA6. All yeast-mating procedures were performed according to the recommendations of the manufacturer (Clontech). These results are not shown.
Colocalization of the interacting proteins and RAB24 in vivo. The 2 μg constructs of pDs Red-N1/CyP-A or pDs Red-N1/GABARAP and 2 μg pEGFP/RAB24 were mixed in 250 μl DMEM (no serum), and 5 μl Lipofectamine 2000 was also mixed gently in 250 μl DMEM (no serum). After 5 min, they were mixed together for 20 min. These mixtures were then co-transfected individually into COS-7 cells and cultured on coverslips in 6-well plates.

GTPase activity detection for human RAB24 and its mutants. Protein expression and purification. E. coli strain BL21 (Rosseta) containing the expression plasmids was grown overnight in LB medium containing 100 mg/l ampicillin. Expression of GST-tagged recombinant protein was induced at OD560 of 0.4-0.6 with 1 mM IPTG for 4 h at 24˚C, then the cells were harvested by centrifugation at 5000 rpm for 15 min at 4˚C. The cells were resuspended in 1X PBS with 1 mM PMSF, then lysed by sonication for 10 min in an ice bath. The lysate was cleared by centrifugation at 14000 rpm for 15 min at 4˚C. After centrifugation, the supernatant was loaded to a glutathione sepharose 4B column. GST fusion proteins were eluted and analyzed by SDS-PAGE.

GTPase activity assay for RAB24 and its mutants. An EnzChek Phosphate Assay kit was purchased from Molecular Probes and utilized according to the manufacturer's protocol. The reaction components are: 50 μl 20X reaction buffer, 200 μl MESG substrate solution, 10 μl PNP (1U), 200 μM GTP, and 5 μM GST-RAB24/GST-RAB24 (Y17A)/GST-RAB24 (S67Q), adding ddH2O to 1 ml (total volume).

Results

Subcellular distribution of RABARAP, CyP-A, RAB24 and RAB24 (D123I) in COS-7. RABARAP and CyP-A are all localized in the cytoplasm of COS-7 cells (Fig. 1). The former also appears to be distributed throughout the cytoplasm, with small amounts in the nucleus; but the latter appears to show a perinuclear reticular localization that often encircles the nucleus (Fig. 1B), which is similar to the distribution pattern of RAB24 (D123I) (Fig. 2B). It is no surprise that subcellular distribution of these two proteins is consistent with previous study results obtained by using other cell lines of human and mouse origin.

As shown in Fig. 2A, RAB24 is predominantly localized in the nucleus except for small amounts distributed in the cytoplasm near the nucleus. However, the mutant RAB24 (D123I) accumulated intracellular inclusions that were always targeted to the perinuclear reticulum in most COS-7 cells (Fig. 2B).

Colocalization of RAB24 and RABARAP or CyP-A in COS-7 cells. RAB24 is well colocalized with CyP-A in a perinuclear position (Fig. 3A). For another interacting protein, GABARAP, there are two kinds of subcellular colocalization with RAB24 (Fig. 3B). In some COS-7 cells, RAB24 and GABARAP can colocalize in the cytoplasm (Fig. 3B, upper two panels); in others, however, their colocalization can also be seen in the nucleus (Fig. 3B, lower panel).

GTPase activity detection of RAB24 and its mutants. Consistent with previous reports, the GTPase activity of
human RAB24 is low (Fig. 4B). The amino acid substitution at two different positions (Y17A and S67Q) resulted in a dramatic increase of GTPase activity of RAB24 (Fig. 4C and D). However, there are no results as yet on GTPase activity of the mutant RAB24 (D123I), which formed inclusion bodies when expressed in E. coli (Fig. 4A).

Discussion

It has been recognized that Rab24 is distributed in the cytoplasm, especially in the endoplasmic reticulum/cis-Golgi region and on late endosomal structures, with cell lines of BHK, HeLa, MDCK, NmuLi, CHO, HEK 293 and NIH 3T3 mouse fibroblasts. Our study showed an unexpected and surprising result that RAB24 is predominantly localized in the cytoplasm near the nucleus (Fig. 2A). It was reported that HIV could use various elements of endocytic/trafficking machinery mediated by Rab proteins to enter and successfully infect the host cells (6). It is known that RAB24 contains a unique arginine-rich insert, 126EEDRRRRR133 (3), whose signal for RAB24 in COS-7 cells remains to be confirmed, but we are convinced that this discrepancy of two distribution patterns suggested RAB24 must play a different role in human and monkey cells.

It is of interest that in most COS-7 cells the mutant RAB24 (D123I) accumulated intracellular inclusions that always target the perinuclear reticulum (Fig. 2B). However, in other cell lines (HEK 293 and NIH 3T3 mouse fibroblasts), it has been reported that the mutant Rab24 GTPase accumulated in nuclear inclusions that disrupted the integrity of the nuclear envelope and appeared to sequester ubiquitin, Hsp70, importin-β and GFP-coupled glucocorticoid receptor, which is not found in other Rab GTPases with a similar GXXXVGK (S/T) motif, and S67, falling beside the DXXG positions (Y17A and S67Q) result in a dramatic increase of GTPase activity of RAB24. Y17, falling within the DXXG motif, is a member of the cyclophilin family, whose functions include protein folding, transport and signaling (11). These immunophilins participate in axonal transport and synaptic vesicle assembly, and may play a role in neuroprotection against abnormal protein aggregation. In other words, they are involved in nervous system degeneration and regeneration (12-14). Other studies showed that CyP-A associates in vitro and in vivo with the dynein/dynactin motor protein complex (including tubulin), which suggests that CyP-A may perform a general function related to the binding of cargo for retrograde movement along microtubules (15). From Fig. 3A, we can see that RAB24 and CyP-A are well colocalized in the perinuclear position, which implied that they may cooperate to function in a signaling pathway at this position or elsewhere. More study results showed that CyP-A plays an essential role in the HIV-1 life cycle, including early and late events in HIV-1 replication (16-18). Moreover, CyP-A has different and even converse effects on HIV-1 infection and replication when expressed in human and monkey cells, i.e., interaction between the capsid protein and host CyP-A protects HIV-1 from restriction in human cells, but is essential for maximal restriction in simian or monkey cells (19-22). Whether or not RAB24 is involved with CyP-A in this process, and the different subcellular distribution of RAB24 is related to the different response to HIV-1 in human and monkey, remains an enigma and needs to be elucidated.

Another interacting protein colocalized with RAB24 (Fig. 3B), GABARAP is a multifunctional protein and could interact with a myriad of binding partners, including the γ2 subunit of the GABAA receptor, tubulin and microtubules, N-ethyl maleimide sensitive factor (NSF), gephyrin and the transferin receptor (23-29). Surprisingly, RAB24 and GABARAP can colocalize in the cytoplasm (Fig. 3B, upper two panels) or nucleus (Fig. 3B, lower panel). A possible explanation is that some GABARAP could translocate into the nucleus by association with RAB24 in COS-7 cells. It is unknown as to whether this is the case or not. Previous study results showed that RAB24 is colocalized with LC3, a family of yeast Atg8-related proteins (the other two families being GABARAP and GATE16). It has been reported that three families of mammalian Atg8-related proteins were able to localize to the autophagosomal membrane depending on their form-II formation (30). LC3 has been used to study autophagy in neurodegenerative and neuromuscular diseases, tumorigenesis, and bacterial and viral infections.

All results from the previous and present studies have strengthened the hypothesis that RAB24 (with CyP-A and/or GABARAP) is engaged in autophagy and neurodegenerative disorders to some extent, since these two cell processes are confirmed to have a significant relationship (31-37). RAB24 has several attributes that are different from other members of the Rab family, one being low GTPase activity, which is known from non-human study models such as the mouse. It was shown that this GTPase activity is attributed to human RAB24, which is consistent to previous reports (Fig. 4B). The amino acid substitution at two different positions (Y17A and S67Q) result in a dramatic increase of GTPase activity of RAB24 (Fig. 4C and D), which suggested that these two amino acid sites could contribute to the low GTPase activity of RAB24. Y17, falling within the GXXXVGK (S/T) motif, and S67, falling beside the DXXG motif, appear to be unique compared with other Rab family members (3). From these results, it can be concluded that these two amino acid sites must have negative effects on the GTPase activity of RAB24.
The mutant RAB24 (D123I) is obviously different from two other mutants since it shifted to form inclusions instead of soluble patterns when expressed in *E. coli* (Fig. 4D). This phenomenon can also be observed when RAB24 (D123I) is expressed in mammalian cells. Therefore, this mutation site (D123I) is able to induce large changes of conformation of RAB24 and influence the function of this protein, resulting in abnormal cellular responses.

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References