Functional and biochemical characterization of a T cell-associated anti-apoptotic protein, GIMAP6

Ching-Huang Ho and Shih-Feng Tsai

From the Department of Life Sciences and Institute of Genome Sciences, National Yang-Ming University, Taipei 112, Taiwan and the Institute of Molecular and Genomic Medicine, National Health Research Institutes, Zhunan, Miaoli 350, Taiwan

GTPases of immunity-associated proteins (GIMAPs) are expressed in lymphocytes and regulate survival/death signaling and cell development within the immune system. We found that human GIMAP6 is expressed primarily in T cell lines. By sorting human peripheral blood mononuclear cells and performing quantitative RT-PCR, GIMAP6 was found to be expressed in CD3+ cells. In Jurkat cells that had been knocked down for GIMAP6, treatment with hydrogen peroxide, FasL, or okadaic acid significantly increased cell death/apoptosis. Exogenous expression of GMAP6 protected Huh-7 cells from apoptosis, suggesting that GIMAP6 is an anti-apoptotic protein. Furthermore, knockdown of GIMAP6 not only rendered Jurkat cells sensitive to apoptosis but also accelerated T cell activation under phorbol 12-myristate 13-acetate/ionomycin treatment conditions. Using this experimental system, we also observed a down-regulation of p65 phosphorylation (Ser-536) in GIMAP6 knockdown cells, indicating that GIMAP6 might display anti-apoptotic function through NF-κB activation. The conclusion from the study on cultured T cells was corroborated by the analysis of primary CD3+ T cells, showing that specific knockdown of GIMAP6 led to enhancement of phosphol 12-myristate 13-acetate/ionomycin-mediated activation signals. To characterize the biochemical properties of GIMAP6, we purified the recombinant GIMAP6 to homogeneity and revealed that GIMAP6 had ATPase as well as GTPase activity. We further demonstrated that the hydrolysis activity of GIMAP6 was not essential for its anti-apoptotic function in Huh-7 cells. Combining the expression data, biochemical properties, and cellular features, we conclude that GIMAP6 plays a role in modulating immune function and that it does this by controlling cell death and the activation of T cells.

This work was supported by Intramural Grant MG-105-PP-01 of the National Health Research Institutes, Taiwan, and two extramural grants from Taiwan’s Ministry of Health and Welfare (106-0324-10-10-05 and Advance Medical Plan 106-0324-01-10-07). The authors declare that they have no conflicts of interest with the contents of this article. This article contains supplemental Figs. S1–S4.

1 To whom correspondence should be addressed. Tel.: 886-2-26534401#35310; Fax: 886-2-28200552; E-mail: petsai@nhri.org.tw.

2 The abbreviations used are: GIMAP, GTPase of immunity-associated proteins; IAN, immune-associated nucleotide-binding protein; NSCLC, non-small-cell lung cancer; PBMC, peripheral blood mononuclear cell; Ctrl, control, AnV, Annexin-V; PI, propidium iodide; OA, okadaic acid; PMA, phorbol 12-myristate 13-acetate; AICD, activation-induced cell death; P-loop, phosphate-binding loop; TLC, thin-layer chromatography; PE, phycoerythrin; P/I, phorbol 12-myristate 13-acetate/ionomycin; PARP, poly(ADP-ribose) polymerase.
in cell-based assays. GIMAP5 is able to function as an anti-apoptotic effector by protecting Jurkat T cells against okadaic acid and γ radiation (13). Similarly, mouse Gimap8 protects NIH 3T3 and CHO-K1 cells from anisomycin-induced apoptosis (14).

T cell activation and differentiation are important steps in the development of adaptive immunity. Naïve T cells require at least two signals, CD3 and CD28, for activation. Following external stimulation, the secondary messenger 1,2-diacylglycerol and inositol triphosphate are released, and these then activate a complicated series of multiple signaling cascades that orchestrate nuclear factor of activated T cells (NFAT), nuclear factor κB (NF-κB), and activator protein 1 (AP-1) to stimulate IL-2 production. Mammalian GIMAPs are known to be expressed predominantly within lymphomyeloid tissues, indicating a role in lymphocyte functionality and homeostasis (2, 3). Some GIMAPs have been reported to function as key regulators during CD4+ T lymphocytes differentiation. It should be noted that GIMAP1 and GIMAP4 have been found to be differentially regulated during Th1 and Th2 differentiation (15). In 2015, a study reported that GIMAP4 affects T cell differentiation via the regulation of interferon γ secretion (16). Previously, we discovered that the RNA expression of the GIMAP family genes, but not of other neighboring non-GIMAP genes, as detected by a microarray analysis, is uniformly lower in human lung tumor tissue samples (17). This finding was confirmed by quantitative PCR assays using a total of 20 non-small-cell lung carcinoma (NSCLC) samples. We also observed that the expression level of GIMAP genes is higher in adjacent non-tumor tissues compared with lung cancer tumor tissue. This finding indicates that GIMAP genes might be involved in either the pathogenesis of NSCLC or in the immune reaction to NSCLC. Among the GIMAPs, GIMAP6 and GIMAP8 showed the highest normal/tumor expression ratio in lung cancer paired samples (17). Compared with GIMAP8, little is known about GIMAP6 regarding its biological function. In this study, we have focused on GIMAP6 and set out to investigate its function in T cells and to characterize its biochemical properties.

Results

Tissue specificity of GIMAP6 expression

To determine the expression pattern of GIMAP6 in different tissues, we analyzed various cell lines from the hematopoietic system, including T lymphocytes (Jurkat), promyeloblasts (HL-60), monocytes (U-937), lymphoblasts (K-562), B lymphocytes (Raji and Ramos), and various cancer derived cell lines of the colon (HCT-15), kidney (293T), liver (Huh-7), and lungs (A549 and H1299). Immunoblotting and quantitative PCR analysis showed that GIMAP6 was detected primarily in T lymphocytes (Fig. 1, A and B). This contrasts with the expression pattern of mouse Gimap6, which is seen equally in T lymphocytes and B lymphocytes (1). To confirm this finding, we isolated the CD3−, CD3+/CD4+, and CD3+/CD8+ fractions of PBMCs (supplemental Fig. S1) and carried out quantitative PCR assays to determine the expression level of GIMAP6 in these different lymphocyte subsets. As shown in Fig. 1C, both CD3+ /CD4+ and CD3+/CD8+ cells showed expression of GIMAP6, whereas CD3− cells were negative. Because CD3− cells within PBMCs are mostly composed of monocytes and B lymphocytes, we concluded that GIMAP6 in humans is predominantly expressed in CD3+ T lymphocytes.

Subcellular localization of GIMAP6 protein

To determine the distribution of GIMAP6 protein in different cellular compartments, subcellular fractionation was performed on Jurkat T lymphocytes to separate the nucleus, the cytosol, and the heavy membrane fraction, with the last containing the mitochondria. Specific antibodies (anti-H3, anti-GAPDH, and anti-PHB1) were used as markers to confirm the purity of the fractions from the different compartments. Immunoblot analysis showed that endogenous GIMAP6 was present primarily in the cytosol of Jurkat T lymphocytes (Fig. 1D). This result is consistent with the results obtained by confocal microscopy, which show that GIMAP6 co-localizes with the cytosolic marker GAPDH (Fig. 1E). Consistent with the result obtained from Jurkat cells, immunofluorescence microscopy of primary CD3+ T cells showed that GIMAP6 co-localized with the cytosolic marker GAPDH (Fig. 1F).

GIMAP6 displays an anti-apoptotic function

Previous studies have reported that members of the GIMAP family play a role in either cell death or cell survival (1, 12–14, 18). Thus we investigated the possible function of GIMAP6 in the programmed death of T cells by comparing wild-type control cells with GIMAP6-deficient cells. Stable knockdown cells (KD1 and KD2) were generated using anti-GIMAP6 shRNA, whereas control cells (Ctl) were prepared using the plKO.1 empty vector plasmid. After single colony selection, the relative -fold of GIMAP6 expression was determined by immunoblot analysis (Fig. 2A). To reveal the extent to which cell apoptosis/death was delayed by GIMAP6, an H2O2-mediated, multiple time point apoptosis assay was performed. Various Jurkat-derived cell lines (Ctl, KD1, and KD2) were treated with 100 μM H2O2 for 4, 8, 24, and 30 h to induce cell apoptosis/death (Fig. 2, B and C). The apoptosis signal, phosphatidylserine exposure, was then measured by flow cytometry using a BD Annexin-V apoptosis detection kit. With Annexin-V (AnV)/propidium iodide (PI) staining, GIMAP6 displayed an apparent anti-apoptotic effect, as indicated by the difference between the control and knockdown lines. In these experiments, the apoptotic group (AnV+/PI−) showed a greater increasing trend in knockdown cells compared with control cells after treatment with H2O2 for 4 h, and this difference became more remarkable at 8 h. Compared with knockdown cells, the increase in apoptotic cells (AnV+/PI−) among control cells was postponed until 24 h after treatment with H2O2. In parallel, the cell distribution of knockdown cells, upon being stained with AnV/PI, shifted from AnV+/PI− to AnV+/PI+ after treatment with H2O2 for 24 h. At 30 h, the AnV+/PI− group continued to decrease, whereas the AnV+/PI+ group remained on the increase. This shift in the cell distribution indicates that the cells were progressing from an apoptotic phase to a dead phase. The results also show that GIMAP6 stable knockdown cells are not only more prone to apoptosis than control cells but also that they progress toward the dead phase earlier. Taken together, it can be concluded that GIMAP6 protects cells from apoptosis by
delaying apoptosis progress when a H$_2$O$_2$-mediated apoptosis assay is carried out with Jurkat T cells.

To further investigate the anti-apoptotic effect of GIMAP6, a number of apoptosis-inducing agents was used to perform sets of similar experiments with appropriately adjusted incubation times. Specifically, incubation was carried out for 6 h with 50 nM okadaic acid (OA) or for 3 h with 8 ng/ml FasL. These results were compared with 8-h treatment with 100 nM H$_2$O$_2$ (Fig. 2, D and E). Similar to the H$_2$O$_2$-mediated apoptosis assay, GIMAP6 also showed anti-apoptotic effects with OA- and FasL-induced apoptosis. To determine whether this finding is related to downstream activation of apoptosis, an alternative assay was performed. Treated and untreated control cells and GIMAP6 stable knockdown cells were immunoblotted using anti-PARP antibodies. Consistently, after apoptotic induction, the knockdown groups had a higher ratio of the cleaved form of PARP compared with the control group (Fig. 2 F). Taken together, these findings indicate that GIMAP6 functions as an anti-apoptotic effector in Jurkat T lymphocytes.

We also used an overexpression approach to demonstrate the anti-apoptotic function of GIMAP6. After stable transfection of the cell line Huh-7, which does not express GIMAP6, single colonies were picked, and the exogenous expression of GIMAP6 was confirmed by immunoblotting (Fig. 3 A). Cells
expressing no (Neg), a low level (TF1), and a high level (TF2) of GIMAP6 were then incubated with various cell death inducers; namely, 50 nM okadaic acid, 50 μM etoposide, 10 ng/ml FasL, and 200 μM H₂O₂. The results were compared with control cells (2% DMSO) after 24 h. It was found that Huh-7 cells show a high resistance to etoposide, FasL, and H₂O₂, and therefore okadaic acid was used as the apoptosis inducer during subsequent experiments (data not shown). An apoptotic marker, the cleaved form of caspase-3, was measured by high-throughput immunofluorescence (Fig. 3B). Under these conditions, protection against okadaic acid-induced apoptosis was observed in cells overexpressing GIMAP6, both TF1 and TF2 (Fig. 3C). To
confirm that this anti-apoptotic effect in the presence of okadaic acid was attributed to GIMAP6, we knocked down GIMAP6 using shRNA treatment in Huh-7 cells expressing large amounts of GIMAP6 (TF2). The knockdown was able to partially restore the sensitivity of these cells to the inducing agent. An increase in the cleaved form of caspase-3 was also observed in the knockdown cell lines (2K2 and 2K3) after okadaic acid treatment compared with the high-level GIMAP6-expressing cell lines (TF2 and 2K1) (Fig. 3D). Thus, it can be concluded that GIMAP6 displays an anti-apoptotic effect during these transfection assays and that GIMAP6 is sufficient to protect non-lymphoid Huh-7 cells against okadaic acid-induced apoptosis.

GIMAP6 has GTP/ATP hydrolysis activity

All GIMAP family members are grouped together within the P-loop NTPase superfamily (NCBI Position-specific Scoring Matrix (PSSM) ID 214148) because of the presence of the AIG1 GTP-binding domain (NCBI Conserved Domain Database (CDD) cd01852) (19). A previous study has revealed that some GIMAP family members are able to bind GTP, whereas others have GTP hydrolysis activity (5, 6, 8). Therefore, the three-dimensional structure of GIMAP6 was predicted from the protein sequence using PHYRE, a structure prediction tool. The prediction result showed that GIMAP6 would seem to have a structure very similar to those of GTP-binding proteins with NTP hydrolase activity. Because GTP hydrolysis plays an important role in regulating the activity of GTP-binding proteins, it is important to measure the GTPase activity of GIMAP6. To characterize the biochemical properties of GIMAP6, GTP hydrolysis assays were performed using two different methods, the traditional isotope-labeled NTP hydrolysis assay and the non-radioactive colorimetric phosphate detection assay. The traditional isotope-labeled NTP hydrolysis assay uses $^{32}$P-labeled ATP, GTP, UTP, and CTP as substrates when performing the experiments. The results showed that GIMAP6 exhibits intrinsic GTPase activity and that this activity is dependent on the presence of Mg$^{2+}$ ions (data not shown). To our surprise, GIMAP6 also showed ATP hydrolysis activity (Fig. 4A). A time course assay of ATP and GTP hydrolysis by GIMAP6 was then carried out to confirm this finding (Fig. 4B). This confirmed that GIMAP6 is the first protein in the GIMAP family to have ATPase activity. Even though such a finding has never been reported previously for the GIMAP family, the...
coexistence of GTP and ATP hydrolysis activity does occur in some other P-loop NTPase superfamily members (20–22). To further investigate the enzyme kinetics of GIMAP6 and to determine the turnover number and the Michaelis-Menten constant of the enzyme, saturation assays were performed (Fig. 4C). The $K_{cat}$ was calculated as $K_{cat} = V_{max}/[E]$, and the ratio $K_{cat}/K_m$ was used to monitor enzyme efficiency. We calculated that, for ATP, $K_{cat}$ was 0.695 (1/min), $K_m$ was 104.44 (μM), and $K_{cat}/K_m$ for ATP was 111 (M⁻¹s⁻¹). For GTP, $K_{cat}$ was 0.102 (1/min), $K_m$ was 46.38 (μM), and $K_{cat}/K_m$ for ATP was 36.3 (M⁻¹s⁻¹). Thus, we concluded that the recombinant GIMAP6 had a higher affinity to GTP but showed a higher catalytic rate to ATP.

The anti-apoptotic function of GIMAP6 is independent of the GTP/ATP hydrolysis activities of the enzyme

To study whether the GTPase/ATPase activity of GIMAP6 is required for the anti-apoptotic function of the enzyme, we generated four mutant enzymes (G47V, G47A, G52V, and G52A) in which GTP/ATP binding by the P-loop motif should be absent. A previous study has shown that site-directed mutagenesis in either of the two glycine residues of the P-loop is able to greatly diminish enzyme activity by reducing the affinity of the enzyme for GTP and/or ATP (20, 23–25). Using these mutants with reduced GTP/ATP binding, it was found that the GTP/ATP hydrolysis activity of GIMAP6, as measured by the non-radioactive colorimetric phosphate assay, had been significantly reduced (Fig. 4D).

To determine the effect of this reduced GTP/ATP hydrolysis activity on the anti-apoptotic function of GIMAP6, the WT and the G52A mutant of GIMAP6 were transiently transfected into Huh-7 cells for 24 h. The cells were then harvested after OA treatment for 24, 36, and 48 h. Compared with cells containing the WT enzyme, no significant difference in the
anti-apoptotic effect of GIMAP6 was observed with the G52A mutant. This was determined by measuring the level of active-form caspase-3 present in the cells by immunoblot analysis (Fig. 4E). These findings indicated that, under our assay conditions at least, the anti-apoptotic function of GIMAP6 is independent of the GTP/ATP hydrolysis activity of the enzyme.

Knockdown of GIMAP6 accelerates PMA/ionomycin-induced T cell activation

T cell activation and differentiation play a major role in adaptive immunity. Small GTPases are known to be involved in these specific processes. For example, GMAP1 and GMAP4 have been reported to be differentially regulated during CD4+ T helper differentiation (15). In this study, we have demon-
strated that PMA/ionomycin-induced T cell apoptosis and activation is modulated by the level of expression of GIMAP6 (Fig. 5). Ctl and knockdown (KD1 and KD2) cells were treated with PMA/ionomycin for 24 h, and the cells were harvested and stained with Annexin-V-FITC to detect apoptosis (Fig. 5, A and B) and with CD69 to detect T cell activation (Fig. 5, A and C). Additionally, culture medium was collected to allow the measurement of IL-2 secretion (Fig. 5, D).

Figure 5. GIMAP6 knockdown accelerates the PMA/ionomycin induction of T cell activation. A. Jurkat-derived cell lines (Ctl, KD1, and KD2) were treated with either PMA/ionomycin or an equal concentration of DMSO to determine the effect of GIMAP6 on T cell activation. To monitor the rate of cell apoptosis/death and T cell activation, PMA/ionomycin-treated cells were harvested at 3, 8, and 24 h and stained for the phosphatidylserine exposure marker Annexin V-FITC and for the T cell activation surface marker CD69-PE. The results were analyzed and quantified by FlowJo 7.6.1. B, the effect of GIMAP6 expression on PMA/ionomycin-induced cell death. All AnV+ cells were taken as apoptotic/dead cells, and the remaining AnV− cells were taken as live cells. C, the effect of GIMAP6 expression on PMA/ionomycin-induced T cell activation. To discriminate activated T cells from inactivated T cells across all live cells, the surface marker CD69 was used to calculate the percentage of CD69+/AnV−. D, the effect of GIMAP6 expression on PMA/ionomycin-induced IL-2 secretion. The data represent the average of at least two independent experiments with triplicate measurements (mean ± S.D.). P/I indicates that the cells were treated with both PMA (10 ng/ml) and ionomycin (1 μg/ml).
of cells activated after treatment for 24 h. Thus we conclude that GIMAP6 knockdown is able to accelerate T cell activation in this experimental system.

We also determined the level of cytokine IL-2 secretion as a measure of T cell activation (Fig. 5D). IL-2 is a potent growth factor and plays an important role in T cell activation and homeostasis (26). In this experiment, IL-2 secretion by GIMAP6 knockdown cells became detectable after treating the cells with PMA/ionomycin for 8 h. We were only able to observe IL-2 secretion after 24 h in control cells. Moreover, the IL-2 concentration released by the knockdown cells was nearly 4-fold higher than that of the control cells. These results totally agree with the surface marker CD69 staining results; both indicate that knockdown of GIMAP6 accelerates the activation of T cells. Our findings are consistent with previous studies showing that IL-2 acts as a sensitizer and renders T cells susceptible to the induction of apoptosis during T cell activation and proliferation (27–29). This might help to explain why we have observed a high apoptotic/dead ratio in high IL-2-secreting GIMAP6 knockdown cells (Fig. 5, B and D).

GIMAP6 displays anti-apoptotic function in PMA/ionomycin-induced T cell activation through the phosphorylation of NF-κB p65

To investigate whether the anti-apoptotic protein GIMAP6 plays a role in T cell functioning, PMA and/or ionomycin were used to induce T cell activation. We found that long-term exposure of Jurkat T cell to PMA greatly induced GIMAP6 expression. After treatment for 24 h, the expression level of GIMAP6 increased 8-fold at the RNA level (Fig. 6A). Consistently, the change in the level of protein was 2.56 times higher after 48 h (Fig. 6B). Previous reports have shown that treating Jurkat T cells with PMA results in an up-regulation of AP-1 and a down-regulation of NF-κB activity (30, 31). We speculated that the induction of GIMAP6 might be controlled via either the protein kinase C/AP-1 or the NF-κB signal transduction cascades. To investigate these possibilities, an inhibitor was used to suppress the induction. When AP-1 and NF-κB activity were suppressed in parallel by treating the cells with 1–3 μM SP100030, there was significant suppression of GIMAP6 at the RNA level (Fig. 6A). We further observed that, when cells were treated with SP100030 alone, there was a slight induction of expression of GIMAP6 rather than a suppression of GIMAP6 expression (data not shown). These findings indicate that SP100030 is not able to suppress the endogenous expression of GIMAP6. SP100030 has been reported to be a potent dual inhibitor of AP-1 and NF-κB and has been used to inhibit cytokine production selectively in T cells (30–32). Because both PMA and SP100030 suppress NF-κB activity and AP-1 activity is absolutely necessary for the induction of GIMAP6, it can be concluded that the
induction of GIMAP6 expression by PMA occurs via the AP-1 signaling cascade.

The NF-κB signal transduction cascade is a key regulator of the life and death of T cells. Loss of NF-κB activation initiates activation-induced cell death (AICD) in T cells (33, 34), which will cause accumulation of reactive oxygen species and suppression of BCL-2 expression, finally leading to the release of BIM and resulting in T cell death through the intrinsic apoptotic pathway (35). In this study, we observed that NF-κB activation is affected by the GIMAP6 expression level in PMA/ionomycin-mediated activation. As shown in Fig. 6C, GIMAP6 knockdown showed a significant decrement of p65 phosphorylation (Ser-536) under the PMA/ionomycin treatment condition compared with control cells. Because loss of NF-κB activation can result in T cell death via apoptosis, the decreased phosphorylation of p65 (Ser-536) in GIMAP6 knockdown cells (KD1 and KD2) might help to explain the high apoptosis/dead ratio observed in the Annexin-V assay (Fig. 5B). Thus, we conclude that GIMAP6 knockdown leads to decreased phosphorylation of p65 (Ser-536) in PMA/ionomycin-stimulated activation, providing a possible mechanism of how GIMAP6 functions as an anti-apoptotic protein.

**Activation with PMA/ionomycin is affected by GIMAP6 expression in human primary T Cells**

As Jurkat cells are derived through human T cell leukemia/lymphoma virus type I infection and have accumulated multiple genetic mutations that contribute to immortalization (36), we further investigated whether GIMAP6 knockdown had an effect on PMA/ionomycin-induced activation in human primary T cells. To this end, CD3+ T cells were enriched from PBMCs (supplemental Fig. S2), and knockdown of GIMAP6 was performed by transfection of either negative control (Neg) or anti-GIMAP6 siRNAs (G6-si599 and G6-si600) into the cells (supplemental Fig. S3). After 48 h, the knockdown efficiency of GIMAP6 was confirmed by quantitative RT-PCR assay. As shown in Fig. 7A, the knockdown efficiency of the anti-GIMAP6 siRNA G6-si600 and G6-si599 group in CD3+ T cells averaged 26.8% and 6%, respectively. Following the conclusion of the analysis in Jurkat cells (Fig. 5), we hypothesized that a higher level of activation should be observed after knocking down GIMAP6 in primary CD3+ T cells. We then measured PMA/ionomycin-induced IL-2 secretion in these three transfectant groups. As shown in Fig. 7B, no significant difference between these three transfectants was observed at 24 and 48 h after treatment with PMA/ionomycin. However, 72 h after stimulation with PMA/ionomycin, IL-2 production enhancement by 53% was observed in the high knockdown efficiency group (G6-si600). Thus, we concluded that the GIMAP6 expression level had an effect on the increased IL-2 production stimulated by PMA/ionomycin.

To confirm that primary T cell activation is sensitive to the GIMAP6 level, we analyzed the PMA/ionomycin-induced expression of the cell surface antigens CD25 (IL-2 receptor α chain) and CD154 (CD40 ligand). Consistent with the IL-2 production level, the high knockdown efficiency group (G6-si600) showed an elevated level of both antigens 72 h after induction (Fig. 7, C and D). These results demonstrate that down-regulation of GIMAP6 leads to enhanced PMA/ionomycin-stimulated activation in various activation signals, supporting the role of endogenous GIMAP6 as a negative regulator of T cell activation.

**Discussion**

Across species, the biological functions of the GIMAP family generally seem to be related to immune reactions against internal or external stimuli. Although the structural features of the GIMAP proteins are evolutionarily conserved in vertebrates and angiosperm plants, the biochemical and molecular properties underlying various functions of the GIMAP proteins are not fully characterized (2). In this study, we have demonstrated that GIMAP6 is a T cell-associated cytosolic protein with an anti-apoptotic function and that it plays a role in regulating T cell activation.

Apoptosis plays a critical role in T cell development and function. Dysregulation of apoptosis in the immune system results in autoimmunity, tumorigenesis, or immunodeficiency. We have clearly shown that GIMAP6 knockdown not only made cells apoptosis-prone but also accelerated T cell activation when treated with PMA/ionomycin. One possibility is that the reduction of GIMAP6 facilitates apoptosis and speeds up the removal of cells without appropriate signals, and this eventually accelerates T cell activation. Therefore, we speculate that GIMAP6 can function as an anti-apoptotic protein as well as a blockade to T cell activation. That is to say, physiologically, GIMAP6 in the immune system might serve as a T cell preserver to keep cells alive and retard or prohibit reactivation.

Taking functional and biochemical approaches, we have gained new insights about the molecular mechanisms of GIMAP6 in modulating T cell functions. We used three different apoptosis inducers, H2O2, FasL, and OA, to treat cultured cells. Although all three inducers are able to trigger cell death, they appear to act via a number of signaling pathways. Therefore, we concluded that the anti-apoptotic function of GIMAP6 might involve the intrinsic apoptosis pathway. A previous report has indicated that mouse IAN1 (mouse Gimap4) binds to Bax and IAN4 (mouse Gimap3) and that IAN5 (mouse Gimap5) binds to Bcl-2 and Bcl-xl, all of which are directly involved in the apoptosis pathway (1). In an attempt to gain further insights into the anti-apoptotic mechanism of GIMAP6, an antibody array was used to screen interaction with GIMAP6 and binding to GIMAP6. So far, we have found no evidence to support the direct binding of GIMAP6 to any appropriate candidate protein (data not shown). While this study was in progress, we became aware of a publication by Pascall et al. (37) that reports that GIMAP6 interacts with the Atg8 homologue GABARAPL2 and that it is recruited to autophagosomes. The biological significance of the GIMAP6-GABARAPL2 protein-protein interaction remains unclear, and its relevance to controlling cell death in T cell development and activation warrants further investigation.

The balance between activation and apoptosis plays an important role in maintaining T cell homeostasis. We observed an elevated level of apoptosis and activation in GIMAP6 knockdown cells when treated with PMA/ionomycin. Also, the anti-apoptotic function of GIMAP6 is related to NF-κB activation.
These findings are in accordance with the knowledge that restimulation of expanded T cells in the absence of co-stimulatory factors can lead to efficient induction of cell death, known as AICD. NF-κB plays an important role in this process, as loss of NF-κB activity results in T cells death through the intrinsic apoptotic pathway (27, 35). In our study, down-regulation of p65 phosphorylation and increase in apoptosis were observed simultaneously in GIMAP6 knockdown cells after treatment with PMA/ionomycin (Figs. 5B and 6C). This implies that phosphorylation of p65 and its downstream cascade AICD is affected by GIMAP6 expression level.

Notably, GIMAP6 is up-regulated in T cells activated by PMA/ionomycin (Fig. 6A), whereas GIMAP6 inhibits T cell activation (Fig. 5). This is expected to lead to an elevated reactivation threshold for activated T cells. It may be envisioned that increased levels of GIMAP6 confer the survival of activated T cells by enhancing resistance to both restimulation and activation-induced cell death. Therefore, GIMAP6 may contribute to T cell immunity through regulating T cell viability during microbe infections and autoimmune diseases. Further studies will help establish the exact physiological role of GIMAP6.

In terms of the biochemical properties of the protein, recombinant GIMAP6 has both GTP and ATP hydrolysis activity. Interestingly, ATPase activity has never been reported previously for any GIMAP protein. ATP usually functions either as a signal molecule for protein phosphorylation via protein kinases...
Anti-apoptosis function of GIMAP6

(38) or as an energy source for the pumps involved in transporting substances inside/outside of cells (39, 40). It also powers the molecular motors of muscle cells during contraction (41). By way of contrast, GTP can act as a molecular switch, and it is a well known part of signal transduction regulation (42). Even though this function is not common, some proteins that contain the P-loop motif, such as msP2 of Semliki Forest virus, 2C of poliovirus, p206 of turnip yellow mosaic virus (20, 21, 43), elongation factor 3 of yeast, FtsZ of Escherichia coli, and Rab14 of Bombyx mori (silkworm), have been reported as possessing nucleotide triphosphate hydrolase superfamily, and it is used during the structural classification of proteins (48, 49).

The elution pattern of GIMAP6 indicates that it is present in cells as a multimer (supplemental Fig. S4). The nature of protein interactions in the GIMAP6 complex remains unclear, and it is reasonable to speculate that such multiprotein interaction(s) might have structural, functional, or regulatory implications (50). As GIMAP6 has been shown to exhibit GTP/ATP hydrolysis activity and has been observed to oligomerize, it is possible to connect these two properties based on previous reports. Many P-loop proteins, including the GTPases atToc33 and IIGP1 and the GIMAP family member GIMAP7 (8, 51, 52), have been shown previously to function as oligomers inside the cell for the purpose of hydrolysis activity. Therefore, further study is required to identify the interaction domain of GIMAP6 and to confirm whether oligomerization is required for the GTP/ATP hydrolysis activity of GIMAP6.

Finally, it appears that GIMAPs, by controlling cell survival and death, are able to modulate the proportion of differentiated cell lineages in the hematopoietic system. Different cell lineages express different levels of GIMAP genes. In addition to the Jurkat cell line, in this study, HL-60 is the only other cell line expressing a detectable level of GIMAP6 (Fig. 1, A and B). HL-60 is a promyelocytic cell line that is derived from peripheral blood leukocytes with a myeloblast morphology; it is often used for studying the molecular mechanism of myeloid differentiation (53–55). The significance of GIMAP6 expression in HL-60 cells remains unclear. Based on the above, we propose that, by turning on and off the GIMAP genes that have pro-apoptotic or anti-apoptotic functions, there is a dynamic regulation of the cell numbers of the various components of the hematopoietic system that make up the immune reaction, and these changes in cell numbers are important during development and disease. Thus, our findings regarding the functional and biochemical properties of GIMAP6 warrant further study to advance the knowledge of how GIMAP proteins maintain lymphocyte homeostasis.

Experimental procedures

Purification of recombinant GIMAP6 protein

The full-length human GIMAP6 (GIMAP6, NM_024711) coding sequence was amplified by PCR using primers containing Ndel and Xhol cutting sites and a commercial cDNA clone (MHS1010-98051901, MCLAB) as a template. The amplified DNA was then cloned using the Ndel and Xhol sites into the vector pET29 (Novagen), which resulted in a coding sequence that contained a six-histidine tag at the C terminus. This plasmid was then transformed into E. coli BL21 (DE3) and expressed in the presence of 1 mm isopropyl 1-thio-β-d-galactopyranoside in LB medium. Purification was performed using a HisTrap HP column (GE Healthcare), which was followed by size exclusion chromatography.

Gel filtration assay

Recombinant GIMAP6 was subjected to size exclusion separation on a Superose 12 10/300 GL prepacked column connected to an AKTA-explorer system (GE Healthcare), which was controlled by the UNICORN software. After automatic sample injection, protein separation was performed using elution buffer (100 mM Tris-HCl and 200 mM NaCl (pH 7.4) at a flow velocity of 0.5 ml/per minute, and 1-ml fractions were collected.

Cells and cell culture

Human cell lines, 293T and Huh-7, were maintained in DMEM (Gibco), whereas the Jurkat cell line was maintained in RPMI 1640 medium (Hyclone). All media were supplemented with 10% fetal bovine serum containing 1% penicillin, 1% streptomycin, and 1% glucose (10378-016, Gibco).

Plasmid preparation and transfection

The previously described GIMAP6 cDNA clone was amplified and cloned into the pcdNA3.1(+) vector (Invitrogen) using EcoRI and Xhol cutting sites. The expression construct was then transformed into E. coli DH5α and grown in LB medium. Plasmids were prepared using a PureLink plasmid midiprep kit (K2100-05, Life Technologies) and then introduced into the various cell lines using transfection reagent (Turbofect R0531, Fermentas) according to the protocol of the manufacturer.

Immunoblotting

Protein samples were separated by electrophoresis using an appropriate percentage SDS-PAGE gel and transferred onto a polyvinylidene difluoride membrane (Millipore). The membrane was blocked with blocking buffer (5% skimmed milk in TBS-T buffer (10 mM Tris, 150 mM NaCl (pH 7.6), and 0.05% Tween 20)) for 1 h at room temperature. The primary antibody was appropriately diluted into the blocking buffer, and the membrane was then incubated with the mixture overnight at 4 °C. Next the membrane was washed with TBS-T buffer. Finally, diluted horseradish peroxidase-conjugated secondary antibody corresponding to the primary antibody was added to the buffer, and the mixture was incubated for 1 h at room temperature. After washing with TBS-T buffer, the membrane was developed using ECL substrate (WBKLS0500, Millipore) and photographed using an ImageQuant™ LAS 4000 (GE Healthcare).

RNA extraction, RT-PCR, and quantitative RT-PCR

Total RNA samples were prepared using TRIzol reagent (Invitrogen) from the indicated cells, and reverse transcription
was carried out using 100 ng of total RNA according to the instructions of the manufacturer (Superscript® III, Life Technologies). Quantitative PCR was performed using TaqMan probe-based gene expression analysis in 48-well plates on a StepOne real-time system (Applied Biosystems). The mRNA levels of the target genes in the samples were normalized against HPRT1 (4326321E, Applied Biosystems). The probe ID for the indicated gene is GIMAP6 Hs00226776_m1.

**Generation of anti-GIMAP6 antibodies**

To characterize GIMAP6 at the protein level, rabbit polyclonal antibodies were raised against recombinant GIMAP6. Rabbits were inoculated with 500 μg of GIMAP6-His fusion protein in 2 ml of PBS mixed with 2 ml of complete Freund adjuvant. Five boosts of 200 μg of protein in 2 ml of PBS plus 2 ml of incomplete Freund adjuvant were administrated over 3 months. All antisera from the rabbit blood used in this study stained only a single protein band with high intensity at the expected molecular size (35 kDa) in immunoblots against GIMAP6-transfected 293T and Huh-7 cells, secondary Jurkat T cells, and human PBMC cell lysates.

**Immunofluorescence microscopy**

Jurkat or primary CD3+ T cells were collected by centrifugation and washed three times with PBS. After resuspension in PBS, the cells were then cross-linked onto poly-L-lysine-coated coverslides at 4 °C for 30 min and fixed with paraformaldehyde. Following penetration with ice-cold methanol and blocking with 0.5% BSA in PBS for 30 min at room temperature, the cells were incubated with a 1:500 dilution of rabbit anti-GIMAP6 and mouse anti-GAPDH antibody for 1 h at room temperature. After three washes with PBS, the secondary antibody was added at a 1:500 dilution. Alexa Fluor 647 or 488 goat anti-rabbit IgG (Life Technologies) and Alexa Fluor 488 or 594 goat anti-mouse IgG (Life Technologies) was used. Confocal laser-scanning microscopy was performed using an Olympus FV10i or Zeiss LSM 700.

**Nucleotide triphosphatase assays**

Nucleotide triphosphatase activity was assessed by either thin-layer chromatography (TLC) or colorimetric phosphate detection. To determine the hydrolysis activities for GIMAP6, α-32P-labeled ATP, GTP, UTP, and CTP (NEG503H, NEG506H, NEG507H, and NEG508H, respectively) were used as substrates. Time course experiments were carried out in a 20-μl reaction mixture containing 50 mM Tris (pH 7.9), 100 mM NaCl, 2.5 mM MgCl₂, and 2.43 mM wild-type GIMAP6 containing 33 nm α-32P of either ATP or GTP (1 μCi) (PerkinElmer Life Sciences) and a specific concentration of ATP or GTP. The reaction was incubated at 37 °C. For TLC, 2 μl of the α-32P-labeled reaction was removed at various time points and mixed with 2 μl of solution containing 0.2% SDS and 5 mM EDTA. Samples were incubated at 70 °C for 2 min to stop the reaction. A fraction of the stopped reaction (0.5 μl) from each time point was spotted on a polyethyleneimine cellulose TLC plate (Merck), and the plate was developed in 1 M formic acid and 0.5 M LiCl. The plate was then dried and exposed to a PhosphorImager for quantitative analysis. The rate of ATP/GTP hydrolysis at each nucleotide concentration was calculated and plotted.

To determine the hydrolysis activities of the G47V, G47A, G52V, and G52A mutants, colorimetric phosphate detection was performed by adding 100 μl of malachite green reagent to the non-radioactive reaction mixture and incubated for 30 min at room temperature to develop a green color. A buffer blank was used for background subtraction, and the absorbance at 650 nm was measured to calculate the ATP/GTP hydrolysis rate by interpolation onto a phosphate standard curve.

**Detection of apoptosis by FACS analysis**

Apoptosis was measured using an apoptosis detection kit (Annexin V-FITC and PI) (BD Biosciences) according to the instructions of the manufacturer. Cells were analyzed on a flow cytometer (BD Biosciences, Canto), and we manually compensated for crossover of FITC fluorescence into the PI detection window. The analysis was completed using FlowJo 7.6.1.

**Knockdown of GIMAP6 in human Jurkat T lymphocytes**

Human Jurkat T lymphocytes were maintained in RPMI 1640 medium with 10% FBS. The cells were transfected with anti-GIMAP6 shRNA plasmids (TRCN0000151921, TRCN0000152080, TRCN0000152413, TRCN0000154009, and TRCN0000156658) or a control plasmid (pLKO.1.nullT) (provided by Academia Sinica, Taiwan) using the Neon transfection system (Invitrogen) in accordance with the recommended protocol. After limiting dilution followed by selection with 4 μg/ml puromycin (Sigma) for 2 weeks, cloning of single cells was achieved. These cells were then amplified for further study.

**Overexpression of GIMAP6 in the HEK293T and Huh-7 cell lines**

HEK293T and Huh-7 cells were cultured separately in Dulbecco’s modified Eagle’s medium supplemented with 10% fetal bovine serum. Transfection of the indicated expression plasmids separately into these cell lines was performed using TurboFect (Fermentas, R0531) according to the instructions of the manufacturer. The transfected cells were then cultured under the indicated conditions for 24 or 48 h. A stably expressing single colony was picked after G418 selection (500 μg/ml) for 2 weeks.

**Peripheral blood mononuclear cell subset cell sorting**

Peripheral blood mononuclear cells (PBMCs) were collected via Ficoll-Paque Plus (GE Healthcare, 18-1152-69) centrifugation from a fresh blood donor sample according to the instructions of the manufacturer. Cell sorting was accomplished by CD3-PE (BD Biosciences, 555333), CD4-FITC (BD Biosciences, 555346), and CD8-allophycocyanin (APC) (BD Biosciences, 555369) triple staining. Three subsets of cells, CD3+/CD4+, CD3+/CD8+, and CD3−, were collected after flow cytometry cell sorting. RNA was extracted from each subset immediately for use in the subsequent experiments.

**Cellular fractionation**

Subcellular fractionation was carried out on cultured Jurkat T lymphocytes using the ProteoJET cytoplasmic and nuclear protein extraction kit (Fermentas, K0311). A heavy membrane...
Anti-apoptosis function of GIMAP6

fraction that contained the mitochondria, a nucleus fraction, and a cytosolic fraction were obtained, and these were analyzed by immunoblotting using the indicated antibodies. For this analysis, histone H3, GAPDH, and PHB1 were used as controls for the nucleus, cytosol, and heavy membrane fractions, respectively.

**Cell content screening**

Cells were seeded on 24-well plates overnight using 1 × 10⁴ cells/well. After treatment with the indicated reagents/drugs for 24 h, the cells were fixed on plates, and apoptosis detection was performed using rabbit anti-cleaved caspase-3 (Cell Signaling Technology) as the primary antibody and Alexa Flour 488 goat anti-rabbit IgG (Invitrogen) as the secondary antibody. With DAPI counterstaining, 16 images were taken (ImageXpress Micro XLS system) for each well, and the data were analyzed using MetaXpress software (Molecular Devices).

**Jurkat T cell activation**

Cells were seeded on 24-well plates overnight using 2 × 10⁵ cells/well. After treatment with either PMA/ionomycin or an equivalent concentration of DMSO, cells were harvested at 72 h and then stained with the T cell activation surface marker Annexin V-FITC (BD Biosciences) and the T cell activation surface marker CD69-PE (Biolegend, 310906). The results were analyzed and quantified using FlowJo 7.6.1.

**IL-2 ELISA**

In the Jurkat T cell activation experiments, supernatants of the media were collected from each well after centrifugation and stored at −80 °C until the relevant assays were carried out. IL-2 was determined using a human IL-2 ELISA kit (Biolegend, 431801).

**Knockdown of GIMAP6 on human primary CD3⁺ T cells**

After PBMC collection from healthy donors, CD3⁺ T cells were further enriched using a human CD3 T cell negative selection kit (Biolegend, 480022). The purity of the CD3⁺ T cell population was examined by flow cytometry with anti-CD3-PE (BD Biosciences, 555333) antibody. The cells were then transfected with anti-GIMAP6 siRNA (Ambion, s54599, s54600) or a negative control (AM4611) using the Neon transfection system (Invitrogen) in accordance with the recommended protocol. The transfection efficiency was monitored by transfecting Cy3-labeled negative control siRNA (AM4621). After transfection for 48 h, the transfection efficiency was checked on fluorescence microscopy. Then cells were either harvested for total RNA extraction or for further experiments.

**Primary CD3⁺ T cell activation**

Previously described GIMAP6 knockdown or negative control CD3⁺ cells were seeded on 24-well plates using 1 × 10⁶ cells/well. After treatment with either PMA/ionomycin or an equivalent concentration of DMSO, cells were harvested at 72 h and then stained with the T cell activation surface marker CD25-APC (Biolegend, 302609) and CD154-Alexa Flour 488 (Biolegend, 310815). The results were analyzed and quantified using FlowJo 7.6.1.

Acknowledgments—We thank Dr. Ming-Zong Lai and Dr. I-Cheng Ho for critical reading of this paper and Dr. Ralph Kirby for editing the manuscript. We thank Hui-Ying Weng for donating the blood samples for the study and the staff of the VGH Genome Research Center at National Yang-Ming University for conducting Sanger sequencing.

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