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T Cell Ig and Mucin Domain-1-Mediated T Cell Activation Requires Recruitment and Activation of Phosphoinositide 3-Kinase^1

Anjali J. de Souza,^* Jean S. Oak,^† Ryan Jordanazy,^* Rosemarie H. DeKruyff,^‡ David A. Fruman,^† and Lawrence P. Kane^2*

Ligation of the transmembrane protein T cell Ig and mucin domain (Tim)-1 can costimulate T cell activation. Agonistic Abs to Tim-1 are also capable of inducing T cell activation without additional stimuli. However, little is known about the biochemical mechanisms underlying T cell stimulation or costimulation through Tim-1. We show that a tyrosine in Tim-1 becomes phosphorylated in a lck-dependent manner, whereupon it can directly recruit p85 adaptor subunits of PI3K. This results in PI3K activation, which is required for Tim-1 function. We also provide genetic evidence that p85 expression is required for optimal Tim-1 function. Thus, we describe a pathway from Tim-1 tyrosine phosphorylation to the PI3K signaling pathway, which appears to be a major effector of Tim-1-mediated T cell activation. The Journal of Immunology, 2008, 180: 6518–6526.

Proteins of the T cell Ig and mucin domain (TIM)^3 family function to modulate T cell activation and effector function. This family consists of four Tim proteins in the mouse and three TIM proteins in the human (1–3). The TIM proteins are cell surface, Ig superfamily receptors that may also be secreted and which share the following domain structure: an extracellular Ig-like domain, followed by a polymorphic mucin-like domain, a transmembrane domain, and a cytoplasmic tail of variable length (1–3).

Of these family members, TIM-1 has been implicated in the immune regulation of asthma and atopic diseases (3, 4). More recent studies have shown that ligation of TIM-1 with an agonistic Ab or with a ligand (TIM-4) can costimulate proliferation of T cells as well as cytokine production, both in vivo and in vitro, and abrogate tolerance in a Th2 model of respiratory tolerance, collectively suggesting that it usually functions as a positive immune regulator (5, 6). Our own prior work also showed that murine Tim-1 can transmit a costimulatory signal for activation and cytokine production of T cells (7). Finally, a recent report demonstrated that human TIM-1 can physically associate with the TCR-CD3 complex to up-regulate T cell activation signals (8). Nevertheless, depending on the affinity and targeted epitope of the cross-linking Ab, and also possibly the timing of its application, opposing conclusions have been reached regarding whether TIM-1 positively or negatively regulates T cell activation and disease outcomes (9–12). Thus, a greater understanding of the biochemical signals mediated by TIM-1 is required to mechanistically explain its role in immune activation.

The TCR/CD3 complex and costimulatory molecules interact with key intracellular signaling proteins that determine which downstream effector pathways are triggered. Of these, the PI3K have been frequently implicated in costimulatory signaling (13). These enzymes regulate diverse biological processes, including cell growth, proliferation, cytoskeletal organization, apoptosis, and vesicular trafficking (14). Among the various classes of PI3K, class IA proteins are linked to tyrosine kinase-dependent signaling downstream of lymphocyte Ag receptors, costimulatory receptors, and cytokine receptors. These proteins generally exist as heterodimers of a regulatory subunit (p85) and a catalytic subunit (p110). Class I PI3K function to phosphorylate inositol rings at the 3' position to generate phosphatidylinositol-3,4-bisphosphate, and phosphatidylinositol-3,4,5-trisphosphate (14). These lipids are stationary at the inner leaflet of the plasma membrane, where their major function appears to be to recruit signaling molecules with pleckstrin homology domains. Additionally, the p85 adaptor subunit contains several other interesting domains, including SH2, SH3, and proline rich domains, and a region with homology to Rho family GTPase-activating proteins. These domains facilitate protein-protein interactions that may be able to mediate signal transduction events independently of p110 (15).

To date, there is little known about the molecular basis for signal transduction downstream of TIM-1. We previously showed that the cytoplasmic tail of Tim-1 could be inducibly tyrosine phosphorylated, and that Tim-1 function in human and murine T cell lines requires a conserved tyrosine in the cytoplasmic tail (7). In this present study, we show that Tim-1 tyrosine phosphorylation and activation of downstream transcription factors require the src family kinase Lck, which is also critical for TCR-mediated signal transduction. Following phosphorylation, Tim-1 recruits the PI3K adaptors p85α and p85β. Finally, we show that Tim-1 can enhance activation of primary murine T cells in a manner that requires p85 expression and PI3K activity.
Materials and Methods

DNA constructs

Murine Tim-1 (from strain C57BL/6), tagged with an extracellular Flag tag, was described previously (7). Similarly, Flag-Tim-1 lacking the cytoplasmic tail (Tim-1 Δ cyto), and Flag-Tim in which tyrosine 276 was mutated to phenylalanine have also been described elsewhere (7).

Cell lines and mice

In addition to parental Jurkat T cells, the lck-deficient variant JCaM1.1 and reconstituted JCaM1/Lck were also used (16, 17). A fast-growing variant of the D10 T cell clone was obtained from M. Krummel (University of California, San Francisco, CA) and was described previously (7, 18). p85β knockout (k.o.) and p85αβ double k.o. mice have been described (19). C57BL/6 mice were purchased from Jackson ImmunoResearch Laboratories. Jurkat, D10, and primary T cells were cultured as previously described (7).

Abs and reagents

Clonotyptic Ab to the Jurkat TCR (C305) was from A. Weiss (University of California, San Francisco). Anti-human CD28, anti-mouse (CD8), anti-mαCD28, and anti-mCD25 were from Caltag Laboratories. Anti-PI3K p85 (rabbit anti- serum) and anti-phospho-tyrosine (4G10) were from Upstate Biotechnology/Chemicon International. Biotin-anti-mCD28 (37,51), bixin-anti-mCD3ε (145-2C11) and FITC-conjugated anti-CD69 were from BD Biosciences. Biotin-anti-mCD4 (L374) was from eBioscience. Streptavidin was from Zymed Laboratories. Rabbit anti-hamster IgG and allophtocyanin-conjugated anti-rat IgG were from Jackson ImmunoResearch Laboratories. Anti-Flag (M2) and β-actin mAbs and ionomycin were from Sigma-Aldrich. Rat anti-mouse Tim-1 (3B3) was described previously (5). Anti-phospho Akt (S473) rabbit mAb was from BioSource International/Invitrogen.

Transient transfections and luciferase assay

T cell lines were transfected by electroporation using a gene pulsar (Bio-Rad), at a setting of 250–260V and 960 μF, in cuvettes containing 1–2 × 10⁶ cells with 0.4 ml antibiotic-free media. The amount and type of DNA plasmids transfected for an experiment are as described in the respective figure legends. Following transfection, cells were cultured in 10 ml complete medium for 16–20 h and then stimulated for 6 h in the bottom well of a 24-well plate. Reporter assays were performed as described previously (20). Luciferase activity was determined with an Orion lumimeter (Zylux).

SH2 domain arrays

A 13 amino acid peptide based on the Tim-1 cytoplasmic tail was synthesized, including Y276 and six amino acids on either side (Peptide Synthesis facility, Biotechnology Center, Molecular Medicine Institute, University of Pittsburgh). The peptide was biotinylated on the N terminus and the cytoplasmic tail was phosphorylated. This was then subjected to HPLC; the end product obtained was greater than 90% pure, as determined by mass spectrometry. The purified peptide was incubated with the SH2 domain array, which was processed as per the manufacturer’s instructions (Panomics). Processed arrays were imaged on a Kodak Image Station 2000R, with accumulation to 2000 gray levels. When used as a competitive inhibitor, the charged residues glutamic acid and aspartic acid upstream of the D10 T cell line (Tim-1 Δ cyto) was described previously (7, 18). p85β knockout (k.o.) and p85αβ double k.o. mice have been described (19).

Interaction of Tim-1 with p85 subunits of PI3K

Regulation of Tim-1 tyrosine phosphorylation

We showed previously that Tim-1 can be inducibly phosphorylated on tyrosine and that tyrosine 276 in the cytoplasmic tail is required for Tim-1-mediated costimulation of NFAT/AP-1-dependent transcription in T cells (7). This tyrosine is conserved in context between different species (1) and is contained in a motif favorable for recognition by src-family tyrosine kinases, as indicated by the charged residues glutamic acid and aspartic acid upstream (22). To ascertain whether src kinases are indeed required for Tim-1 phosphorylation, we examined tyrosine phosphorylation in the presence of PP2, a potent inhibitor of src kinases (23). Jurkat cells were transfected with epitope-tagged Tim-1, and stimulated with pervanadate, in the presence or absence of PP2, followed by immunoprecipitation (IP) of Tim-1 and anti-phosphotyrosine Western blotting. As shown in Fig. 1A, tyrosine phosphorylation of Tim-1 was induced by treatment with pervanadate. Inclusion of PP2, along with stimulation by pervanadate, led to a significant decrease in the phosphorylation intensity, which was similar to basal levels. This indicates a requirement of src kinases for Tim-1 tyrosine phosphorylation, even under conditions where potent tyrosine kinase activation is induced, i.e., by pervanadate. Because Lck is a prominent src family tyrosine kinase essential for TCR-mediated T cell activation, we used Lck-deficient mutant variant of Jurkat cells, JCaM1 (17), to probe the role of Lck in Tim-1 tyrosine phosphorylation and function. The ability of Tim-1 to become tyrosine phosphorylated in JCaM1 cells was severely compromised (Fig. 1B), even after treatment with the potent stimulant pervanadate. In contrast, JCaM1 cells that have been stably reconstituted with Lck (JCaM1-Lck) showed a recovery of inducible Tim-1 phosphorylation (Fig. 1B) similar to what was observed in wild-type cells (Fig. 1A). To establish the functional relevance of lck-dependent Tim-1 tyrosine phosphorylation, we examined Tim-1-mediated NFAT/AP-1 activation in JCaM1 cells. Consistent with our previously published data with parental Jurkats (7), Tim-1 expression in Lck-reconstituted JCaM1 cells resulted in enhanced transcription of an NFAT/AP-1 luciferase reporter (Fig. 1C), both basally as well as in conjunction with stimulation through the TCR. In contrast, Lck-deficient JCaM1 cells displayed little, if any, reporter activity either with or without stimulation, despite the fact that Flag-Tim-1 expression was equivalent on both cell lines (data not shown). These data suggest that Tim-1 enhances T cell activation, at least in part, by coupling to phosphotyrosine-dependent signaling pathways.

Interaction of Tim-1 with p85 subunits of PI3K

One mechanism by which phosphotyrosine-dependent signaling complexes are formed is through phosphotyrosine-SH2 domain interactions. To identify SH2 domain-containing signaling proteins...
that might specifically bind to phosphotyrosine 276 of the Tim-1 tail, a solid phase screen was performed. The TranSignal SH2 domain array from Panomics contains 38 SH2 domains of various signaling proteins, folded in their native conformation and immobilized in duplicate on a membrane. A 13 amino acid peptide was synthesized (Fig. 2A), which contained the tyrosine 276 of Tim-1 tail and 6 amino acids on either side. The peptide was tyrosine phosphorylated and biotinylated and incubated with the SH2 domain array membrane (for key see Fig. 2B). As shown in Fig. 2C, two prominent sets of spots were obtained, which correspond to the N-terminal SH2 domain of p85α/H9251 and p85α/H9252. Other spots that were not consistent and of a much lower signal intensity corresponded to SH2 domains from Ras GTPase-activating protein (RAS-GAP) and the tyrosine kinase Fyn. We next performed the assay in the presence of a competitive inhibitor—phenylphosphate—to confirm that the binding observed was dependent on phosphotyrosine. As shown in Fig. 2D, a much weaker signal was obtained when the SH2 array was incubated with phosphorylated Tim-1 peptide in the presence of phenylphosphate. By comparing the p85 signal with the positive control spots along the right and bottom edges, we determined that binding of the peptide to the p85α and p85β N-terminal SH2 domains was inhibited by 99 and 95%, respectively.

To validate the Tim-1-p85 association with full-length proteins, the interaction was also established in T cells by conventional IP. Flag-Tim-1-transfected Jurkat cells were stimulated with pervanadate, followed by anti-Flag IP and Western blotting with a pan-p85 Ab. Although there was a reproducibly strong interaction of Tim-1 with p85 under conditions of pervanadate stimulation, a weaker and less consistent association was also detected under resting conditions (Fig. 3A). Consistent with the functional importance of

FIGURE 1. Role of Lck in Tim-1 tyrosine phosphorylation and function. A, Jurkat T cells expressing Flag-Tim-1 were stimulated as indicated, with or without the Src kinase inhibitor PP2. Anti-Flag immunoprecipitates were separated by SDS-PAGE and Western blotted with anti-Flag (lower panel), then stripped and re-probed with anti-phosphotyrosine Ab 4G10 (P-Tyr; upper panel). B, Lck-deficient JCaM1 or Lck-reconstituted JCaM1 cells expressing Flag-Tim-1 were stimulated and analyzed as in A. C, JCaM1 or JCaM1-Lck cells were transfected with an NFAT/AP-1 luciferase reporter and either empty vector or Flag-Tim-1. Cells were stimulated the next day and luciferase activity was determined. Each experiment is representative of two (A and B) or three (C) that were performed. stim, Stimulated; and unstim, unstimulated.

FIGURE 2. Specificity of SH2 domain binding to phosphorylated Y276 in the Tim-1 cytoplasmic tail. A, Sequence of the peptide, and modifications, used to determine the SH2 specificity of phospho-Y276. B, Layout of the SH2 domain array. SH2 domains are present as duplicate spots. For proteins with more than one SH2 domain, “D1” indicates the more N-terminal SH2. C, Results of the SH2 domain array screen, using the peptide illustrated in A. The dashed box indicates the location of negative control spots; the bottom and right edge contain positive control spots. These results are representative of those obtained in four separate experiments, using two different lots of membrane. D, Competitive inhibition of Tim-1 peptide binding by the phosphotyrosine analog phenylphosphate. The blot shown is representative of two experiments.
tyrosine 276 in the Tim-1 cytoplasmic tail (7), there was a loss of inducible p85 binding to mutant forms of Tim-1 either lacking the cytoplasmic tail or with Y276 mutated to phenylalanine (Fig. 3B). In agreement with data discussed above, this interaction was also significantly diminished in cells treated with PP2 (Fig. 3C). A major target for PI3K-derived lipids is the serine/threonine kinase Akt, activation of which is absolutely dependent upon PI3K. We therefore examined whether endogenous Tim-1 ligation results in activation of Akt, as assayed by its phosphorylation at serine 473. As shown in Fig. 3D, we observed inducible Akt activation, in both primary T cell blasts and the Th2 T cell clone D10, after stimulation (of untransfected cells) with Abs to CD3 and CD28 or Tim-1. Akt activation by either CD3/CD28 or Tim-1 was completely inhibited by pretreatment with the PI3K inhibitor LY294002 (data not shown). These results demonstrate that Tim-1...

FIGURE 3. Recruitment and activation of PI3K by Tim-1. A, Jurkat T cells expressing Flag-Tim-1 were stimulated as indicated, lysed, and immunoprecipitated with anti-Flag. Whole cell lysates (WCLs) and IPs were separated by SDS-PAGE and Western blotted for p85α/β. B, Jurkat T cells expressing the indicated wild-type or mutant Flag-Tim-1 constructs were stimulated and analyzed for p85 binding as described for A. C, Jurkat T cells expressing wild-type Flag-Tim-1 were stimulated in the presence of 10 μM PP2 (Src kinase inhibitor) or 10 μM Ly294002 (PI3K inhibitor) then analyzed for p85 binding as above. D, T cell blasts (left panels) or D10 Th2 T cells (right panels) were stimulated as indicated. Lysates were separated by SDS-PAGE and Western blotted with a rabbit mAb to phospho-Akt (S473; upper panels). Blots were stripped and re-probed with a mouse mAb to β-actin (lower panels), to control for protein loading. The location of phospho-Akt is indicated by the arrows, whereas the star indicates a nonspecific band that is only observed in primary T cells. Results in each part are representative of three experiments.

FIGURE 4. Role of the PI3K pathway in Tim-1 activation of NFAT/AP-1. T cells were transfected with an NFAT/AP-1 luciferase reporter and the indicated constructs, then, stimulated the next day, as indicated, followed by determination of luciferase activity. A, Cells were stimulated in the presence or absence of the PI3K inhibitor LY294002. B, Cells were stimulated in the presence or absence of the Akt inhibitor Akti1/2 (20 μM). C and D, T cells were transfected with an NFAT/AP-1 reporter, plus either empty vector or Flag-Tim-1, and the indicated amounts of short interfering RNA SmartPool oligos (Dharmacon) specific for p85α and p85β. Twenty-four-hour transfected cells were stimulated for 6 h, followed by determination of luciferase activity. Error bars indicate SD for triplicate points in a single experiment. Results shown in each part are representative of three experiments.
can interact with p85 and activate PI3K signaling, when expressed in T cells, in a phosphorylorysin-dependent manner.

Role of p85 and PI3K in Tim-1 function

p85 proteins function as regulatory subunits of class IA PI3K and, as such, control localization and activation of the associated catalytic p110 proteins. To determine whether the catalytic property of PI3K is required for Tim-1-mediated costimulation of downstream NFAT/AP-1 dependent transcription, D10 cells (which possess a normally regulated PI3K pathway) (18) were cotransfected with an NFAT/AP-1 luciferase reporter, along with either empty vector or Tim-1. These cells were then treated with a range of stimuli, in the presence or absence of LY294002, an inhibitor of PI3K activity (24). As shown in Fig. 4A, stimulation through CD3 and CD28 induced an enhanced activation of the reporter in Tim-1-transfected cells, compared with cells transfected with empty vector. However, in the presence of LY294002, the effect mediated by Tim-1 was markedly reduced. These data suggest that one mechanism by which Tim-1 costimulates T cell activation is through the PI3K/Akt pathway.

One of the earliest activation markers expressed by T cells is the cell surface glycoprotein CD69, which appears within several hours following TCR cross-linking and persists for ~3 days (27, 28). To assess the effect of Tim-1 on CD69 up-regulation, purified CD4+ T cells from C57BL/6 mice were activated with a range of concentrations of anti-CD3 Ab (and a fixed concentration of anti-CD28 Ab, 0.5 μg/ml), with or without anti-Tim-1 Ab (6 μg/ml), in the presence or absence of PI3K inhibitor LY294002 (10 μM). Cells were stained with fluorescently labeled Abs to CD69 (A) or CD25 (B) and analyzed by flow cytometry. Results in each panel are representative of three experiments that were performed.

FIGURE 5. PI3K activity is required for up-regulation of early activation markers by endogenous Tim-1 on primary T cells. Purified CD4+ T cells from C57BL/6 spleen and lymph node were stimulated for 24 h with the indicated concentrations of anti-CD3 Ab (and a fixed concentration of anti-CD28 Ab, 0.5 μg/ml), with or without anti-Tim-1 Ab (6 μg/ml), in the presence or absence of PI3K inhibitor LY294002 (10 μM). Cells were stained with fluorescently labeled Abs to CD69 (A) or CD25 (B) and analyzed by flow cytometry. Results in each panel are representative of three experiments that were performed.

FIGURE 6. p85 is required for Tim-1-mediated costimulation of CD69 expression. A. Expression of Tim-1 on resting or activated CD4+ T cells from spleens of p85β or p85αβ k.o. mice. Splenocytes from p85β (B) or p85αβ (C)-deficient mice were stimulated overnight with various concentrations anti-CD3 Ab (0.05, 0.2, and 1 μg/ml), along with a fixed concentration of anti-CD28 (1 μg/ml). Anti-Tim-1 Ab was also added to some cultures, at a concentration of 8 μg/ml. Cells were stained the next day for CD4 and CD69 and analyzed by flow cytometry. Results shown were gated on CD4+ cells.
alone. In light of data discussed above, we were interested in determining whether PI3K catalytic activity is required for Tim-1-mediated CD69 up-regulation. As shown in Fig. 5A, LY294002 significantly inhibited CD69 up-regulation induced by Tim-1 ligation, either alone or in conjunction with anti-CD3/CD28. Cells were harvested and stained for CD69 (A, C, and E) or CD25 (B, D, and F) expression, and analyzed by flow cytometry. Results in A and B, and E and F, are single experiments, representative of three, or two, separate experiments, respectively. C and D, Effects of Tim-1 costimulation (± SD) averaged over those experiments, with p values indicated.

Another inducible cell surface marker essential for T cell commitment to effector function is the high affinity IL-2 receptor α-chain (CD25). To evaluate whether Tim-1 selectively enhanced expression of the early activation Ag CD69, or whether this phenomenon was a more general consequence of its effects on T cell activation, we also examined CD25 up-regulation (Fig. 5B). As observed with CD69, Tim-1 cross-linking costimulated CD25 surface expression and this effect was more evident with lower concentrations of anti-CD3. Again, as with CD69, Tim-1 ligation alone led to some up-regulation of CD25 expression. Treatment with LY294002 significantly inhibited Tim-1-mediated CD25 up-regulation, both by itself and in conjunction with CD3/CD28 signals. Thus, data presented in Fig. 5 suggest that Tim-1 induces expression of the activation markers CD25 and CD69 in a PI3K-dependent manner.

To determine more conclusively the relevance of the p85 interaction for endogenous Tim-1 function, a mouse model of p85-deficiency mice was used (19). These mice were generated by crossing Pik3r1f mice, which have floxed alleles of Pik3r1 (encoding p85α and p50α), with Pik3r2n mice, which are deficient in the germline for Pik3r2 (encoding p85β). The resulting mice were then bred to Lck-Cre transgenic mice, yielding mice with T cells specifically lacking expression of PI3K regulatory subunits from the double negative stage of thymocyte development forward. T cells from p85β null mice develop normally and possess no detectable defects in PI3K signaling or any compensatory changes in PI3K isoform expression, so these mice served as controls (19, 29). p85 double k.o. mice also possess T cells with no gross developmental problems but have selective defects in TCR-dependent signaling and cytokine production (19).

We first assessed expression of Tim-1 on T cells from p85-deficient mice. Importantly, expression of Tim-1 was not impaired on p85β/−/− k.o. T cells, with expression in the resting (low levels) and activated (higher levels) state indistinguishable from p85−/− or wild-type mice (Fig. 6A). To confirm and extend the inhibitor studies performed with T cells from C57BL/6 mice, we examined co-stimulatory Tim-1 effects on surface expression of CD69 and CD25 in CD4−/− T cells from these p85 mutant mice. Representative flow cytometry plots for results obtained with CD69 are shown in Fig. 6B and C. Thus, while stimulation with anti-CD3/CD28 alone was relatively unaffected by the complete lack of p85 in this experiment (compare C with B), the ability of Tim-1 to costimulate CD69 up-regulation was severely impaired.

Because we and others have observed both stimulatory and co-stimulatory effects of Tim-1 ligation, we wanted to determine whether both or either of these activities would be affected by p85 deficiency. We therefore examined the effects of Tim-1 Ab 3B3 on up-regulation of CD69 and CD25 in CD4−/− T cells from these p85 mutant mice. Representative flow cytometry plots for results obtained with CD69 are shown in Fig. 7, A–D. Thus, while stimulation with anti-CD3/CD28 alone was relatively unaffected by the complete lack of p85 in this experiment (compare C with B), the ability of Tim-1 to costimulate CD69 up-regulation was severely impaired.

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paired in the p85 double k.o. cells, compared with T cells lacking cells and found that Tim-1 induction of IL-2 secretion was im-
expected, addition of IL-2 to the cultures had no effect on the differences did not underlie the effects we had observed. Thus, as
was largely corrected with the addition of exogenous IL-2 (19).

Although activation markers were assessed relatively early in the activation of surface activation markers and cytokines that are important for T cell activation and function.

The data we present here reveal a requirement for the Src family kinase Lck for Tim-1 tyrosine phosphorylation and subsequent NFAT/AP-1-dependent transcription. Our current model is that Lck first phosphorylates the cytoplasmic tail of Tim-1, which allows subsequent recruitment of signaling proteins to Y276, a residue that is required for Tim-1 function in T cell lines (7). It should be noted that our data are also consistent with a possible role for the related tyrosine kinase Fyn in the regulation of Tim-1 function. Thus, the JCaM.1 cell line, while lacking expression of Lck, also contains little, if any, Fyn protein (30). In addition, the kinase inhibitor PP2, although relatively specific for Src family kinases, cannot distinguish between possible functions for Lck or Fyn. Fi-
nally, in some SH2 domain array experiments, we noted weak binding of the Tim-1 phospho-Y276 peptide to the SH2 domain of Fyn. Further investigation is therefore warranted to address whether Fyn contributes to Tim-1 function.

Using an unbiased approach, we find that phosphorylation of Y276 leads to recruitment of the p85 subunits of PI3K, and we further confirm this interaction with endogenous p85 in T cells expressing Tim-1. Interestingly, interaction with p85 is a common feature shared by other T cell accessory molecules, such as CD28, ICOS, and CTLA-4. However, in contrast to these molecules, which preferentially bind the C-terminal SH2 domain of p85 (31–35), Tim-1 binds the amino-terminal SH2 domain, as revealed by the SH2 domain array. Another difference is that p85 binds to the CD28 family members via a consensus YYXM motif, whereas it binds to Tim-1 at a noncanonical sequence (YIVE). Although the amino-terminal p85 SH2 domain has a 10-fold lower avidity for the YYXM motif than does the C-terminal SH2 domain (31), work from Songyang and colleagues (36) suggests that YMXM is the preferred target for both domains. We have attempted to coimmuno-
noprecipitate an isolated N-terminal p85a SH2 domain construct with Tim-1, but have been unable to do so (unpublished data), possibly due to the low affinity of the interaction in the absence of other sequences. Further investigation will be required to more precisely define how the Tim-1-p85 complex forms and what other proteins may be involved. Nonetheless, our ability to coimmuno-
precipitate p85 with Tim-1 in a phosphotyrosine-dependent manner (Fig. 3) and the observed interaction between the purified N-
SH2 and peptide (again, phosphotyrosine-dependent; Fig. 2) suggest that this is a direct interaction.

Consistent with its recruitment of p85, Tim-1 requires PI3K and Akt activity to enhance T cell activation, because LY294002 and Akti 1/2 inhibit Tim-1 function. We also used a recently described genetic model, in which there is a complete lack of expression of the p85 regulatory subunits of PI3K selectively in T cells (19).

This is accompanied by a significant reduction in levels of the catalytic p110 subunit isoforms (which are normally stabilized by p85 binding), resulting in specific abrogation of class IA PI3K activity in T cells. Consistent with previous data (19), CD25 and CD69 up-regulation triggered by TCR stimulation was only mildly affected in the p85β and double k.o. T cells. However, the ability of Tim-1 to costimulate CD25 and CD69 is severely impaired in the double k.o. T cells, when compared with p85β-deficient T cells. Again, consistent with previous findings (19), p85β-deficient T cells behave identically to wild-type T cells with regard to CD25 and CD69 induction (data not shown).

T cell activation induced by Tim-1 alone was observed in our earlier studies performed in Jurkat T cells and is also in agreement

the p85α/β double k.o. T cells, had a profound effect on the ability of Tim-1 to induce increases in CD69 and CD25, either by itself or in conjunction with CD3/CD28 signaling. Previous studies with p85-deficient mice had demonstrated that the proliferation defect was largely corrected with the addition of exogenous IL-2 (19). Although activation markers were assessed relatively early in the course of T cell activation, we wanted to ensure that proliferation differences did not underlie the effects we had observed. Thus, as expected, addition of IL-2 to the cultures had no effect on the results (data not shown).

Finally, we examined cytokine production by the p85 mutant T cells and found that Tim-1 induction of IL-2 secretion was im-
paired in the p85 double k.o. cells, compared with T cells lacking p85β alone (Fig. 8). In contrast to the more potent stimulatory effects of Tim-1 Ab on CD69 and CD25 induction, Tim-1 crosslinking by itself had a small effect on IL-2 secretion (note difference in scales between A and B), which was partially im-
paired in the p85 α/β-deficient T cells (Fig. 8A). However, in the presence of sub-optimal anti-CD3/CD28 stimulation, Tim-1 liga-
tion significantly enhanced the production of IL-2, an effect that is severely impaired in the p85 α/β-deficient T cells (Fig. 8B).

**Discussion**

Recent studies have demonstrated that the transmembrane protein Tim-1 can influence the course of an immune response through direct effects on T cell activation. However, the biochemical sig-
naling pathways underlying these effects of Tim-1 ligation are not understood. Using a combination of biochemical, pharmacologi-
cal, and genetic approaches, we delineate a signaling pathway,

![FIGURE 8.](http://www.jimmunol.org/)

- **A** Anti-Tim-1 Only
- **B** Anti-CD3/CD28 +/- Anti-Tim-1

starting with Src kinase-mediated phosphorylation of Tim-1, to recruitment and activation of PI3K and Akt, which are required for induction of NFAT/AP-1-dependent transcription and up-regulation of surface activation markers and cytokines that are important for T cell activation and function.

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with other studies where hyperproliferation of T cells was observed in mice treated with soluble mTim-1 molecules, as well as activation of primary human cells treated with anti-Tim-1 Ab (6–8). A recent study has shown that Tim-1 can associate with the TCR/CD3 complex in the basal state, an interaction that is increased by Tim-1 or TCR/CD3 ligation (8). Tim-1 association with the TCR/CD3 complex could lead to aggregation of both, which may facilitate Tim-1 phosphorylation and activation of downstream signaling pathways through both TCR-dependent and -independent pathways. Indeed, we show that increasing amounts of agonistic anti-Tim-1 alone are accompanied by increased surface expression of CD69 and CD25. Although there is a significant dampening of this response in T cells from p85 double k.o. mice, higher concentrations of anti-Tim-1 are eventually sufficient to induce CD69 and CD25 expression. Though these T cells completely lack p85 regulatory subunits, proximal TCR-induced signaling seems relatively unaffected, because ZAP-70 phosphorylation is intact (19). Given that Tim-1 may also induce ZAP-70 phosphorylation through its interaction with the TCR (8), the residual effects of Tim-1 observed in p85-deficient T cells may be TCR-dependent.

Data presented here support the model that Tim-1 functions as a positive costimulatory molecule. However, anti-Tim-1 Abs targeting different epitopes of the Tim-1 extracellular domain have been shown to initiate either positive or negative effects on T cell activation (9, 10, 12). Thus, two anti-Tim-1 Abs, RMT1-10 and 3B3, differing by 17-fold in binding affinity to the same or closely related Tim-1 epitope in the IgV domain, exert opposite effects on T cell function (9). Although Tim-1 ligation by RMT1-10 apparently inhibits T cell activation, crosslinking with the higher affinity Ab 3B3 potentiates T cell activation. Recent structural studies also predict different ligand binding modes for the TIM family members (37, 38), consistent with direct and indirect evidence for multiple ligands for these proteins (2, 6, 7, 39–41). Thus, it is possible that more than one ligand might interact with Tim-1 concurrently on different faces of the molecule. Furthermore, homotypic interactions between individual Tim-1 molecules are mediated through their Ig domains, interactions that may be affected by glycosylation of the polymorphic mucin domain (38). The ultimate outcome of Tim-1 engagement may therefore depend upon the nature of the ligand(s) with which it interacts, as well as the timing and stoichiometry of such interactions. Differences in functional outcomes after Tim-1 ligation may result from divergent effects of Tim-1 ligation on T cell signaling pathways under such circumstances.

Although our studies indicate that PI3K plays an important role in mediating Tim-1 signaling, this does not exclude the possibility that additional molecules can also bind to Tim-1 and transmit signals to other downstream pathways. Based on data presented here, we suggest that ligation of Tim-1, when coupled with TCR stimulation, enhances downstream signaling pathways that induce T cell activation, at least in part through PI3K and its downstream effector Akt. Future studies will focus on the mechanisms by which Tim-1-mediated PI3K activation affects T cell activation.

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