**Vav Regulates Peptide-specific Apoptosis in Thymocytes**

By Young-Yun Kong,*‡§ Klaus-Dieter Fischer,* Martin F. Bachmann,** Sanjeev M. Ariathasan,‡§ Ivona Kozieradzki,*‡§ Mai P. Nghiém,*‡§ Dennis Bouchard,*‡§ Alan Bernstein,‡‡ Pamela S. Ohashi,*‡§ and Josef M. Penninger*‡§

From the *Amgen Institute, ‡Ontario Cancer Institute, §Department of Medical Biophysics and D Department of Immunology, and ID Department of Medical Genetics, University of Toronto, Toronto, O ntario, Canada M5G 2C1; the Institute for Radiation and Cell Research, University of W ürzburg, D-97078 W ürzburg, Germany; the **Basel Institute for Immunology, CH 4005 Basel, Switzerland; and the ‡‡Samuel Lunenfeld Research Institute, Mount Sinai Hospital, Toronto, Ontario, Canada M5G 1X5

**Summary**

The protooncogene Vav functions as a GDP/GTP exchange factor (GEF) for Rho-like small GTPases involved in cytoskeletal reorganization and cytokine production in T cells. Gene-targeted mice lacking Vav have a severe defect in positive and negative selection of T cell antigen receptor transgenic thymocytes in vivo, and vav-/- thymocytes are completely resistant to peptide-specific and anti-CD3/anti-CD28-mediated apoptosis. Vav acts upstream of mitochondrial pore opening and caspase activation. Biochemically, Vav regulates peptide-specific Ca2+ mobilization and actin polymerization. Peptide-specific cell death was blocked both by cytochalasin D inhibition of actin polymerization and by inhibition of protein kinase C (PKC).

Vav was found to bind constitutively to PKC-ø in thymocytes. Our results indicate that peptide-triggered thymocyte apoptosis is mediated via Vav activation, changes in the actin cytoskeleton, and subsequent activation of a PKC isoform.

**Key words:** Vav • negative selection • actin cytoskeleton • signaling transduction • protein kinase C

Engagement of the TCR initiates a cascade of molecular events resulting in tyrosine phosphorylation of cytoplasmic proteins, Ca2+ mobilization, activation of the mitogen-activated protein kinase (MAPK)1 and stress-activated protein kinase (SAPK) pathways, and reorganization of the cytoskeleton. The protooncogene product Vav is expressed in hematopoietic cells and is rapidly phosphorylated after activation of T cells by various growth factors or by cross-linking of antigen receptors (1–3). Vav contains a collection of structural motifs, including a pleckstrin homology (PH) domain, known to facilitate membrane localization; a calponin homology (CH) domain, involved in actin binding; one Src homology (SH)2 domain and two SH3 domains, known to mediate protein–protein interactions; and a Dbl homology (DH) domain (1, 4). In the protooncogene Dbl, the DH domain confers the capacity for guanine nucleotide exchange for the Rho family of small GTPases, which regulate cytoskeletal organization and SAPK/JNK signaling (5–10). Although Vav has been implicated in Ras and MAPK signaling (1, 11, 12), recent biochemical and genetic studies have established a role for Vav in the activation of Rac1 and other members of the Rho family of small GTPases (13–15).

T cells from vav-/- mice exhibit a block in cell cycle progression and fail to produce IL-2 in response to anti-CD3 cross-linking (16–18). Although Vav has no apparent role in TCR-mediated signaling pathways leading to MAPK or SAPK activation after CD3e cross-linking, Vav has been shown to regulate TCR-mediated Ca2+ flux and reorganization of the actin cytoskeleton. Consistent with this role, the functional defects observed in vav-/- T cells can be mimicked using the actin polymerization inhibitor cytochalasin D (CytD [19, 20]). In addition, Vav has been found...
to have a crucial role in thymocyte development and positive selection of both MHC class I- and MHC class II-restricted TCR transgenic (Tg) thymocytes (16-18, 21). However, it has been reported that superantigen-reactive and alloreactive vav-/- thymocytes could still undergo negative selection (21), suggesting a differential requirement for Vav in positive versus negative thymocyte selection.

To examine the role of Vav in the selection and activation of peptide-specific thymocytes, we introduced the H-Y TCR (22) and the lymphocytic choriomeningitis virus (LCMV) p33 peptide-specific P14 TCR transgenes (23) into a vav-/- background. We report that Vav is essential for peptide-specific clonal deletion and TCR-triggered apoptosis. Vav was found to regulate peptide-specific Ca$^{2+}$ mobilization and actin polymerization in thymocytes. Peptide-triggered apoptosis could be blocked using the actin polymerization inhibitor CytD and a global protein kinase C (PKC) blocker. Among all PKC isoforms tested, only PKC-@ was found to associate with Vav in thymocytes. These results suggest that TCR-mediated changes in the actin cytoskeleton and PKC-@ are crucial prerequisites for negative selection and peptide-triggered apoptosis.

### Materials and Methods

**Mice.** Gene-targeted mice made deficient in Vav by homologous recombination (19), and H-Y and P14 TCR Tg, and P14 Tg β2-microglobulin (β2m)-/- mice have been described previously (22, 24-27). Mice were screened for the Tg TCR αVβ8 chain. The TCR mutation was identified using genomic Southern blotting and PCR. Care of animals was in accordance with institutional guidelines.

**Reagents.** The PKC blockers R0-31-8220 (which blocks all PKC isoforms) and GF109023X (which inhibits the activity of PKC-@) were purchased from Calbiochem Corp. (La Jolla, CA [28]). The phosphatidylinositol 3'-kinase (PI3K)-specific inhibitor Wortmannin and the fungal metabolite CytD (which blocks actin polymerization) were obtained from Sigma Chemical Co. (St. Louis, MO). Phalloidin binds specifically to polymerized filamentous actin (F-actin) and was directly labeled with FITC (Sigma Chemical Co.).

**Induction of Apoptosis.** Freshly isolated thymocytes from C57BL/6 mice were cultured in RPMI 1640 medium containing 10% FCS and 10⁻⁵ M β-mercaptoethanol in the absence or presence of dexamethasone (Sigma Chemical Co.), anti-CD95 (clone Jo91; PharMingen, San Diego, CA), anti-CD3e (clone 145-2C11; Pharmingen), anti-CD3e plus anti-CD28 (clone 37.51), or PMA (12.5 ng/ml) plus Ca²⁺ ionophore A23187 (100 ng/ml), for different time periods and at different concentrations as indicated in figure legends. Optimal concentrations and activation regimes for the induction of apoptosis were determined in pilot studies (29, and data not shown).

**Immunocytochemistry.** Blood samples (20 µl) were collected in heparinized capillary tubes and washed once in immunofluorescence staining buffer (1% FCS, 0.01% NaN₃ in PBS). Single cell suspensions of thymocytes, spleen cells, and mesenteric lymph node cells were prepared as described (29), resuspended in PBS, and incubated with the appropriate mAbs for 30 min at 4°C. The following mAbs were used: anti-CD4 (FITC- or PE-labeled; PharMingen); anti-CD8 (FITC- or PE-labeled, or biotinylated; PharMingen); anti-pan TCR-α/β (FITC- or PE-labeled; PharMingen); anti-TCR Vβ8.1 and 8.2 (clone K16, PE-labeled or biotinylated; Pharmingen); anti-H-2Kb (FITC-labeled or biotinylated; Pharmingen); anti-H-2Kb (FITC-labeled or PE-labeled; Pharmingen); anti-H-2Db (biotinylated; Pharmingen); anti-CD8 (biotinylated; Pharmingen); anti-CD3e (FITC- or PE-labeled; Pharmingen); anti-Vav (FITC- or PE-labeled; Pharmingen); anti-CD69 (biotinylated; Pharmingen); anti-CD45RA (biotinylated; Pharmingen); and anti-TCR Vγ1Vδ2 (biotinylated; Pharmingen). All staining combinations were as indicated in figure legends. Biotinylated mAbs were visualized with Streptavidin-R-ED670 (GIBCO BRL, Gaithersburg, MD), and samples were analyzed using a FACScan® and Lysis II software (Becton Dickinson, Mountain View, CA).

**Negative Selection In Vitro.** Thymocytes were purified from vav-/-, vav-/-, and vav-/- P14 Tg mice. P14 Tg mice express an αβ TCR (TCR Vα2Vβ8) specific for the strong agonist p33 peptide of LCMV. Thymocytes were cultured on EL4 cells (H-2b/b) in RPMI medium containing 5% FCS and 10⁻³ M β-mercaptoethanol (2 ml final vol/well of 24-well flat-bottomed plates; Costar Corp., Cambridge, MA). EL4 cells (H-2b/b) were pulsed with different concentrations of the deleting p33 peptide or the weak agonist p33 peptide analogue 8.1 for 2 h before coculture with thymocytes (26, 27, 30). Thymocytes were harvested after 10, 16, and 22 h of incubation and stained with anti-CD4-PE, anti-CD8-PFITC, or biotinylated; anti-TCRVα1 (FITC-labeled or biotinylated; PharMingen); anti-TCRVβ1 (FITC-labeled; PharMingen); anti-CD3 (FITC-labeled or biotinylated; PharMingen); anti-CD28 (biotinylated; PharMingen); or anti-HSA (biotinylated; PharMingen). All staining combinations were as indicated in figure legends. Biotinylated mAbs were visualized with Streptavidin-R-ED670 (GIBCO BRL, Gaithersburg, MD), and samples were analyzed using a FACScan® and Lysis II software (Becton Dickinson, Mountain View, CA).

**Signal Transduction.** Thymocytes were isolated from nonselecting β2m⁻/⁻ vav-/-, positively selecting vav-/-, and vav-/- P14 Tg mice and cultured on a monolayer of confluent and adherent MC57G fibroblasts (H-2b/b) in RPMI medium containing 5% FCS and 10⁻³ M β-mercaptoethanol. MC57G APCs were pulsed with different concentrations of the agonist LCMV-p33 peptide or the control AV peptide for 2 h before coculture with thymocytes. Nonadherent thymocytes (2 × 10⁶ P14 Tg thymocytes/well) were harvested after 15, 30, 60, and 180 min of
cocculture with adherent fibroblasts and assayed for MAPK and SAPK activation, nuclear factor κB (NF-κB) activation (IkB phosphorylation), and phosphotyrosine signaling (19). In brief, tyrosine phosphorylation was monitored in total cell lysates and immunoprecipitates using an antiphosphotyrosine mAb (Upstate Biotechnology, Inc., Lake Placid, N.Y.). Activation of SAPKs and MAPKs was detected using phospho-SAPK- and phospho-MAPK-specific Abs (New England Biolabs Inc., Beverly, MA). The levels of SAPK and MAPKs (extracellular signal–regulatory kinase [ERK]1/2) were determined by immunoblotting (New England Biolabs Inc.). Phosphorylated IkB was detected using an mAb specific for phosphoserine (Ser32) of IkBα (New England Biolabs Inc.). Ser32 phosphorylation of IkBα targets IkB for degradation and is essential for the release of active NF-κB (34).

Immunoprecipitation. T lymphocytes and peripheral lymph node T cells were activated with anti-CD3ε (1 μg/ml) for different time periods as indicated in figure legends. Cells (10^7/lane) were harvested and lysed, and Vav was immunoprecipitated using an anti-Vav1 Ab reactive against amino acids 576–589 (Santa Cruz Biotechnology Inc., Santa Cruz, CA). Immunocomplexes were separated by SDS-PAGE, transferred onto a membrane, and Western blotted to detect the presence of Vav and the PKC isozymes PKC-α, -β (I-βII, -βII), -γ, -δ, -ε, -ζ, -θ, -ι, -λ, -μ, and receptor for activated C-kinase (RACK1). In parallel, total cell lysates (2 × 10^6 cells/lane) were Western blotted to ascertain the relative protein levels of these PKC isozymes in thymocytes. PKC isozyme Abs were purchased from Transduction Laboratories (Lexington, KY).

Peptide-specific Ca2+ Mobilization. Peptide-specific Ca2+ mobilization ([Ca2+]i) in P14 Tg thymocytes was determined as described (35). In brief, freshly isolated P14 Tg thymocytes (2 × 10^6) were loaded with 3 mM INDO-1 (Molecular Probes Inc., Eugene, OR) in IMDM (pH 7.4) for 4 h at 37°C. Thymocytes were then incubated with peptide-presenting EL4 cells loaded with either p33 or AV peptide and centrifuged (1,500 rpm, 4°C) to allow conjugate formation between thymocytes and EL4 cells. Increases in intracellular free Ca2+ were recorded in real time on live-gated thymocyte–EL4 conjugates using a FACScalibur® (Becton Dickinson).

A din Polymerization. T lymphocytes were preincubated with PMA (12.5 ng/ml), and EL4 cells were loaded with either p33 or AV peptide (10 μg/ml) for 30 min on ice. Actin polymerization was initiated by placing the cells at 37°C for different time periods as indicated in figure legends. Actin polymerization was stopped by addition of 4% paraformaldehyde. After fixation, cells were incubated for 30 min with FITC-labeled phalloidin, which specifically binds to polymerized F-actin. Cells were washed three times in PBS, and phalloidin staining was analyzed using a FACScan®. Percentage of peptide-specific actin polymerization (percent F-actin) was calculated as (mean value of phalloidin fluorescence intensity of p33-activated cells at a certain time point)/(mean value of phalloidin fluorescence intensity of AV-activated cells at the same time point) × 100.

Results

Impaired TCR-mediated Positive Selection of vav⁻/⁻ T lymphocytes. To analyze the role in vivo of Vav in thymocyte selection, we introduced two rearranged TCRα/β transgenes, H-Y and P14, into a vav⁻/⁻ background. The H-Y TCR recognizes a male-specific peptide in the context of MHC class I. Thymocytes expressing the H-Y TCR are positively selected in female H-2k mice (22, 24, 25). The P14 TCR is specific for a peptide epitope of the LCMV glycoprotein p33 in the context of the MHC haplotype H-2D^b (23). In positively selecting vav⁻/⁻ mice, the development of P14⁺ and H-Y⁺ thymocytes is skewed towards the CD8⁺ lineage (Fig. 1, A and B). In vav⁻/⁻ mice, this bias in favor of mature CD8⁺ thymocytes did not occur in either H-Y Tg or P14 Tg thymocytes (Fig. 1, A and B), and development was in fact blocked at the immature CD4⁺/CD8⁺ stage of differentiation. Consistent with a block in positive selection, TCR Tg vav⁻/⁻ thymocytes expressed high levels of HSA and did not upregulate surface expression of the maturation and selection markers CD69, CD5, CD5 (Fig. 1C), H-2Kb, or CD45RB (not shown). Expression of the P14 Tg TCR Vβ8 and TCR Vα2 chains was significantly lower in immature CD4⁻/CD8⁺ thymocytes of vav⁻/⁻ mice compared with CD4⁺/CD8⁺ thymocytes from vav⁻/+ mice (Fig. 1C). Similarly, expression of the H-Y Tg-specific TCR Vβ8 and TCR Vα3 chains detected by the T3.70 Ab was significantly lower in H-Y Tg vav⁻/⁻ mice (not shown). Importantly, P14 Tg vav⁻/⁻ (and H-Y Tg vav⁻/⁻ [not shown]) thymocytes displayed a phenotype that is similar to that of P14 Tg (and H-Y) vav⁻/+ β2m⁻/⁻ mice, which have a defect in positive selection of MHC class I-restricted thymocytes (Fig. 1C). These results show that Vav regulates the positive selection of MHC class I-restricted thymocytes and confirm previous data demonstrating the crucial role of Vav in positive thymocyte selection (16–18, 21).

Impaired Negative Selection and Peptide-specific Apoptosis. To examine the role of Vav in peptide-specific clonal deletion in vivo and in vitro, we investigated the negative selection of H-Y and LCMV TCR Tg thymocytes. In the H-Y Tg system, thymocytes expressing the H-Y TCR are positively selected in female H-2k mice but negatively selected in male H-2b mice. Negative selection in male H-Y Tg mice results in a small thymus due to deletion of CD4⁻/CD8⁺ thymocytes (24; Fig. 1B). However, male vav⁻/⁻ mice contained a large number of CD4⁺/CD8⁺ thymocytes, indicating that negative selection of H-Y Tg thymocytes was severely impaired in the absence of Vav.

To further elucidate the requirement for Vav in negative selection, an in vitro model of peptide-specific negative selection was used (30). In the LCMV-p33 peptide system, APCs are loaded with different concentrations of the LCMV glycoprotein strongly agonist p33 or the weak agonist p33 analogue 8.1. Although vav⁻/⁻ P14 Tg thymocytes readily underwent apoptosis in a dose-dependent manner in response to treatment with either the p33 or 8.1 peptide (Fig. 2A), vav⁻/⁻ P14 Tg thymocytes were completely resistant to peptide-mediated apoptosis, even at very high peptide concentrations (Fig. 2B). Since P14 vav⁻/⁻ thymocytes are blocked in positive T cell selection, we used β2m⁻/⁻ vav⁻/+ P14 TCR Tg mice (H-2b) as an additional control since these mice have a block in the positive selection of the P14 TCR due to deletion of the MHC class I ligand (β2m⁻/⁻). The kinetics and extent of
p33-induced apoptosis of $\beta_{2m}^{-/-}$ CD4$^+$CD8$^+$ P14 Tg thymocytes were similar to those of vav$^{-/-}$ P14 Tg mice, indicating that the defect of P14 Tg vav$^{-/-}$ thymocytes to undergo peptide-specific apoptosis was not due to differences in the composition of thymocyte populations (Fig. 2C). These results show that Vav is required for the negative selection of peptide-specific thymocytes.

vav$^{-/-}$ thymocytes are resistant to CD3/CD28-mediated apoptosis. Immature CD4$^+$CD8$^+$ thymocytes are highly susceptible to cell death induced by many apoptotic stimuli (36). We evaluated the ability of vav$^{-/-}$ and vav$^{-/-}$ thymocytes to undergo apoptosis in response to the following stimuli: dexamethasone, PMA/Ca$^{2+}$ ionophore, anti-CD95 (FAS), anti-CD3e, and anti-CD3e/anti-CD28 (29, 37). Although treatment with anti-CD3e/anti-CD28 induced the cell death of vav$^{-/-}$ CD4$^+$CD8$^+$ thymocytes, vav$^{-/-}$ thymocytes were strikingly resistant to anti-CD3e/anti-CD28-triggered apoptosis (Fig. 3). No significant differences were observed between vav$^{-/-}$, vav$^{-/-}$, and vav$^{-/-}$ thymocytes in the extent or kinetics of cell death in response to anti-CD95 (FAS), PMA plus Ca$^{2+}$ ionophore (Fig. 3), or dexamethasone (not shown), implying that the cellular apoptotic machinery is functional in the absence of Vav. These data indicate that Vav is a crucial signal transduction molecule involved in TCR-mediated thymocyte apoptosis.

Vav Links TCR Signaling to Mitochondrial ($\Delta W_m$) Disruption. Disruption of the mitochondrial transmembrane potential ($\Delta W_m$) due to the opening of mitochondrial membrane pore complex has been identified as the earliest common denominator of apoptosis (36). Alterations in mitochondria lead to the release of "apoptosis-inducing factor" (AIF) or cytochrome c, resulting in the activation of effector caspases such as caspase 3 (Cp32) (32, 38–41). To assess whether Vav links TCR signaling to the opening of mitochondrial pores, we determined the mitochondrial transmembrane potential

Figure 1. Impairment of positive and negative thymocyte selection in vav$^{-/-}$ mice. (A) Impaired positive selection in P14 Tg vav$^{-/-}$ mice. TOP panels) and P14 TCR-$/\alpha\beta$ Tg (bottom panels) thymocytes were isolated from 6-wk-old H-2$^{b}$ vav$^{-/-}$ and vav$^{-/-}$ mice and stained with anti-CD4-PE and anti-CD8-FITC. Percentages of positive cells within quadrants are indicated. Total thymocyte numbers were as follows (mean values ± SD; n = 3; 6 wk of age): vav$^{-/-}$ non-Tg: 112 ± 5.2 × 10$^6$; vav$^{-/-}$ non-Tg: 1.2 ± 0.6 × 10$^6$; P14 Tg: 12.0 ± 3.4 × 10$^6$; vav$^{-/-}$ P14 Tg: 7.2 ± 0.6 × 10$^6$. 1 result representative of 10 experiments is shown. (B) Impaired selection of the Tg H-Y TCR in vav$^{-/-}$ mice. Thymocytes were isolated from 5-wk-old positively selecting female (top panels) and negatively selecting male (bottom panels) H-2$^{b}$ H-Y TCR Tg mice and stained with anti-CD4-PE and anti-CD8-FITC. Percentages of positive cells within quadrants are indicated. One result representative of five experiments is shown. Total thymocyte numbers were as follows (mean values ± SD; n = 3; 6 wk of age): vav$^{-/-}$ female: 19.6 ± 3.3 × 10$^6$; vav$^{-/-}$ female: 7.5 ± 0.9 × 10$^6$; vav$^{-/-}$ male: 3.6 ± 0.8 × 10$^6$; vav$^{-/-}$ male: 13.2 ± 2.6 × 10$^6$. (C) Phenotype of CD4$^+$CD8$^+$ thymocytes. Thymocytes were isolated from P14 Tg vav$^{-/-}$, P14 Tg vav$^{-/-}$, and P14 Tg vav$^{-/-}$/vav$^{-/-}$ mice and triple stained using anti-CD4 (PE), anti-CD8 (FITC), and a third biotinylated Ab reactive to TCRV8, TCVa2, CD28, CD69, CD5, or HSA. Histograms are shown for the indicated surface markers (solid line) on gated CD4$^+$CD8$^+$ thymocytes. Broken lines, background staining using unspecific Abs. One result representative of three experiments is shown.
Vav regulates peptide-specific apoptosis of P14 TCR Tg thymocytes. P14 Tg thymocytes were purified from vav−/− (A), vav−/− (B), and vav+/+ β2m−/− mice and cultured on EL4 (H-2Dk) APCs pulsed with the indicated concentrations of p33 or 8.1 peptides. Cells were harvested after 22 h of culture, and cell death was determined by FACs+staining using anti-CD4-PE, anti-CD8-FITC, and a vital dye, either 7-AAD or annexin V. Percent survival was determined from the number of viable CD4+CD8+ cells remaining after culture in a given concentration of p33 or 8.1 peptide compared with the number of control thymocytes remaining after culture in an equal concentration of the control peptide AV. The AV peptide has high affinity for H-2Dk but very low affinity for the P14 TCR. Peptide concentrations on the x-axis indicate a range of 10−6 to 10−11 M. No result representative of five experiments is shown.

Figure 3. vav−/− thymocytes are resistant to anti-CD3/anti-CD28-mediated apoptosis. Cell death was induced in vav+/− and vav−/− thymocytes by treatment with anti-CD3ε, anti-CD28, anti-CD95 (FAS), or PMA plus Ca2+ ionophore. Percentage of viable CD4+CD8+ thymocytes remaining after treatment is indicated on the y-axis. The mean result of a triplicate culture (±SD) representative of three independent experiments is shown for each activation.

(A) (% viable CD4+CD8+ cells)
with CytD (Fig. 5 E) or chelation of extracellular Ca\(^{2+}\) by EGTA (Fig. 5 F) significantly decreased p33 peptide-triggered Ca\(^{2+}\) mobilization. The extent of the CytD-mediated decrease in peptide-specific Ca\(^{2+}\) flux was dependent on the concentration of the p33 peptide (not shown). Treatment of P14 thymocytes with CytD had no apparent effect on the extent of tyrosine phosphorylation, NF-κB activation, or MAPK activation (not shown). These results...
suggest that peptide-specific actin polymerization has an important role in the extent and duration of TCR-mediated Ca\(^{2+}\) mobilization in thymocytes.

Inhibition of actin polymerization blocks peptide-specific thymocyte apoptosis. To further elucidate the role of Vav-regulated changes in the actin cytoskeleton during negative selection, we tested whether the actin polymerization inhibitor CytD could interfere with peptide-specific apoptosis of P14 TCR Tg thymocytes. Interestingly, CytD blocked the peptide-specific cell death of vav\(^{+/-}\) P14 thymocytes in a dose-dependent manner (Fig. 7). Similarly, anti-CD3/anti-CD28-mediated, but not dexamethasone- or CD95-mediated, apoptosis of vav\(^{+/-}\) thymocytes could be inhibited by CytD (not shown). These data suggest that TCR-mediated actin polymerization plays a role in peptide-specific apoptosis of thymocytes.

**Figure 6.** Impaired peptide-specific actin polymerization (F-actin formation) in P14 vav\(^{+/-}\) thymocytes. Thymocytes from P14 vav\(^{+/-}\) (diamonds) and vav\(^{+/-}\) (squares) littermates were isolated and activated with EL4 APCs loaded with the deleting p33 peptide (10\(^{-9}\) M) or the control AV peptide (10\(^{-9}\) M) for the indicated time periods. Percentage of actin polymerization (mean percent increase in F-actin ± SD) is shown on the y-axis. Values were calculated as described in Materials and Methods. The x-axis shows the kinetics of induction in minutes after p33 stimulation. One result representative of three independent experiments is shown.

**Figure 7.** Inhibition of actin polymerization blocks peptide-specific thymocyte apoptosis. (A) The actin polymerization inhibitor CytD blocks peptide-specific apoptosis of P14 TCR Tg vav\(^{+/-}\) thymocytes induced by different concentrations of the high-affinity p33 peptide. The percentage of viable CD4\(^{+}\)CD8\(^{+}\) cells remaining after treatment is shown on the y-axis. Peptide concentrations are indicated on the x-axis and range from 10\(^{-8}\) to 10\(^{-9}\) M. The concentrations of CytD used to inhibit thymocyte apoptosis are indicated. (B) Dose-response curve of CytD inhibition of peptide-specific apoptosis of P14 Tg vav\(^{+/-}\) thymocytes. Percent inhibition of apoptosis is shown on the y-axis. The concentration of the deleting p33 or control AV peptide was 10\(^{-9}\) M. Apoptosis of CD4\(^{+}\)CD8\(^{+}\) thymocytes was detected as described in Materials and Methods. One result representative of six experiments is shown.

**Figure 8.** PKC activation can restore peptide-specific apoptosis of P14 vav\(^{+/-}\) thymocytes (A and B) The specific PKC activator PMA, but not Ca\(^{2+}\) ionophore, can rescue peptide-specific apoptosis of vav\(^{+/-}\) thymocytes. P14 vav\(^{+/-}\) thymocytes (2 × 10\(^{6}\) well) were incubated with EL4 APCs loaded with 10\(^{-9}\) M AV or p33 peptide plus varying concentrations of the Ca\(^{2+}\) ionophore A23617 (A) or PMA (B). Higher concentrations of the Ca\(^{2+}\) ionophore A23617 plus p33 peptide also failed to trigger apoptosis (not shown). Viability of CD4\(^{+}\)CD8\(^{+}\) thymocytes was assessed 16 h after activation as described in Materials and Methods. It should be noted that PMA alone induced apoptosis of CD4\(^{+}\)CD8\(^{+}\) vav\(^{+/-}\) thymocytes (not shown). One result representative of three independent experiments is shown. (C) Inhibition of p33 peptide-mediated apoptosis of P14 TCR Tg vav\(^{+/-}\) thymocytes by the global PKC inhibitor RO-31-8220. Apoptosis was induced by treatment with 10\(^{-9}\) M p33 peptide in the presence of the global PKC blocker RO-31-8220, GF109203X (which only inhibits the activity of the Ca\(^{2+}\)-dependent PKC-\(\alpha\), -\(\beta\), and -\(\gamma\) isoforms), or Wortmannin (a specific PI3K blocker). The percentage of apoptotic cells is shown on the y-axis. Concentrations of pharmacological inhibitors are shown on the x-axis (\(\mu\)M). Similar results were obtained using different concentrations of p33 peptide or higher concentrations of the inhibitors (not shown). One result representative of five different experiments is shown. (D and E) Inhibition of p33 peptide-mediated or PMA-mediated apoptosis of P14 TCR Tg vav\(^{+/-}\) thymocytes by kinase or actin polymerization blockers (D) T thymocytes were treated with p33 peptide (10\(^{-9}\) M) in the absence or presence of RO-31-8220 (RO, 1 \(\mu\)M), the actin polymerization blocker CytD (2.5 \(\mu\)M), or Wortmannin (WT, 1 \(\mu\)M). (E) T thymocytes were treated with PMA (12.5 nM) in the absence or presence of RO-31-8220, CytD, or Wortmannin at the concentrations described for D. Unstimulated thymocytes served as controls for both D and E. Results for D and E are shown as mean percent viable CD4\(^{+}\)CD8\(^{+}\) cells ± SD. Percentage of apoptosis was calculated as described in Materials and Methods. One result representative of three different experiments is shown.

Cooperation between Vav and PKC in Negative Selection. The primary biochemical defects associated with signaling in peptide-specific vav\(^{+/-}\) thymocytes were impaired Ca\(^{2+}\) flux and reduced actin polymerization. Addition of Ca\(^{2+}\) ionophore, which restores Ca\(^{2+}\) flux in P14 vav\(^{+/-}\) and
vav-/- thymocytes (not shown), did not restore p33 peptide-mediated apoptosis even when used at very high concentrations (100 ng/ml; Fig. 8 A). In contrast, activation of vav-/- P14 Tg cells with the deleting p33 peptide plus activation of PKC via the phorbol ester PMA significantly shifted the dose-response curve of cell death in a dose-dependent manner (Fig. 8 B). Thus, activation of PKC via PMA, but not rescue of Ca\(^{2+}\) flux by Ca\(^{2+}\) ionophore, was able to restore peptide-specific apoptosis in P14 vav-/- thymocytes, suggesting that the Vav and PKC signaling cascades cooperate in the induction of TCR-mediated thymocyte apoptosis. However, it should be noted that our results do not preclude a role for Ca\(^{2+}\) elevation in thymocyte selection and clonal deletion in vivo.

Recently, it has been shown that in order for Vav to act as a GDP/GTP exchange factor (GEF) for Rac, RhoA, or CDC42, PI3K activity and the binding of PI3K-generated phospholipid products to the PH domain of Vav are required. Therefore, we investigated whether the specific PI3K inhibitor Wortmannin could block peptide-specific apoptosis. As shown in Fig. 8, C and D, the inhibition of PI3K did not affect the apoptosis of P14 thymocytes triggered by the deleting p33 peptide, suggesting that the role of Vav in thymocyte apoptosis is independent of PI3K phospholipid-dependent Vav activation. However, inhibition of PKC by the compound RO-31-8220 (which blocks all PKC isoforms) prevented peptide-specific apoptosis in wild-type thymocytes (Fig. 8, C and D). Peptide-specific apoptosis of P14 thymocytes was not blocked by the pharmacological inhibitor GF109203X, which prevents activation of Ca\(^{2+}\)-dependent PKC isoforms (Fig. 8 C). PMA-triggered apoptosis was blocked by the global PKC blocker RO-31-8220 but not by CytD or Wortmannin (Fig. 8 E). These data indicate that Vav links TCR signaling to activation of a Ca\(^{2+}\)-independent PKC isoform required for the induction of peptide-specific thymocyte apoptosis.

To further evaluate the link between Vav and PKC, we immunoprecipitated Vav from wild-type thymocytes and analyzed the binding of various PKC isoforms to Vav. Surprisingly, Vav coimmunoprecipitated with the Ca\(^{2+}\)-independent PKC isoform PKC-\(\theta\) but not with any other PKC isoform tested (Fig. 9, left). The association between Vav and PKC-\(\theta\) did not change after CD3 cross-linking, suggesting that Vav/PKC-\(\theta\) binding is constitutive in thymocytes. Constitutive Vav/PKC-\(\theta\) association was also observed in mature peripheral T cells (not shown). Our results do not preclude associations between Vav and low abundance PKC isoforms.

**Discussion**

Our genetic and functional analyses of vav-/- mice show that Vav is a crucial regulator both of positive and negative selection of CD8\(^{+}\) thymocytes and the peptide-triggered apoptosis of developing T cells. Vav is involved as a regulator of TCR-mediated cytoskeletal reorganization and Ca\(^{2+}\)...
mobilization after peptide-specific activation by APCs. Our results suggest that peptide-triggered thymocyte apoptosis is mediated via Vav activation, changes in the actin cytoskeleton, and subsequent activation of a PKC isoform. This hypothesis is based on the following findings: (a) Vav-deficient thymocytes do not undergo peptide-MHC-mediated cell death in vitro or in vivo; (b) vav-/- thymocytes exhibit a defect in actin polymerization, and inhibition of cytoskeletal changes by CytD blocks peptide-MHC-mediated and anti-CD3/anti-CD28-mediated thymocyte apoptosis. However, inhibition of actin polymerization does not inhibit dexamethasone-, CD95-, or PKC-mediated thymocyte apoptosis; (c) activation of PKC restores the susceptibility to apoptosis of p33 peptide-triggered Tg vav-/- thymocytes; and inhibition of a C2a-independent PKC isoform inhibits TCR-mediated thymocyte apoptosis and (d) Vav associates constitutively with PKC-ð but not with any other PKC isoform. Thus, the PKC isoform PKC-ð is a good candidate for the effector kinase of negative thymocyte selection.

The PKCs are a family of 11 phospholipid-dependent serine/threonine kinases. These closely related isoenzymes differ in their structural and biochemical properties, tissue distribution, subcellular localization, and substrate specificity (45, 46). According to primary structure and binding to Ca2+ or phorbol esters, different PKC subgroups exist: conventional PKCs (α, βI, βII, and γ) bind Ca2+ and are activated by PMA; novel PKCs (δ, ε, θ, and η) are activated by PMA but do not bind Ca2+; atypical PKCs (ζ, λ, ι, and μ) bind to diacylglycerol (DAG) but are not activated by PMA or Ca2+ ionophores. Activation of PKC molecules by lipid second messengers requires membrane recruitment (47). T cell activation requires PKC activity. Previously, it has been reported using PCR and Northern blotting that PKC-α, -β, -γ, -ε, -ζ, and -θ are expressed in thymocytes (48–50). Our study shows for the first time that all PKC isoforms are expressed in thymocytes, albeit at different levels. Although most PKC isoforms are expressed ubiquitously, the isoform PKC-θ is predominantly expressed in the hematopoietic system, particularly in T cells (48–50), and has been placed upstream of IL-2 transactivation and AP1 (Fos/Jun) activity in T lymphoma cells (51, 52). Recently, it has been shown in Jurkat cells that calcineurin and PKC-θ cooperate in inducing IL-2 gene transcription (53), a function that is reminiscent of the coordination of IL-2 gene expression by calcineurin and Vav/Rac1 (20, 54–57). Interestingly, the small Ras-like molecules Rac1, CDC42, and Rho, whose activation is regulated by Vav, may act upstream of PKC (58, 59), suggesting that Vav, Rac1, and PKC-θ may form a complex. Although Vav and PKC-θ clearly associate in thymocytes and peripheral T cells, Rac1, CDC42, and RhoA do not coimmunoprecipitate with these complexes.

Vav and Vav-regulated Rac1 and CDC42 have been previously shown to mediate changes in the actin cytoskeleton and to activate SAPK/JNK K (13–15, 60, 61). Similarly, PKC-θ has been placed upstream of SAPK/JNK in T cell lines (53). However, TCR/CD28-mediated SAPK/JNK activation appears normal in vav-/- thymocytes and mature T cells after anti-CD3 cross-linking (19, 20). Similarly, a genetic mutation in the SAPK/JNK signaling pathway had no effect on the induction of thymocyte apoptosis (29), suggesting that Vav and perhaps PKC-ð-mediated negative selection are independent of SAPK/JNK. Importantly, although SAPK/JNK activity is readily induced in thymocytes in response to anti-CD3 and anti-CD28 Abs (62), we failed to detect SAPK/JNK activation in thymocytes after specific peptide-MHC-mediated stimulation. Thus, Vav may define a novel signaling pathway that leads to negative thymocyte selection and IL-2 production in peripheral T cells. This signaling pathway appears to be independent of MAPK, SAPK/JNK, or NF-κB activation.

Rac1, RhoA, or CDC42 activation by Vav requires PI3K activity in vitro (44). However, inhibition of PI3K has no apparent effect on the peptide-triggered thymocyte apoptosis of thymocytes, suggesting either that the role of Vav in thymocyte apoptosis is independent of its GTP exchange activity for Rac and CDC42, or that Vav can function as an exchange factor even in the absence of PI3K activity in vivo. Similarly, it has been shown that a pharmacological PKC inhibitor, but not the PI3K inhibitor Wortmannin, can block anti-CD3/anti-CD28-mediated thymocyte apoptosis (63). These latter results are consistent with our observation that Vav and PKC regulate anti-CD3/anti-CD28-mediated apoptosis and imply that PKC activation is a crucial signal in the pathway of TCR-triggered apoptosis in developing thymocytes. Whether PKC-θ and/or other PKC isoforms are the crucial downstream kinases in thymocyte selection needs to be determined in genetic experiments.

Both we (19) and Holsinger et al. (20) have previously shown that Vav regulates Ca2+ mobilization and actin reorganization in peripheral T cells after anti-CD3 Ab cross-linking. The results of the present study provide the first genetic evidence that Vav is also a crucial regulator of the TCR-mediated Ca2+ flux and actin polymerization required for thymocyte selection of peptide-MHC-stimulated thymocytes. In particular, actin polymerization was found to be crucial for the induction of apoptosis. Various cytoskeletal proteins and regulators of cytoskeletal changes, such as gelsolin (64–66), β-catenin (67), PAK2 (68), actin (67, 69–76), fodrin (77), and Gα2 (76), are proteolytically cleaved after the induction of apoptosis. Cleavage of cytoskeletal proteins, which occurs after caspase activation, has an important role in the death effector phase of apoptosis, particularly in the regulation of membrane alterations, morphological changes, and DNA fragmentation (68, 69). In contrast, inhibition of actin polymerization by CytD resulted in impaired TCR-mediated apoptosis. These cytoskeletal changes were found to be upstream of the opening of the mitochondrial membrane pore complex and caspase activation. However, CytD could not block apoptosis after PKC activation (which induces rapid changes in the actin cytoskeleton) and CytD could not protect thymocytes
from dexamethasone- or CD95-mediated apoptosis, suggesting that actin changes per se are not the principle mechanism for conveying a cell death signal in thymocytes. Rather, we propose that TCR-mediated and Vav-regulated changes in the actin cytoskeleton lead to the recruitment of PKC-\(\mu\) to the site of contact between T cells and APCs. All other PKC isoforms (\(\alpha, \beta_1, \delta, \eta, \) and \(\zeta\)) are excluded from the contact site, suggesting that PKC-\(\mu\) has a specific role in T cell activation (78). Since Vav regulates TCR clustering and PKC-\(\mu\) is found in these clusters, Vav-regulated actin polymerization may be required for recruitment of PKC-\(\mu\) to the TCR and activation of PKC at the cell membrane. How Vav and possibly PKC-\(\mu\) link TCR-mediated signals to mitochondrial apoptosis and caspase 3 activation remains to be determined. Importantly, our results show that Vav and actin polymerization also regulate the extent and duration of peptide-specific \(Ca^{2+}\) mobilization, an event thought to be involved in thymocyte apoptosis and clonal selection (79–83).

Our data show that Vav regulates positive and negative selection of MHC class I–restricted thymocytes. We have previously reported that the progression of CD4\(^+\)CD8\(^-\) T cell precursor cells to CD4\(^+\)CD8\(^+\) thymocytes is impaired in vav\(^-/-\) mice on a C57BL/6 (H-2b) background but not on a C57BL/6 (H-2d) background (19). Thus, Vav clearly has a role in preTCR-mediated expansion of early thymocytes, but the absence of Vav can be partially compensated by other signaling molecules. However, this compensatory mechanism does not appear to be operational or sufficient to rescue positive and negative selection of TCR Tg thymocytes in vav\(^-/-\) mice. Thus, Vav either plays a different role in TCR signaling at different stages of development and/or different compensatory mechanisms are available to CD4\(^-\)CD8\(^-\) versus CD4\(^+\)CD8\(^-\) thymocytes that make up for the Vav mutation in the expansion from CD4\(^-\)CD8\(^-\) to CD4\(^+\)CD8\(^+\) cells but cannot compensate for selection of CD4\(^+\)CD8\(^+\) thymocytes.

Conclusion. Our data provide the first genetic evidence for a role for Vav in peptide-MHC-triggered cytoskeletal reorganization in vivo. These results indicate that Vav-regulated actin reorganization is a crucial prerequisite for antigen receptor-mediated selection and apoptosis in peptide-specific thymocytes. We have shown that Vav and the actin cytoskeleton regulate the extent and duration of \(Ca^{2+}\) mobilization after peptide-specific activation, and that Vav functions upstream of mitochondrial pore opening and caspase activation. The defect in peptide-specific apoptosis in vav\(^-/-\) thymocytes can be overcome by activation of PKC, and the inhibition of PKC blocks peptide-specific cell death in wild-type thymocytes, indicating that PKC activation is the trigger for thymocyte apoptosis in response to peptide. Of all PKC isoforms tested, Vav associated only with the PKC-\(\mu\) isoform expressed primarily in T cells, suggesting that PKC-\(\mu\) is the crucial kinase involved in thymocyte selection and clonal deletion.

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Address correspondence to Josef M. Penninger, The Amgen Institute, Ontario Cancer Institute, Department of Medical Biophysics and Immunology, University of Toronto, 620 University Ave., Toronto, Ontario M5G 2C1, Canada. Phone: 416-204-2241; Fax: 416-204-2278; E-mail: jpenning@amgen.com

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