Positive and negative regulation by SLP-76/ADAP and Pyk2 of chemokine-stimulated T-lymphocyte adhesion mediated by integrin α4β1

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ABSTRACT Stimulation by chemokines of integrin α4β1-dependent T-lymphocyte adhesion is a crucial step for lymphocyte trafficking. The adaptor Vav1 is required for chemokine-activated T-cell adhesion mediated by α4β1. Conceivably, proteins associating with Vav1 could potentially modulate this adhesion. Correlating with activation by the chemokine CXCL12 of T-lymphocyte attachment to α4β1 ligands, a transient stimulation in the association of Vav1 with SLP-76, Pyk2, and ADAP was observed. Using T-cells depleted for SLP-76, ADAP, or Pyk2, or expressing Pyk2 kinase–inactive forms, we show that SLP-76 and ADAP stimulate chemokine-activated, α4β1-mediated adhesion, whereas Pyk2 opposes T-cell attachment. While CXCL12-promoted generation of high-affinity α4β1 is independent of SLP-76, ADAP, and Pyk2, the strength of α4β1-VCAM-1 interaction and cell spreading on VCAM-1 are targets of regulation by these three proteins. GTPase assays, expression of activated or dominant-negative Rac1, or combined ADAP and Pyk2 silencing indicated that Rac1 activation by CXCL12 is a common mediator response in SLP-76–, ADAP–, and Pyk2-regulated cell adhesion involving α4β1. Our data strongly suggest that chemokine-stimulated associations between Vav1, SLP-76, and ADAP facilitate Rac1 activation and α4β1-mediated adhesion, whereas Pyk2 opposes this adhesion by limiting Rac1 activation.

INTRODUCTION Trafficking of T-lymphocytes from blood circulation to lymphoid tissue and to sites of injury and infection depends on rapid and transient activation of αLβ2 and α4β1 integrin function by chemokines located on the endothelium and inside tissues (Luster et al., 2005; Ley et al., 2007). Following the binding of chemokines to their receptors, an inside-out signaling that impinges on the cytoplasmic domains of integrins is generated (Thelen and Stein, 2008). The final consequence of this signaling is the generation of high-affinity extra- and intracellular αLβ2- and α4β1-integrin conformations that are competent for binding to their ligands ICAM-1 and VCAM-1, respectively, which are expressed on lymphoid or inflamed endothelium, and also facilitates α4β1 interaction with the CS-1 region of tissue fibronectin.

While talin binding to cytoplasmic motifs on β1- and β2-integrin subunits represents a common and central event for transmission of inside-out signaling from chemokine receptors to generate active α4β1 and αLβ2 (Moser et al., 2009b; Hogg et al., 2011; Kim et al., 2011; Calderwood et al., 2013), important differences exist in the molecular pattern of this signaling to either integrin. αLβ2-dependent leukocyte adhesion stimulated through inside-out signaling shows a crucial requirement for kindlin 3, another important β subunit–binding protein, and to a lesser extent for α4β1 (Malinin et al., 2009; Manevich-Mendelson et al., 2009; Moser et al., 2009a; Svensson et al., 2009; Stein et al., 2010). These data suggest that chemokines can modulate leukocyte adhesion through distinct integrin pathways, potentially influencing the adhesive responses of different immune cell types to chemokines.
et al., 2009). Of note, the RAPL-Rap1 and Rho/Rac-phospholipase D pathways are critical for chemokine-stimulated T-cell adhesion involving αβ2, with no or lesser roles in αβ1-mediated adhesion (Katagiri et al., 2003; Ghandour et al., 2007; Bolomini-Vittori et al., 2009). Instead, the Vav1-Rac1 pathway is required for T-cell adhesion involving αβ1 upon exposure to chemokines (Garcia-Bernal et al., 2005). Vav1 is a multidomains adapter protein that functions as a guanine nucleotide exchange factor (GEF) predominantly for the Rac1 GTPase (Bustelo, 2000; Turner and Billadeau, 2002; Tybulewicz, 2005). In T-cells, chemokine-induced, ZAP-70-dependent Vav1 tyrosine phosphorylation and Vav1 dissociation from talin represent an important event involved in αβ1 activation and in adhesion strengthening mediated by this integrin (Garcia-Bernal et al., 2009).

Vav1 binds key molecules involved in T-lymphocyte activation, such as SLP-76 and Pyk2 (Tuosto et al., 1996; Wu et al., 1996; Katagiri et al., 2000; Gismondini et al., 2003). SLP-76 is an adaptor molecule that is essential for T-cell activation and actin reorganization following T-cell receptor (TCR) stimulation (Koretzky et al., 2006). On TCR activation, ZAP-70-mediated phosphorylation of SLP-76 on N-terminal tyrosine residues allows its binding to the SH2 domain of Vav1 (Tuosto et al., 1996; Wu et al., 1996; Raab et al., 1997), whereas the C-terminal SH2 domain of SLP-76 interacts with several tyrosines on ADAP (da Silva et al., 1997; Musci et al., 1997; Geng et al., 1999; Boerth et al., 2000), a key molecule that also regulates T-cell activation (Griffiths and Penninger, 2002).

Pyk2 is a non receptor tyrosine kinase that is homologous to FAK (Avraham et al., 1995; Lev et al., 1995; Sasaki et al., 1995) and is activated following TCR stimulation and cell exposure to chemotactants (Dikic et al., 1996; Davis et al., 1997; Ganju et al., 1997; Qian et al., 1997). Early data showed that Pyk2 is capable of associating with Vav1 in T and NK cells (Katagiri et al., 2000; Gismondini et al., 2003; Gao and Blaystone, 2009). There are three Pyk2 isoforms of 110 kDa, 105 kDa, and a third species called PRNK (Pyk2-related nonkinase) that only contains 238 amino acids from the C-terminus and lacks the catalytic and N-terminal regions (Xiong et al., 1998; Ivankovic-Dikic et al., 2000). A Tyr-402 residue is a target of Pyk2 autophosphorylation, which in turn recruits and activates Src kinases, leading to Src-dependent phosphorylation of several tyrosine residues on Pyk2 (Dikic et al., 1996; Li et al., 1999; Avraham et al., 2000).

The role of SLP-76, ADAP, and Pyk2 in integrin-mediated T-lymphocyte adhesion has been mostly studied in TCR-activated cells. Thus SLP-76 mediates αβ2-dependent T-cell adhesion in response to TCR stimulation (Baker et al., 2009; Horn et al., 2009). In addition, it has been demonstrated that ADAP is required for TCR-stimulated cell adhesion mediated by β1 and β2 integrins (Griffiths et al., 2001; Peterson et al., 2001), a process regulated by an ADAP/SKAP-55 signaling module (Wang et al., 2003, 2007; Kliche et al., 2006). As for Pyk2, its tyrosine phosphorylation as a consequence of integrin-dependent outside-in signaling was earlier demonstrated (Li et al., 1996; van Severen et al., 1998; Zheng et al., 1998), and it was shown that Pyk2 regulates TCR-stimulated, LFA-1-dependent CD8+ cell adhesion (Beinke et al., 2010).

Little is known on the potential involvement of SLP-76 and Pyk2, as well as of ADAP, in chemokine-stimulated T-cell adhesion dependent on integrins. For instance, SLP-76 is required for optimal T-lymphocyte arrest on ICAM-1 under flow conditions (Lee et al., 2012). In addition, it has been recently shown that CCR7-mediated αβ2 functions in T-cells are regulated by two independent ADAP/SKAP55 modules (Kliche et al., 2012). The integrin αβ1 is especially involved in chemokine-stimulated T-lymphocyte trafficking to sites of inflammation, with lesser roles in cell migration to lymphoid tissues (Berlin-Rufenach et al., 1999), thus highlighting the structural and functional differences between αβ1 and αβ2. The role of SLP-76, ADAP, and Pyk2 in chemokine-activated T-cell adhesion mediated by αβ1 has not yet been addressed. In the present work, we have studied the potential involvement of these signaling molecules in T-lymphocyte adhesion dependent on αβ1.

RESULTS
Role of SLP-76, ADAP, and Pyk2 on chemokine-activated T-cell adhesion mediated by αβ1

Coimmunoprecipitation analyses using anti–SLP-76 monoclonal antibody (mAb) revealed that CXCL12 stimulates a transient increase in SLP-76 association with Vav1 and ADAP (Figure 1A). In agreement with previous work (Fischer et al., 1998), talin was found in anti–SLP-76 immunoprecipitates, and CXCL12 also up-regulated this association. No β1-integrin subunit was found in anti–SLP-76 immunoprecipitates (unpublished data). Cell fractionation experiments showed that SLP-76 is mainly found on the membrane-associated fraction, and its localization was not altered by CXCL12 (Supplemental Figure S1A). Instead, ADAP mostly displayed a cytosolic localization in nonstimulated cells, but it was rapidly and transiently recruited to the membrane following incubation with CXCL12, coincident with the stimulation by this chemokine of SLP-76-ADAP association.

To study potential connections between SLP-76 and ADAP in chemokine-activated T-cell adhesion involving αβ1, we knocked them down using RNA interference in Molt-4 and peripheral blood T-lymphocytes (PBL-T). SLP-76 was depleted with a pool of SLP-76 small interfering RNA (siRNA; see Materials and Methods), whereas ADAP was found to be more efficiently silenced by ADAP-2 than ADAP-1 siRNA (Figure 1B), and we therefore used ADAP-2 siRNA to carry out the subsequent functional assays. Control experiments indicated that SLP-76 or ADAP depletion did not affect cell viability or the expression of CXCR4, α4 or β1 (unpublished data). ADAP knocking down did not alter the coprecipitation between Vav1 and SLP-76 in nonstimulated cells (t = 0), but their increased association in CXCL12-incubated cells was delayed and of smaller magnitude (Figure 1C), suggesting that a critical level of ADAP expression and/or its localization was needed for enhanced Vav1-SLP-76 association.

Previous data showed that the kinase Pyk2 binds to the SH3 domain of Vav1 in Jurkat T-cells (Katagiri et al., 2000). Likewise, Vav1 associated with Pyk2 in Molt-4 cells, and CXCL12 triggered a transient increase in Vav1-Pyk2 binding (Figure 1D, top, and Supplemental Figure S1B), in line with previous findings (Okabe et al., 2005). In addition, low amounts of SLP-76 were also detected in anti-Pyk2 immunoprecipitates. ADAP was minimally detected only in Pyk2 immunoprecipitates from cells not exposed to CXCL12 (unpublished data). Furthermore, we found that Vav1-Pyk2 coimmunoprecipitation was independent of SLP-76 or ADAP, as this association was similarly detected in control, SLP-76 (Supplemental Figure S1C, left), or ADAP siRNA transfectants (unpublished data). Interestingly, antibodies to talin, an additional Vav1 binding partner (Fischer et al., 1998; Garcia-Bernal et al., 2009), coprecipitated Pyk2 in addition to Vav1 in Molt-4 and PBL-T (Figure 1D, bottom). Supporting the specificity of talin-Pyk2 assembly, talin-Pyk2 coprecipitation was diminished in Pyk2 knockout cells (Figure 1, E and F). Whether Pyk2 directly binds to talin, as in FAK-talin interaction (Mitra et al., 2005), or indirectly through its association with Vav1 has not been addressed in the present study. In addition, Pyk2 knocking down did not significantly alter Vav1-talin association or CXCL12-promoted Vav1-talin dissociation (Figure 1G), an event contributing to the initial steps of αβ1-mediated T-cell adhesion (Garcia-Bernal et al., 2009). Moreover, Vav1 and SLP-76 (Supplemental Figure S1C, right)
and SLP-76-talin (unpublished data) interactions were not altered following Pyk2 silencing. We also observed that Pyk2-depleted cells had similar Erk1/2 MAP kinase activation at early time points following stimulation with CXCL12, but this activation was defective at the longest times tested (Supplemental Figure S1D). Instead, CXCL12-promoted Akt stimulation was not altered in Pyk2-knockdown transfectants.

We next tested SLP-76−, ADAP−, and Pyk2-depleted T-cells in α4β1-mediated static cell adhesion assays. As chemokine-up-regulated T-cell attachment is rapid and transient (Grabovsky et al., 2000), the adhesion assays to α4β1 ligands coimmobilized with chemokines were performed using short incubation times (2 min) upon a brief cell centrifugation (15 s) to place cells immediately in contact with ligands. Under these conditions, we observed that SLP-76−-depleted Molt-4 and PBL-T had modest reductions in adhesion to FN−H89 (CS-1/FN), whereas their attachment to VCAM-1 displayed small but nonsignificant decreases compared with control siRNA transfectants (Figure 2A). Similarly, SLP-76−-deficient J14 cells displayed chemokine-promoted adhesion to VCAM-1 comparable with parental Jurkat cells, whereas activated adhesiveness of Lck-deficient JCaM1.6 cells was fully abrogated (Figure 2B), consistent with previous data (Feigelson et al., 2001). ADAP depletion generally affected to a higher degree the stimulation of α4β1-dependent T-cell attachment compared with SLP-76 knocking down (Figure 2A). In addition, ADAP-depleted Molt-4 cells exhibited a substantial inhibition of CXCL12-activated, α4β1-mediated migration across tumor necrosis factor α−treated HUVEC monolayers under static conditions (Supplemental Figure S2), whereas transfectants knocked down for SLP-76 (Supplemental Figure S2) and J14 cells (unpublished data) displayed partial impairments in migration.

Of note, Pyk2 knocking down resulted in significant increases in chemokine-triggered T-cell adhesion to both FN−H89 and VCAM-1 relative to control siRNA transfectants (Figure 2A). Instead, we were unable to detect alterations in attachment to ICAM-1 with CXCL12-incubated, Pyk2-silenced cells (Figure 2C), in line with previous results using Pyk2−/− T-cells exposed to standard doses of anti-CD3 antibodies (Beinke et al., 2010). Moreover, overexpression of PRNK, a Pyk2 C-terminal region isoform that lacks the kinase domain (Xiong et al., 1998) and inhibits the endogenous Pyk2 kinase activity, led to up-regulation of α4β1-dependent attachment of nonstimulated T-cells, which achieved a further enhancement in the presence of CXCL12 (Figure 2D). In support of an involvement of Pyk2 kinase activity in the regulation of α4β1-mediated T-cell adhesion, we found that the expression of the Pyk2-K457A kinase–inactive mutant (Katagiri et al., 2005), we observed similar VCAM-1-Fc binding to SLP-76−, ADAP−, or Pyk2-depleted cells relative to control siRNA transfectants (Figure 3, B and C), a positive control for integrin-affinity regulation. These results indicate that chemokine-stimulated high-affinity α4β1 does not require SLP-76, ADAP, or Pyk2 functions.

To further examine a potential involvement of these three proteins in early steps of CXCL12-stimulated, α4β1-dependent cell adhesion, we used VCAM-1-Fc in soluble binding assays to determine the efficiency of α4β1-VCAM-1 interaction without the subsequent cell spreading step. Using short-incubation conditions to obtain a two- to threefold induction in VCAM-1-Fc binding (20-s stimulation with CXCL12 followed by 75-s incubation with VCAM-1-Fc; Garcia-Bernal et al., 2005), we observed similar VCAM-1-Fc binding to SLP-76−, ADAP−, or Pyk2-depleted cells compared with parental Jurkat cells (unpublished data). When we tested longer incubations with VCAM-1-Fc, we observed that control and ADAP siRNA transfectants (Figure 3D, right) displayed a consistent sustainment of α4β1-VCAM-1-Fc interaction that was inhibited by PTx. All transfectants displayed comparable VCAM-1-Fc binding upon exposure to Mn2+ (Figure 3, B and C), a positive control for integrin-affinity regulation. These results indicate that chemokine-stimulated high-affinity α4β1 avidity, which likely contributes to maintaining α4β1-VCAM-1 interaction.

Role of SLP-76, ADAP, and Pyk2 in adhesion strengthening and spreading mediated by α4β1

To analyze the α4β1-dependent adhesion strength of transfectants depleted for SLP-76, ADAP, or Pyk2, we subjected cells that had been attached to VCAM-1/CXCL12 at 1 dyne/cm2 to detachment at sequential increases of shear stress. ADAP- and SLP-76−-depleted PBL-T and Molt-4 transfectants, as well as J14 cells, were found to develop lower resistance to detachment at high shear stress than control siRNA transfectants, although ADAP silencing consistently led to higher detachment rates than SLP-76 knockdown (Figure 4, A and B). Instead, Pyk2-depleted T-cells exhibited higher resistance to detachment than control siRNA transfectants (Figure 4A).

In the adhesion process, cells undergo spreading after the strengthening of the adhesion. Both SLP-76− and ADAP-knockdown
Molt-4 and PBL-T transfectants developed a time-dependent progressive reduction in the spreading on VCAM-1/CXCL12 compared with control counterparts (Figure 4C, left and middle). On the contrary, Pyk2 knockdown and PRNK transfectants displayed higher spreading than control cells (Figure 4C). These data indicate that ADAP and Pyk2 regulate the chemokine-stimulated T-cell adhesion strengthening and spreading steps mediated by α4β1, whereas SLP-76 mainly contributes to cell spreading, with lesser roles in adhesion strengthening.

**Rac1 involvement in SLP-76, ADAP, and Pyk2 regulation of CXCL12-activated T-cell adhesion dependent on α4β1**

The strength of α4β1-mediated adhesion and the subsequent spreading process in chemokine-stimulated T-cells depends on the activation of the GTPase Rac1 (Garcia-Bernal et al., 2005). Compared with control cells, knocking down SLP-76 led to a less efficient Rac1 activation in Molt-4 cells in response to CXCL12, which was rapidly lost at the longest incubation time tested (Figure 5A, top). Moreover, CXCL12-exposed J14 cells displayed no stimulation of Rac activation relative to parental Jurkat cells (Figure 5A, bottom). In addition, ADAP-depleted transfectants displayed a blockade with or without CXCL12 (n = 3). (D) Cells were transfected with empty (Mock) or PRNK vectors, and transfectants were tested by Western blotting for PRNK expression (left) or in adhesion assays (middle and right) (n = 4). (E) Cells were transfected with control GFP vector or with the indicated GFP-fused Pyk2 mutants, and transfectants were subjected to immunoblotting or to adhesion assays (n = 4). Adhesions were significantly inhibited (**, p < 0.001; *, p < 0.01; n.s., nonsignificant).
of CXCL12-stimulated Rac activation (Figure 5B). Furthermore, inhibition of Rac activation in SLP-76- and ADAP-deficient transfectants was associated with impaired stimulation by CXCL12 of F-actin polymerization (Supplemental Figure S3A). In contrast, Pyk2 silencing led to a more sustained Rac1 activation and F-actin polymerization than control transfectants (Figure 5C and Supplemental Figure S3B).

Chemokine-stimulated Rac1 activation in T-cells was previously shown to involve Vav1 tyrosine phosphorylation (Garcia-Bernal et al., 2002). Furthermore, the CXCL12-triggered enhancement of Rac activation in SLP-76-silenced cells was detected in ADAP-depleted cells (Figure 5D), similar to the unaltered Vav1 phosphorylation in ADAP knockout mice (Reynolds et al., 2002). Furthermore, the CXCL12-triggered enhancement of Rac activation seen in Pyk2-silenced transfectants did not correlate with higher Vav1 tyrosine phosphorylation than control counterparts, and this phosphorylation generally displayed lower intensity compared with control transfectants (Figure 5D). Similar to Molt-4 cells, SLP-76 knockdown in PBL-T resulted in inefficient Vav1 tyrosine phosphorylation in response to CXCL12, whereas phosphorylation was not affected in ADAP-depleted cells (Figure 5E). Again, Pyk2-silenced PBL-T exhibited CXCL12-stimulated Vav1 tyrosine phosphorylation but of lower magnitude and less maintained than control siRNA transfectants (Figure 5E).

Next we tested the relevance of the deficient Rac activation shown by SLP-76- and ADAP-deficient T-cells on their impaired adhesion. For this, we transfected green fluorescent protein (GFP)-fused, constitutively active forms of Rac (GFP-Rac V12) in J14 and parental Jurkat cells or expressed GFP-Rac V12 together with ADAP siRNA, and then we tested the transfectants in adhesion assays to VCAM-1 coimmobilized with CXCL12. Flow-chamber adhesion assays revealed that the reduction in stable adhesion shown by J14-GFP cells relative to parental Jurkat-GFP was significantly rescued upon expression of GFP-Rac V12 (Figure 6A). Likewise, GFP-Rac V12 expression reversed the inhibitory effects of ADAP siRNA transfection on the stimulation of cell adhesion to VCAM-1/CXCL12 (Figure 6B). Moreover, the up-regulated adhesion to VCAM-1/CXCL12 shown by Pyk2-knockdown Molt-4 and PBL-T transfectants was partially reversed when dominant-negative, GFP-fused Rac1 (GFP-Rac N17) was transfected together with Pyk2 siRNA (Figure 6C and D). These results strongly suggest that altered Rac1 activation in SLP-76- and ADAP-, or Pyk2-knockdown T-cells represents an important mechanism mediating their changes in chemokine-stimulated, α4β1-dependent cell adhesion.

As the adhesion data indicate that T-cell attachment involving α4β1 is especially altered upon knocking down ADAP or Pyk2, we next analyzed the effect of their combined depletion on cell adhesion to VCAM-1. Results revealed that the inefficient stimulation of adhesion displayed by ADAP-silenced cells could be rescued by Pyk2 siRNA cotransfection, although the double-knockdown transfectants did not reach the adhesion values exhibited by cells transfected with Pyk2 siRNA alone (Figure 6A). Moreover, the up-regulated cell attachment following Pyk2 silencing was not detected when both SLP-76 and Pyk2 were knocked down (Figure 6B), suggesting that cooperative responses between ADAP and SLP-76 exist during their positive regulation of α4β1-dependent cell adhesion. Similar to ADAP-Pyk2 siRNA cotransfection, PRNK transfection together with ADAP siRNA led to a recovery of adhesion to VCAM-1/CXCL12 (Figure 6C). Notably, the rescue in adhesion shown by cells knocked down for both ADAP and Pyk2 correlated with longer, sustained Rac1 activation in these transfectants (Figure 7D). The small decrease in activation of Rac at early time points might reflect initial inhibition due to ADAP depletion that is later recovered by siPyk2 silencing. These results suggest that the stimulatory actions of ADAP and the inhibitory activity of Pyk2 during CXCL12-activated adhesion mediated by α4β1 are connected through Rac1.

**FIGURE 3:** Flow-chamber adhesion assays and determination of high-affinity α4β1 expression in SLP-76-, ADAP-, or Pyk2-depleted T-cells. (A) Control, SLP-76, ADAP, or Pyk2 siRNA transfectants or Jurkat or J14 cells were perfused in flow chambers coated with VCAM-1 and analyzed for rolling and stable cell arrest (n = 3–4). Data are presented as mean ± SD of cell percentages from the total cell population. Adhesions were significantly inhibited or stimulated in comparison with those of control siRNA transfectants or parental Jurkat cells, *p < 0.05 or **p < 0.005, respectively. (B and C) The indicated siRNA Molt-4 transfectants or cells transfected with PRNK or empty vector were tested by flow cytometry for HUTS-21 mAb binding after stimulation with CXCL12 or Mn^2+. (D) Following exposure to CXCL12 for 20 s, transfectants were analyzed by flow cytometry for VCAM-1-Fc binding after the indicated times. PTx denotes cells preincubated with pertussis toxin.

|        | Rolling | Stable adhesion |
|--------|---------|-----------------|
| Control | 80      | 20              |
| SLP-76   | 50      | 50              |
| ADAP    | 30      | 70              |
| Pyk2    | 20      | 80              |

|        | Fluorescence Intensity (fold induction) |
|--------|----------------------------------------|
| Medium |                                        |
| CXCL12 |                                        |
| Mn^2+  |                                        |

|        | Fluorescence Intensity |
|--------|------------------------|
| Mock   | 20                     |
| PRNK   | 40                     |

|        | Fluorescence Intensity (fold induction) |
|--------|----------------------------------------|
| Medium |                                        |
| CXCL12 |                                        |
| Mn^2+  |                                        |

|        | Fluorescence Intensity |
|--------|------------------------|
| 0      | 20                     |
| 75     | 40                     |
| 120    | 60                     |
| 180    | 80                     |
| 240    | 100                    |

\[ n = 4 \]
proteins with Vav1, strongly suggesting that the dynamics of this protein module can regulate the extent of α4β1-mediated T-cell attachment.

Alterations in adhesion to α4β1 ligands exhibited by SLP-76–, ADAP–, or Pyk2-depleted cells were independent of changes in α4β1 affinity following CXCL12 stimulation. Rather, flow-chamber adhesion assays revealed that the strengthening of α4β1-dependent attachment was regulated by SLP-76 and ADAP, though the strength of ADAP-knockdown transfectant attachment was consistently lower than SLP-76–depleted counterparts. Moreover, we show that spreading on VCAM-1/CXCL12 was similarly decreased in T-cells silenced for SLP-76 and ADAP. Compared with the absolute requirement of talin and Vav1 in the control of α4β1 affinity and strength of

FIGURE 4: Analyses of T-cell adhesion strengthening and spreading on VCAM-1. The indicated siRNA transfectants (A) or Jurkat and J14 cells (B), preattached onto coimmobilized VCAM-1 and CXCL12 in flow chambers, were subjected to cell detachment after sequential increases of shear stress. Data show mean ± SD of cell percentages from the initial number of bound cells remaining attached at the indicated shear stresses (A, n = 4; B, n = 3). (C) Molt-4 and PBL-T were transfected with the indicated siRNA or vectors, and transfectants were allowed to attach to VCAM-1 immobilized with CXCL12. Spreading was evaluated from Nomarski images at the indicated times. Percentage of cell spreading was determined from cells (2500–3000) from different fields of view. Spreading was significantly inhibited (**, p < 0.01; *, p < 0.05) or stimulated (Δ, p < 0.05) compared with control siRNA or mock transfectants (n = 3). n.s., nonsignificant.

DISCUSSION

Vav1 is needed for chemokine-stimulated T-cell adhesion involving α4β1 (Garcia-Bernal et al., 2005, 2009), and thus it is conceivable that Vav1-binding proteins such as SLP-76 and Pyk2 could modulate this process. Using Molt-4 and primary human PBL-T depleted for SLP-76 or for ADAP, an SLP-76–interacting partner, as well as SLP-76–deficient J14 cells, we show here that these two proteins contribute to optimal stimulation by CXCL12 of α4β1-mediated T-cell attachment under flow conditions. Instead, Pyk2 negatively regulates this adhesion, as its depletion or expression of kinase-inactive Pyk2 forms led to increased cell attachment. Of note, modulation of cell adhesion by SLP-76, ADAP, and Pyk2 correlated with a chemokine-stimulated transient increase in association of these proteins with Vav1, strongly suggesting that the dynamics of this protein module can regulate the extent of α4β1-mediated T-cell attachment.

Alterations in adhesion to α4β1 ligands exhibited by SLP-76–, ADAP–, or Pyk2-depleted cells were independent of changes in α4β1 affinity following CXCL12 stimulation. Rather, flow-chamber adhesion assays revealed that the strengthening of α4β1-dependent attachment was regulated by SLP-76 and ADAP, though the strength of ADAP-knockdown transfectant attachment was consistently lower than SLP-76–depleted counterparts. Moreover, we show that spreading on VCAM-1/CXCL12 was similarly decreased in T-cells silenced for SLP-76 and ADAP. Compared with the absolute requirement of talin and Vav1 in the control of α4β1 affinity and strength of
**FIGURE 5:** Analyses of Rac1 activation and Vav1 tyrosine phosphorylation in SLP-76–, ADAP–, or Pyk2-deficient T-cells. Control, SLP-76, ADAP, or Pyk2 siRNA transfectants in Molt-4 cells, or Jurkat and J14 cells (A–C) were stimulated with CXCL12 and subjected to GTPase assays to detect Rac1 activation. Activation was significantly increased compared with untreated cells ($^{\Delta \Delta \Delta}$, $p < 0.001$; $^{\Delta \Delta}$, $p < 0.01$; $^{\Delta}$, $p < 0.05$; A, $n = 7$; B and C, $n = 3$). The indicated Molt-4 (D) or PBL-T (E) siRNA transfectants incubated with or without CXCL12 for the indicated times were analyzed by immunoprecipitation and Western blotting. Vav1 tyrosine phosphorylation was significantly up-regulated compared with untreated cells ($^{\Delta \Delta}$, $p < 0.01$; $^{\Delta}$, $p < 0.05$; D, $n = 6$; E, left, $n = 2$; E, right, $n = 3$).
FIGURE 6: Rac1 involvement in SLP-76, ADAP, and Pyk2 regulation of CXCL12-activated T-cell adhesion dependent on α4β1. (A) Cells were transfected with GFP or GFP-Rac V12 vectors and were analyzed by immunoblotting (top) or were subjected to flow-chamber adhesion assays to VCAM-1 coimmobilized with CXCL12 (bottom). Adhesion was significantly rescued in comparison with J14-GFP transfectants (Δ, p < 0.05; n = 3). (B) Molt-4 cells were transfected with the indicated siRNA and GFP vector combinations and were analyzed by Western blotting (top) or in static adhesion assays to VCAM-1 immobilized with or without CXCL12 (bottom). Adhesion was significantly inhibited (*, p < 0.05) or recovered (Δ, p < 0.05). (C and D) The indicated Molt-4 or PBL-T transfectants were tested by immunoblotting or were subjected to adhesion assays to VCAM-1 immobilized with or without CXCL12. Adhesion was significantly stimulated (ΔΔ, p < 0.01) or inhibited (*, p < 0.05) (B–D, n = 3).
chemokine-activated T-cell attachment (Garcia-Bernal et al., 2006, 2009; Manevich et al., 2007), the contribution of SLP-76 and ADAP in these processes appears to be of lesser magnitude. However, both adaptors are clearly needed to achieve optimal stability of this adhesion.

VCAM-1-Fc soluble binding to Pyk2-silenced cells was maintained longer than with control transfectants, suggesting that Pyk2 can regulate the avidity of α4β1. Sustained VCAM-1 binding likely represents one of the mechanisms accounting for the stronger cell arrest and resistance to detachment under shear stress shown by Pyk2-knockdown cells relative to control counterparts. Associated with increased strength of attachment to VCAM-1/CXCL12 exhibited by Pyk2-depleted cells, they displayed enhanced spreading, an observation in line with previous results showing high spreading correlating with defective cell detachment of Pyk2−/− macrophages (Okigaki et al., 2003). Therefore, our data indicate that Pyk2 negatively regulates the strength and spreading phases of chemokine-activated T-cell attachment involving α4β1.

Our previous results revealed that Vav1-mediated Rac1 activation in response to CXCL12 in T-cells represents an important event for strengthening of α4β1-VCAM-1 interaction, as well as for cell spreading (Garcia-Bernal et al., 2005). The present data indicate that Rac1 is a common mediator of SLP-76, ADAP, and Pyk2-modulated T-cell adhesion to α4β1 ligands. Thus SLP-76 and ADAP likely function as adaptors with Vav1 to optimize chemokine-promoted Rac activation and stimulation of α4β1-dependent cell adhesion. Instead, Pyk2 restricts this attachment by limiting Rac activation. As knocking down any of these three proteins influences the outcome of Rac activation, the data strongly suggest that depletion-mediated changes in the composition of the complex formed by Vav1, SLP-76, ADAP, and Pyk2 could account for alterations in Rac activation. In this regard, we show that knocking down ADAP delays Vav1-SLP-76 association and impairs Rac activation in response to CXCL12, suggesting that the Vav1-SLP-76-ADAP module facilitates Rac1 activation upon chemokine stimulation and optimizes the strength and spreading of α4β1-mediated T-cell adhesion. On this point, SLP-76 assembly with Vav1 and Nck contributes to TCR-stimulated Rac1 activation (Bubeck Wardenburg et al., 1998; Barda-Saad et al., 2010), and importantly, ADAP associates with these complexes together with WASP (Krause et al., 2000; Coppolino et al., 2001) and cooperates with Nck in the recruitment of SLP-76 for regulation of Rho GTPase–mediated actin rearrangement (Pauker et al., 2011).

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Earlier results reported that SLP-76 depletion in Jurkat cells did not affect the CXCL12-stimulated Rac1 activation (Horn et al., 2009). The basis for the discrepancies between these results and our observation of inefficient Rac1 activation by CXCL12 in SLP-76–deficient cells is not known at present.

FIGURE 7: Analysis of functional connections between ADAP and Pyk2 in CXCL12-stimulated adhesion mediated by α4β1. (A–C) Molt-4 cells were transfected with the indicated individual or combined siRNA and/or vectors, and transfectants were analyzed by immunoblotting or in adhesion assays to VCAM-1 immobilized with or without CXCL12. Adhesions were significantly inhibited (*p < 0.05) or significantly stimulated or rescued (ΔΔΔ, p < 0.001; ΔΔ, p < 0.01; or Δ, p < 0.05) (A, n = 4; B and C, n = 3). (D) The indicated transfectants were tested in GTPase assays to detect active Rac.
Characterization of the mechanics of Pyk2-regulated, α4β1-mediated cell attachment reveals that the Pyk2 kinase domain negatively controls this adhesion. Thus expression of PRNK, which lacks the Pyk2 kinase domain, as well as of a point mutant, kinase-inactive Pyk2 form, mimicked the effects of Pyk2 depletion on the increase in α4β1-mediated activation, strongly suggesting that Pyk2-dependent tyrosine phosphorylation activity in T-cells contributes to keep Rac1 activation levels under control. Of note, negative regulation of Rac activation and cell adhesion to VCAM-1 by Pyk2 directly opposed the stimulatory activity of ADAP in these processes, as detected in cells silenced for both proteins. In this experimental setting, the absence of Pyk2 might rescue the activation of Rac1 concurrent with the inhibitory effects of ADAP knockdown, leading to stimulation of adhesion to VCAM-1, but we cannot exclude that recovery of these activities occurs subsequent to the negative actions of ADAP depletion. Our data are in line with early work proposing that Pyk2 regulates Rac1 activation (Tokiwa et al., 1996) and with results showing that macrophages from mice knocked out for the Pyk2 closely related kinase FAK have elevated Rac activation (Owen et al., 2007). However, the level of activation of this GTPase based on Pyk2 activity might be cell-context dependent (Chapman and Houtman, 2014), as NK cells expressing a kinase-dead Pyk2 displayed reduced Rac activation (Gismondi et al., 2003).

CXCL12 promotes tyrosine phosphorylation of Vav1 associated with Rac1 activation and stimulation of α4β1-mediated T-cell adhesion (Garcia-Bernal et al., 2005). Inefficient phosphorylation of Vav1 in response to CXCL12 in SLP-76-depleted T-cells correlated with their impaired Rac activation, whereas ADAP-knockdown cells displayed fully competent chemokine-promoted phosphorylation of this GEF, in spite of their defective Rac activation. These results might reflect the fact that ADAP does not directly interact with Vav1 but associates with the Vav1-SLP-76 module, possibly in a SLP-76-dependent manner. Interestingly, the increased Rac activation following CXCL12 stimulation in Pyk2-silenced cells was not coupled with enhanced Vav1 phosphorylation, but rather phosphorylation was not as efficient as in control transfectants. These data reveal that SLP-76 and Pyk2 but not ADAP contribute to optimal Vav1 tyrosine phosphorylation in T-cells in response to CXCL12, and indicate that net Rac activation is the result of Vav1-dependent and Vav1-independent mechanisms. It remains puzzling that inefficient Vav1 phosphorylation in SLP-76- and Pyk2-depleted T-cells led to opposite effects on Rac activation. The molecular basis underlying these distinct responses is not known at present. SLP-76 and Pyk2 knockdown might similarly affect recruitment and/or activity of tyrosine kinases involved in Vav1 tyrosine phosphorylation, such as ZAP-70 and Lck (Guibins et al., 1993; Ottoson et al., 2001). But it is also clear that the subsequent Rac activation is differentially regulated and likely involves distinct GEFs and GTPase-activating proteins (GAPs). GEFs and GAPs that are known to regulate CXCL12-stimulated Rac activation and α4β1-mediated T-cell adhesion include the GEF DOCK2 and the GAP β-chimerin. Of note, lymphocytes from DOCK2 knockout mice display minimal chemokine-induced Rac activation compared with control cells (Fukui et al., 2001). Furthermore, we showed that DOCK2 cooperates with Vav1 for CXCL12-promoted Rac activation, and its silencing inhibits T-cell attachment mediated by α4β1 under flow conditions (Garcia-Bernal et al., 2006). Instead, β-chimerin overexpression in T-cells results in impaired Rac activation in response to CXCL12 associated with inhibition of α4β1-dependent T-cell adhesion (Siliceo et al., 2006). Although it has not yet been tested, the possibility exists that DOCK2 and β-chimerin might be recruited to the Vav1-Pyk2-SLP-76-ADAP module, thus contributing to the regulation of Rac activity and subsequent cell attachment mediated by α4β1. Other GEFs and GAPs expressed on T-lymphocytes that are known to regulate Rac activation include PlekH2, Pix, and ArhGAP15 (Seoh et al., 2003; Phee et al., 2005; Runne and Chen, 2013), though their role in the control of α4β1-dependent T-cell adhesion has not been addressed.

SKAP-55 is an additional molecule that could modulate ADAP-controlled T-cell attachment to α4β1 ligands following chemokine stimulation. Thus ADAP binds SKAP-55, an adaptor protein that mediates up-regulation of T-cell adhesion to ICAM-1 and fibronectin upon TCR activation (Marie-Cardine et al., 1998; Wang et al., 2003, 2007). The ADAP-SKAP-55 module interacts with RIAM through SKAP-55, facilitating TCR-mediated integrin activation (Kliche et al., 2006; Menasche et al., 2007; Horn et al., 2009). Furthermore, independent ADAP-SKAP-55 modules regulate CCR7-mediated αLβ2 functions in T-cells (Kliche et al., 2012), and recent data indicate that ADAP can also bind talin and kindlin to promote integrin-αIIβ3 activation and fibrinogen binding (Kasirer-Friede et al., 2014). Whether the ADAP-SKAP-55 association contributes to chemokine-stimulated Rac activation and T-cell adhesion involving α4β1 is currently unknown.

A model can be proposed from the present results (Figure 8). The net Rac1 activation levels needed to develop sufficient adhesion strength and cell spreading mediated by α4β1 are possibly the result of opposing activities inside the Vav1-SLP-76-ADAP-Pyk2 module. Thus Pyk2 kinase activity would oppose the Rac1 activation

![Figure 8: Model for regulation by Vav1-associated proteins of chemokine-stimulated, α4β1-dependent T-cell adhesion. Rac1 activation by CXCL12 is a well-known event for stimulation of the adhesion strength and spreading mediated by α4β1 in T-cells. A transient complex formed by Vav1, SLP-76, ADAP, and Pyk2 regulates this adhesion. While Vav1-SLP-76-ADAP stimulates chemokine-promoted Rac1 activation (green arrow), the presence of Pyk2 in the complex opposes this activation (red blocking symbol). The result of these opposing actions is a net Rac activation and up-regulation of T-cell adhesion strengthening and spreading dependent on α4β1. Talin can also be found in this complex, perhaps as an independent pool from β1-associated talin. Its role in the assembly and function of this complex has not been addressed in the present study.](image-url)
stimulated by Vav1-SLP-76-ADAP and possibly additional costimulatory molecules, potentially by activating a GAP or by inhibiting a GEF for Rac1. Talin can also be found in this protein module, but its role in the stability and function of the complex and whether it represents a different pool from β1-associated talin has not been addressed here. Furthermore, the present data, together with our previous results (Garcia-Bernal et al., 2009), raise the possibility that separate pools of SLP-76 with either Vav1 or talin might exist. Our data should add molecular mechanistic insights into the process of chemokine-stimulated T-cell adhesion mediated by α4β1, a key step during T-lymphocyte trafficking to sites of inflammation.

MATERIALS AND METHODS

Cells, antibodies, and reagents

The human Molt-4 and Jurkat T-cell lines were cultured in RPMI 1640 medium (Lonza, Verviers, Belgium) and 10% fetal bovine serum (Life Technologies, Paisley, UK) (complete medium). The Jurkat-derived cell lines JCaM1.6 (American Type Culture Collection) and J14 (a gift from Arthur Weiss, University of California, San Francisco, CA) were grown in complete medium. Peripheral blood T-lymphocytes (PBL-T) were prepared as previously described (Garcia-Bernal et al., 2005). The Consejo Superior de Investigaciones Científicas Ethics Committee (Madrid, Spain) approved the protocols used to obtain and process the human blood samples. Control P3×63 and anti-β1 TS2/16 mAb were gifts from Francisco Sánchez-Madrid (Hospital de la Princesa, Madrid, Spain), and polyclonal anti-β1A antibodies were from Guido Taron (Turin University, Italy). Antibodies to SLP-76 and Vav1 (C14) were from Santa Cruz Biotechnology (Santa Cruz, CA), anti-phospho-Vav (Y174; ab47282) was from Abcam (Cambridge, MA), anti-Rac1 and anti-ADAP were from BD Biosciences (San Diego, CA), and anti-GFP was from Molecular Probes (Eugene, OR). Antibodies to β-actin, vinculin, and talin clone 8D4 were from Sigma-Aldrich (St. Louis, MO), and antibody to Pyk2 was from Millipore (Billerica, MA). The HUTS-21 anti-β1 mAb has been previously characterized (Luque et al., 1996). CXCL12 was purchased from R&D Systems (Minneapolis, MN) and CCL21 was purchased from Peprotech (Rocky Hill, NJ).

Transfections and RNA interference

Vectors coding for Rac wild-type (wt), Rac V12, Rac N17, Pyk2-Y402F, Pyk2-K457A, and Pyk2-wt fused with GFP were provided by Francisco Sánchez-Madrid. The vector pRKS5-myc-PRNK (Ivanovic-Dikic et al., 2000) was a gift from Ivan Dikic (Goethe University School of Medicine, Frankfurt, Germany). The siRNA against human SLP-76 (ON-TARGETplus, SMARTpool) was purchased from Dharmacon (Lafayette, CO). The siRNAs for ADAP-1, sense: CAAUUGGUUGACCUAGCAAtt; ADAP-2, sense: GAGAUAUUAAGGUGUCCUtt; Pyk2, sense: GAUGUGGUCGUUUAUGUAtt; and control siRNA (Garcia-Bernal et al., 2005) were purchased from Sigma-Aldrich. Vectors and siRNA were nucleofected (Amaxa,ologne, Germany) following a previously described procedure (Garcia-Bernal et al., 2005). Molt-4 or PBL-T siRNA transfecteds were transfected 22–24 h posttransfection, and transfection did not affect cell viability, as assessed in cell cycle analyses by flow cytometry.

Cell adhesion and spreading assays

For static adhesions, we used the reported method (Garcia-Bernal et al., 2005). Briefly, cells were plated in triplicate on wells coated with the fibronectin fragment FN-H89, which contains the CS-1 region, or with VCAM-1 together with CXCL12 or CCL21 (650 ng/ml), and plates were incubated and adhesion measured. Adhesion data are presented relative to control untreated cells, which have been given an arbitrary value of 100. For flow-chamber adhesion assays, we followed the described protocol (Garcia-Bernal et al., 2006). In brief, cells were infused at a flow rate of 1 dyne/cm² into flow chambers containing coimmobilized VCAM-1 and CXCL12. Rolling cells that subsequently firmly attached were expressed as stable arrest, whereas tethering cells that did not arrest at any moment were expressed as rolling cells. For evaluation of shear resistance, cells were allowed to attach and then were subjected to sequential increases of the flow rate. The number of cells remaining bound was determined as the percentage of total adhered cells after the adhesion step. For cellular spreading, CXCL12-stimulated cells attached on VCAM-1 were fixed and analyzed as previously described (Garcia-Bernal et al., 2006).

Soluble VCAM-1 binding and flow cytometry

For cell binding to VCAM-1-Fc (R&D Systems), cells were stimulated for 20 s with CXCL12 before VCAM-1-Fc was added, which was detected by flow cytometry using phycoerythrin-conjugated AffiniPure F(ab′)2 fragment goat anti-human immunoglobulin G (IgG), Fcy-fragment specific (Jackson ImmunoResearch, West Grove, PA). For the detection of high-affinity β1, cells were stimulated for 20 s with CXCL12 before HUTS-21 anti-β1 mAb (10 μg/ml) was added for 30 min at 4°C. After washing, cells were incubated with FITC-conjugated rabbit anti-mouse IgG (Jackson ImmunoResearch). Fluorescence intensity data indicate fold-induction values relative to those from control untreated cells, which were given an arbitrary value of one.

Immunoprecipitation, immunoblotting, and GTPase assays

For immunoprecipitation, we followed the described method (Garcia-Bernal et al., 2009). In brief, cell lysate supernatants were incubated with antibodies followed by coupling to protein G-Sepharose. Proteins were separated by SDS–PAGE and transferred to membranes that were sequentially incubated with primary antibodies and with horseradish peroxidase–conjugated secondary antibodies. Protein visualization was achieved using Immobilon Western chemiluminescent substrate (Millipore). For GTPase assays, we used the reported method (Garcia-Bernal et al., 2005). Briefly, cells exposed to CXCL12 were lysed, and aliquots from extracts were separated for total lysate controls and for incubation with GST-PAK-CD fusion protein, followed by incubation with glutathione-Sepharose beads (GE Healthcare, Piscataway, NJ). Bound proteins were eluted and subjected to immunoblotting using anti-Rac1 antibodies.

Statistical analyses

Data were analyzed by one-way analysis of variance, followed by Tukey-Kramer multiple comparisons. In both analyses, the minimum acceptable level of significance was p < 0.05.

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