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T cell specific adaptor protein (TSAd) promotes interaction of Nck with Lck and SLP-76 in T cells

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Abstract

Background: The Lck and Src binding adaptor protein TSAd (T cell specific adaptor) regulates actin polymerization in T cells and endothelial cells. The molecular details as to how TSAd regulates this process remain to be elucidated.

Results: To identify novel interaction partners for TSAd, we used a scoring matrix-assisted ligand algorithm (SMALI), and found that the Src homology 2 (SH2) domain of the actin regulator Non-catalytic region of tyrosine kinase adaptor protein (Nck) potentially binds to TSAd phosphorylated on Tyr²⁸⁰ (pTyr²⁸⁰) and pTyr³⁰⁵. These predictions were confirmed by peptide array analysis, showing direct binding of recombinant Nck SH2 to both pTyr²⁸⁰ and pTyr³⁰⁵ on TSAd. In addition, the SH3 domains of Nck interacted with the proline rich region (PRR) of TSAd. Pull-down and immunoprecipitation experiments further confirmed the Nck-TSAd interactions through Nck SH2 and SH3 domains. In line with this Nck and TSAd co-localized in Jurkat cells as assessed by confocal microscopy and imaging flow cytometry. Co-immunoprecipitation experiments in Jurkat TAG cells lacking TSAd revealed that TSAd promotes interaction of Nck with Lck and SLP-76, but not Vav1. TSAd expressing Jurkat cells contained more polymerized actin, an effect dependent on TSAd exon 7, which includes interactions sites for both Nck and Lck.

Conclusions: TSAd binds to and co-localizes with Nck. Expression of TSAd increases both Nck-Lck and Nck-SLP-76 interaction in T cells. Recruitment of Lck and SLP-76 to Nck by TSAd could be one mechanism by which TSAd promotes actin polymerization in activated T cells.

Keywords: TSAd, T cell specific adaptor protein, Nck, Lck, SH2 domain, adaptor protein, SH2D2A, SLP-76

Background

Regulation of actin dynamics is important for several aspects of T cell function, including differentiation, migration through tissues and proliferation. T cell activation initiates multiple molecular events including activation of protein tyrosine kinases, the formation of protein signaling complexes, and cytoskeletal actin reorganization leading to establishment of the immunological synapse (IS) at the T cell-antigen presenting cell (APC) interface [1, 2]. Briefly, following T cell receptor (TCR) ligation the

tyrosine kinases Lck and Zap-70 become activated, leading to the formation of a signaling complex containing the adaptor proteins linker for activation of T cells (LAT), SH2 domain containing leukocyte protein of 76 kDa (SLP-76) and non-catalytic region of tyrosine kinase adaptor protein (Nck) [2].

The molecular aspects concerning actin dynamics in T cells are complex, and many questions remain to be solved. One of the more recently identified players of actin reorganization is the T cell specific adaptor protein (TSAd) [3] encoded by the *SH2D2A* gene. TSAd interacts with and modulates the activity of the Src family protein tyrosine kinase Lck [4, 5] as well as Src itself [6]. TSAd has been found to control actin polymerization events in T cells and endothelial cells. More specifically,

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in response to VEGF-A stimulation, TSA_d is required for stress fiber formation and migration of endothelial cells [7]. Moreover, we have also shown that TSA_d regulates CXCL12-induced migration and actin cytoskeletal rearrangements in T cells by promoting Lck dependent tyrosine phosphorylation of IL2-inducible T-cell kinase (Itk) [8].

To better understand the function of TSA_d, we used an algorithm for identification of SH2 domain-ligand pairs (SMALI) to identify possible binding partners for the TSA_d phosphotyrosines. SMALI pointed to a possible interaction between TSA_d and the adaptor Nck. Nck is known to regulate the actin cytoskeleton. It consists of one C-terminal Src homology 2 (SH2) domain and three N-terminal SH3 domains which allows for multiple protein-protein interactions. More than 60 interaction partners for Nck have been identified [9, 10]. Nck interacts constitutively with the guanine nucleotide exchange factor Vav1 [11]. Upon TCR-triggering, Nck and Vav1 interacts with SLP-76, leading to the activation of the actin rearrangement at the T-cell APC interface. Thus, Nck is a key adaptor in T cell activation-dependent actin filament formation through its interactions with components of the TCR/CD3 complex and cytoskeletal regulators including Vav1 and SLP-76 [9, 12–14]. Nck plays a universal role in regulation of the signaling networks critical for organizing the actin cytoskeleton; including formation of the IS following TCR engagement, cell proliferation and cell migration [9, 15, 16].

Here we explored the possible interaction between TSA_d and Nck using intact and mutated TSA_d and Nck constructs. We found that the Nck SH2 domain binds to both TSA_d pTyr²⁸⁰ and TSA_d pTyr³⁰⁵, with pTyr²⁸⁰ as the preferred binding site. Additionally, two of the three Nck SH3 domains were found to interact with the PRR on TSA_d, presumably in a cooperative manner. Our data indicate the existence of a direct interaction between of Nck and TSA_d. When TSA_d is co-expressed, interaction of Nck with Lck is increased. Moreover, TSA_d also enables Nck to interact with SLP-76, an interaction previously shown to be important for actin polymerization and rearrangement [17]. TSA_d promoted actin polymerization in Jurkat cells, and this was dependent on TSA_d exon 7 encoding interaction sites for both Nck and Lck. Thus, the Nck-TSA_d interaction may represent an additional link whereby TSA_d contributes to the regulation of the actin cytoskeleton in T cells.

Results

The Nck SH2 domain interacts with TSA_d-pTyr²⁸⁰ and -pTyr³⁰⁵

TSA_d possesses several protein interaction motifs, including an N-terminally located SH2 domain, and a C-terminal part consisting of a PRR and several tyrosine

phosphorylation sites. TSA_d is tyrosine phosphorylated in non-stimulated Jurkat cells [4, 18] and in peripheral blood mononuclear cells [3] while increased amount of tyrosine phosphorylated TSA_d may be seen upon TCR stimulation [18]. To identify novel SH2 domain containing binding partners for TSA_d, we performed an *in silico* scan using the SMALI algorithm [19, 20]. A relative SMALI score >1.0, strongly indicates potential binding between an SH2 domain and a phosphotyrosine containing ligand. SMALI identified the Nck SH2 domain as a possible interaction partner for TSA_d pTyr²⁶⁰, pTyr²⁸⁰ and pTyr³⁰⁵, (relative SMALI scores: 1.05, 1.37 and 1.10 respectively) (Fig. 1a). In comparison, the Lck-SH2 domain already known to bind to the three C-terminal TSA_d phosphotyrosines [5], displayed a SMALI score of 1.21 for TSA_d-pTyr³⁰⁵. To test the SMALI predictions, we carried out a peptide array analysis of 15-mer TSA_d phosphotyrosine peptides spotted to a nitrocellulose membrane. Recombinant GST-Nck SH2 protein showed direct binding to both TSA_d pTyr²⁸⁰ and pTyr³⁰⁵ while the TSA_d pTyr²⁶⁰ phosphopeptide only revealed a weak signal (Fig. 1b). In accordance with the SMALI predictions, the pTyr²⁸⁰ interaction displayed the strongest signal (Fig. 1b).

Interaction between Nck and TSA_d was then assessed by *in vitro* pull-down interaction analysis in lysates from 293T cells transiently expressing TSA_d as well as the kinase Lck to ensure phosphorylation of TSA_d [4, 18]. The constructs used for transfections are depicted in Fig. 1c and d. TSA_d exists in different phosphorylation forms, with different migration in SDS-PAGE [5]. In the absence of Lck, the lower band predominates (Fig. 1e, lane 1). Nck SH2 protein pulled down TSA_d or TSA_d lacking its PRR from 293T cells only when co-expressed with Lck (Fig. 1e, lane 3 and 6). In contrast TSA_d lacking the three penultimate tyrosines known to be phosphorylated by Lck [5], failed to interact (Fig. 1e, lane 4). Moreover, Nck SH2 did not interact with Lck in this experiment (data not shown). We previously showed that removal of the TSA_d PRR is associated with reduced phosphorylation of TSA_d [5]. In line with this observation, the absence of TSA_d PRR results in reduced interaction between the GST-Nck SH2 domain and TSA_d (Fig. 1e, lane 6) compared to TSA_d WT.

Further analysis revealed that substantially less of the TSA_d Tyr²⁸⁰ → Phe²⁸⁰ (Y280F) mutant was pulled down with Nck SH2, while Nck SH2 could still pull down the TSA_d Y305F protein (Fig. 1f). Elimination of Tyr²⁹⁰ did not affect TSA_d's binding to Nck SH2 in 293T cells (Fig. 1f), as predicted by SMALI and shown in the peptide array analysis (Fig. 1b). GST alone did not bind to TSA_d (data not shown). The results presented in Fig. 1f revealed that single mutations of TSA_d Tyr²⁸⁰, Tyr²⁹⁰ and Tyr³⁰⁵ did not completely abolish binding between TSA_d and the Nck SH2 domain. We thus proceeded to

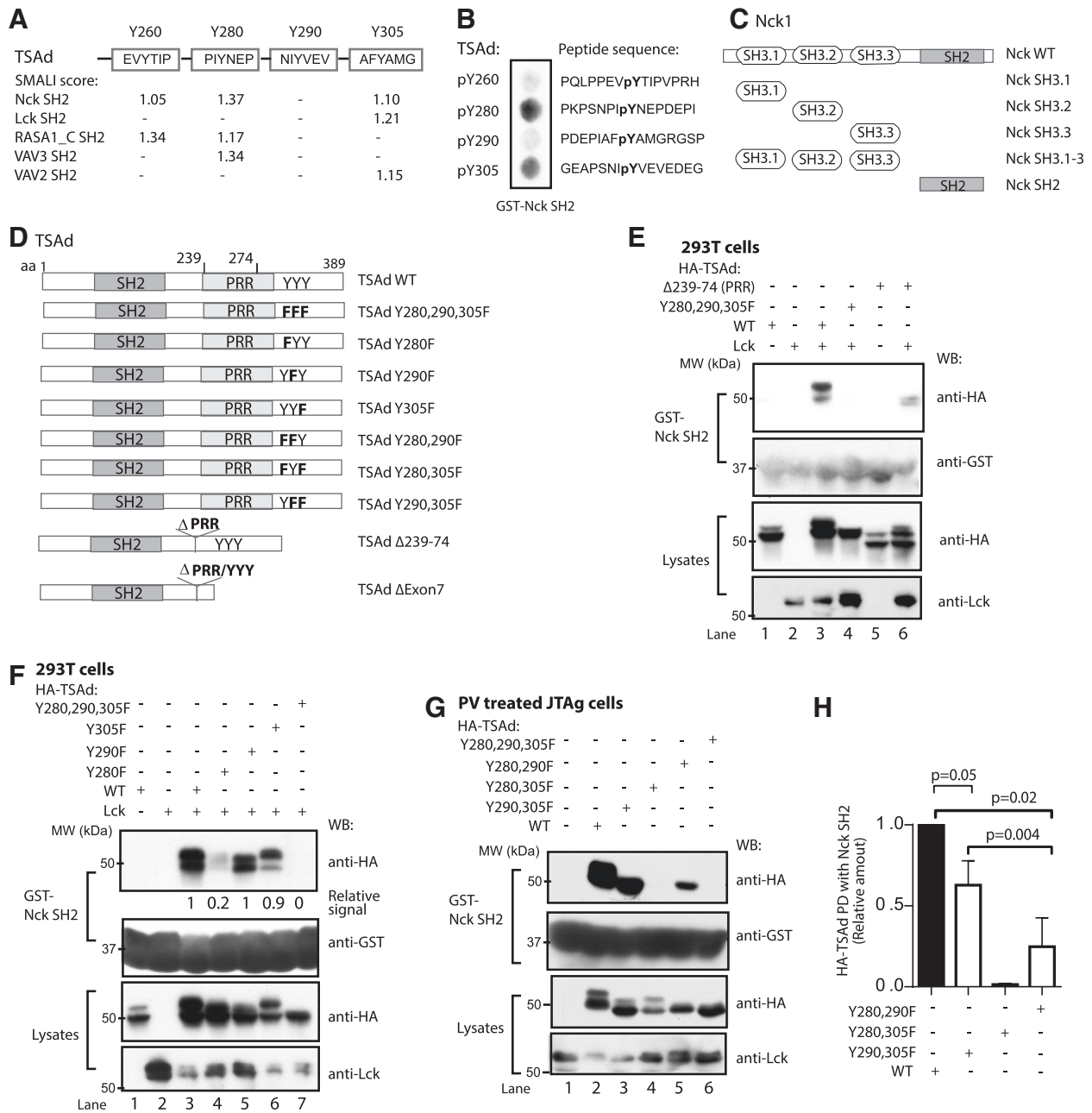


Fig. 1 TSAd-pTyr²⁸⁰ and -pTyr³⁰⁵ interact with the Nck SH2 domain. **a** SH2 domain interaction partners to TSAd phosphotyrosines predicted by SMALI. The five best candidates are listed. Scores > 1 indicate a potential interaction. "-" indicates a SMALI score < 1. **b** Peptide spot array spotted with indicated TSAd phosphopeptides probed with GST-Nck SH2 and developed with anti-GST antibody. SH2: Src homology 2 domain, Y: tyrosine phosphorylation site. **c** and **d** Overview of the Nck (**c**) and TSAd (**d**) Constructs used. WT: wild type, SH3: Src homology 3 domain, PRR: proline rich region, Y: tyrosine phosphorylation site and F: phenylalanine. **e-g** Proteins pulled down by GST-Nck SH2 were resolved on SDS-PAGE, and immunoblotted with the indicated antibodies. **e** GST-Nck SH2 pull-down from 293T cells transiently transfected with Lck and the indicated HA-tagged TSAd cDNA constructs. Lck immunoblot of the GST-Nck SH2 pull-down was negative (data not shown). **f** As in (e), including TSAd encoding single Y → F mutations. Relative TSAd binding to GST-Nck SH2 domain was analysed by ImageJ. Amount of pulled down HA-TSAd WT was set to 1. **g** GST-Nck SH2 pull-down from pervanadate treated JTAG cells transiently expressing indicated TSAd Y → F double mutations. Control GST pull-downs were negative (data not shown). Data shown are representative of at least three experiments (triple Y → F TSAd mutant and double mutants) or two experiments (Y → F single mutants). **h** Graph shows relative amounts of HA-TSAd that were pulled down (PD) with GST-Nck SH2 as in (g) measured by ImageJ. Amount of pulled down HA-TSAd WT was set to 1. Mean values ± SD of three independent experiments (2-tailed paired *t*-test)

analyze double Y to F mutants of TSAad for interaction with Nck SH2 using pull-down experiments in Jurkat TAG (JTAG) cells transiently expressing HA-tagged TSAad (Fig. 1g). To ensure maximal tyrosine phosphorylation of TSAad we chose to treat the TSAad transfected JTAG cells with the strong tyrosine phosphatase inhibitor pervanadate (PV) prior to cell lysis and pull down with Nck SH2. Mutation of both Tyr²⁸⁰ and Tyr³⁰⁵ on TSAad (Y280, 305F) resulted in disrupted interaction between TSAad and GST-Nck SH2 (Fig. 1g, lane 4), while TSAad molecules with either Tyr²⁸⁰ or Tyr³⁰⁵ intact retained ability to interact with Nck SH2. This pattern of Nck SH2 interaction could reflect differences in tyrosine phosphorylation as TSAad pTyr²⁸⁰ is reported twice as frequently in the phosphosite.org database as TSAad pTyr³⁰⁵ (Gopalakrishna et al. *submitted*). However, both TSAad Tyr²⁸⁰ and Tyr³⁰⁵ are phosphorylated by Lck to a similar extent *in vitro* [21], thus the difference in amount of TSAad mutants pulled down could also reflect differences in binding affinity, as indicated by SMALI (Fig. 1b). Taken together, these results confirm the SMALI predicted interaction between the Nck SH2 domain and TSAad pTyr²⁸⁰ and pTyr³⁰⁵, with pTyr²⁸⁰ as the preferred binding site.

The proline rich region of TSAad interacts with the SH3 domains of Nck

In addition to one SH2 domain, Nck contains three N-terminally located SH3 domains (Fig. 1c). PRRs provide possible interaction sites for SH3 domains, and we have previously shown that TSAad aa 239–274 harbors binding sites for Lck and Itk SH3 domains [5, 8]. We thus examined whether the TSAad PRR could bind to any of the three Nck SH3 domains. A Scansite search [22] at low stringency identified the Nck SH3.2 domain as a potential binding partner for TSAad prolines aa 245 (EPSQLLRPKP-PIPAK) and aa 353 (SVIGQGPPPLPHQPPP). Pull-down experiments in 293T cells (similar to Fig. 1e) expressing Lck and various TSAad constructs revealed binding of Nck SH3.1-3 domains to TSAad (Fig. 2a, lane 1). Neither presence of Lck (Fig. 2a, lane 1 and 3) nor mutation of the three penultimate tyrosines (Y280,290,305 F) of TSAad (Fig. 2a, lane 4) affected the interaction with Nck SH3.1-3 while the PRR (Δ 239-274) was required for binding to GST-Nck SH3.1-3 (Fig. 2a, lane 5). Lck has been reported to interact with Nck-SH3 domains [23]. However, Lck was not found to interact with Nck-SH3.1-3 in this experiment (data not shown). These results indicate binding of the Nck SH3 domains to the PRR of TSAad.

We then studied the TSAad interaction to individual Nck SH3 domains in more detail. Peptide spot arrays of 20-mer peptides representing the TSAad PRR showed direct binding of full length GST-Nck SH3.1-3 domains and all three GST-Nck SH3 single domains to TSAad peptides (Fig. 2b). Several classical SH3-ligand motifs are found

within the TSAad PRR peptides that bind to the Nck SH3 domains, including the minimal consensus sequence PxxP, and the canonical K/RxxPxxP and PxxPxK/R motifs. Binding of Nck SH3 domains to TSAad prolines were found to occur primarily within the TSAad PRR aa 239–274. This observation is in agreement with results from pull-down data (Fig. 2a), showing abolished Nck SH3 domain interaction with TSAad Δ 239-274. In comparison, although the C-terminal TSAad sequence aa 359-366 (P359) displayed some reactivity with the Nck-SH3 domains, this part of TSAad probably does not represent important Nck SH3 interaction sites. There are no classical SH3 domain binding motifs within this region, and pull-down experiments showed no binding of Nck SH3 domains to TSAad Δ 239-274 (Fig. 2a) where aa 359–366 are present.

To confirm interaction of Nck SH3 domains with TSAad, we performed pull-down assays from 293T cell expressing TSAad using single GST-Nck SH3 domain constructs. GST-Nck SH3.1 and SH3.3 both interacted with TSAad (Fig. 2c). The Nck SH3 interaction with TSAad was confirmed in JTAG cells transfected with plasmids encoding intact HA-tagged TSAad. All three Nck SH3 domains together (1–3) pulled down similar amounts of TSAad as Lck-SH3 (Fig. 2d, compare lane 1 and 5), while single Nck SH3 domains did not pull-down TSAad (Fig. 2d, lane 2–4). The discrepancy between Nck SH3 binding in 293T and JTAG cells regarding the single Nck SH3 domains may be due to other competing Nck-SH3 ligands present in JTAG cells. The different migration of HA-TSAad that interacted with Nck SH3.1-3 versus single Nck, Lck or Itk SH3 domains could be due to in-gel interference between the GST-fusion proteins and TSAad, either due to the molecular size of the GST protein, or the number of binding sites for TSAad. Taken together, these results show that Nck may interact with TSAad also through its SH3 domains, and that the three SH3 domains interact cooperatively with TSAad.

Both TSAad phosphotyrosines and PRR contribute to TSAad-Nck interaction

The interaction between TSAad and Nck was further analyzed in Nck immunoprecipitates (IPs) from 293T cells expressing Lck together with wild type (WT) or mutated HA-tagged TSAad molecules and GFP-tagged Nck molecules. TSAad WT co-precipitated with Nck (Fig. 2e, lane 1 and 2). Exon7 encodes most of the PRR and the three C-terminal TSAad tyrosines (Fig. 1d), which includes the Nck interaction sites identified in Fig. 1e-g and Fig. 2a-c. TSAad encoded by cDNA lacking exon7 did not co-immunoprecipitate with Nck (Fig. 2e, lane 3). Removal of either the Nck SH2 or SH3 domain interaction sites on TSAad (Δ 239-274 or Y280,290,305F), did not fully abolish TSAad-Nck interaction (Fig. 2e, lane 4 and 5). Collectively, these results show that both the phosphotyrosines and the PRR of TSAad may be engaged in Nck binding.

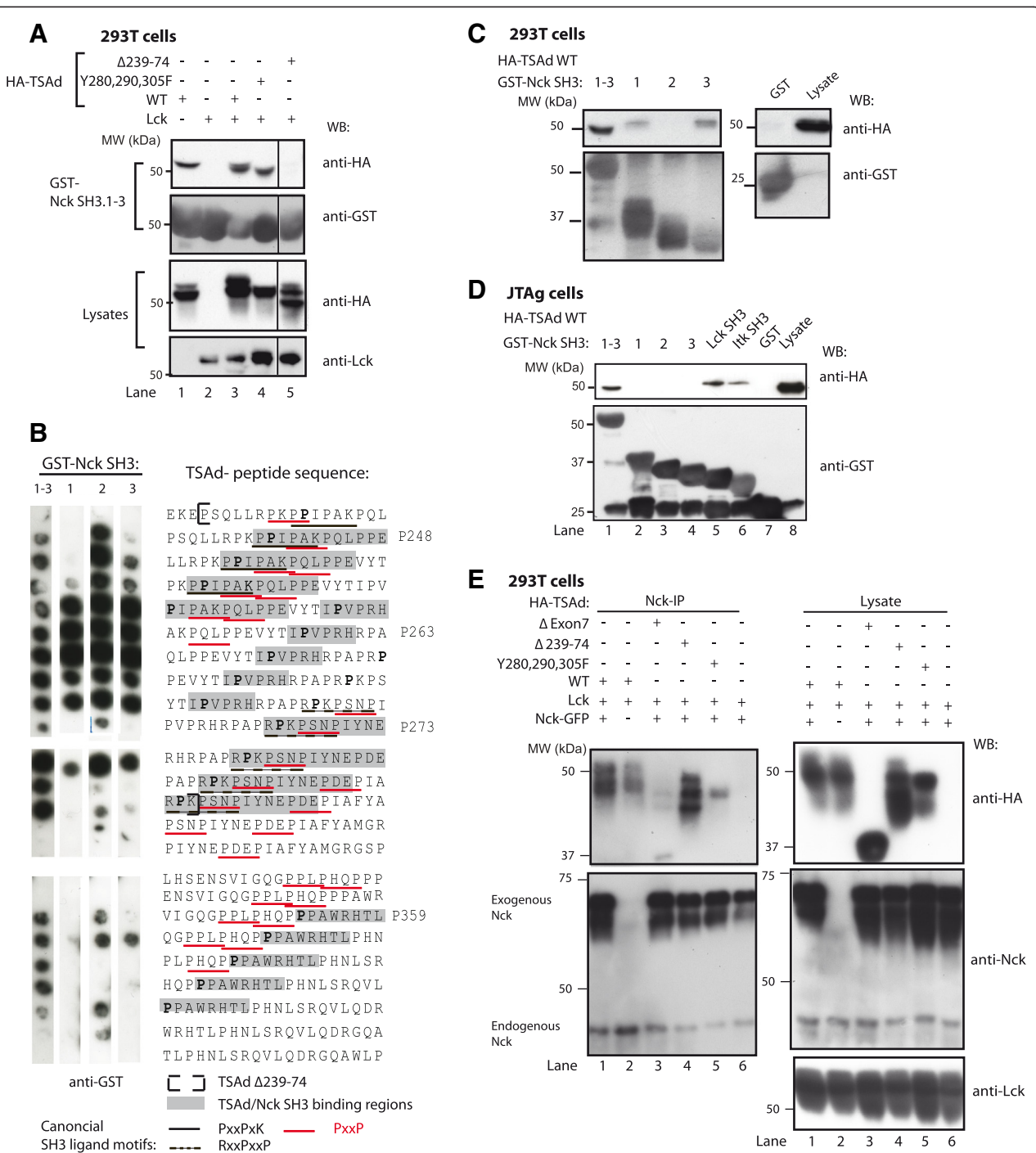


Fig. 2 The proline rich region (PRR) of TSAd interacts with the Nck SH3 domains. **a** GST-Nck SH3 pull-downs from 293T cells transiently expressing the indicated HA-tagged TSAd constructs. Proteins were resolved on SDS-PAGE, and immunoblotted with the indicated antibodies. Control GST pull-downs were negative (data not shown). Lck immunoblot of the GST-Nck SH3 pull-down was negative (data not shown). Results shown are from the same experiment as shown in Fig. 1e and are representative of at least three experiments. **b** Peptide spot array spotted with proline containing TSAd peptides probed with GST-Nck SH3 peptides and developed using anti-GST antibody. Nck SH3 domain interaction sequences on TSAd are boxed in grey, the TSAd region aa 239–274 is outlined and canonical SH3 ligand motifs and the core PXXP motifs are underlined with black or red lines respectively. **c** Pull-down with single GST-Nck SH3 domains and control GST performed as in (a). Results are one representative of three independent experiments. **d** Pull-down of GST-Nck SH3 domains in JTA9 cells transiently expressing HA-TSAd WT. Positive controls: Itk SH3 and Lck SH3. Negative control: GST. Blots were immunoblotted with the indicated antibodies. Data is one representative of two experiments. **e** Nck co-immunoprecipitation of TSAd expressed in 293T cells. Cells were transiently transfected with Nck-GFP, Lck and the indicated HA-tagged TSAd constructs. Cell lysates and anti-Nck IP were separated by SDS-PAGE and immunoblotted with indicated antibodies. Data are representative of three independent experiments

TSAd and Nck co-localize

To further assess whether TSAd and Nck interact *in vivo*, we initially performed co-immunoprecipitation experiments from transiently transfected JTAG cells expressing exogenous TSAd and Nck. TSAd and Nck were found to co-immunoprecipitate in these cells (Fig. 3a). We then used confocal microscopy to examine the localization of endogenous Nck and TSAd in CD3/CD28 activated human CD3+ T cells (Fig. 3b), as well as in JTAG cells transiently expressing Nck-GFP and TSAd-mCherry (Fig. 3c). This analysis indicated that TSAd and Nck co-localized in both human CD3+ T cells and in transfected JTAG cells. TSAd lacking both the Nck SH2 and SH3 interaction sites (TSAd Δ Exon7), did not co-immunoprecipitate nor co-localize with Nck (Fig. 3a and c). A faint band was observed for the TSAd Δ Exon7 co-immunoprecipitate with Nck (Fig. 3a). This band is not specific since it was similar to background (without the Nck antibody, data not shown).

Co-localization between Nck and intact or mutated TSAd molecules in JTAG cells was then quantified by

imaging flow cytometry (IFC) (Fig. 3d) using the bright detail similarity feature of the IDEAS software (Fig. 3e and f). Compared to TSAd WT, mutation of the Nck SH2 interaction site TSAd Tyr²⁸⁰, did not significantly affect Nck co-localization (Fig. 3f). However, when expressing TSAd mutated for all three C-terminal TSAd-tyrosines (i.e. Y280,290,305F), co-localization with Nck was significantly reduced. Again, absence of both the PRR and the pTyr sites on TSAd (Δ Exon7) resulted in disruption of Nck and TSAd co-localization (Fig. 3f). Taken together, these results support the notion that TSAd and Nck interact *in vivo*, and that both Nck SH2 and SH3 domains contribute to this interaction.

TSAd promotes interaction between Nck and Lck

We and others have previously shown that the Src kinase Lck interacts with TSAd [4, 5, 18, 21, 24]. Additionally, Nck has been reported to bind to a proline motif in the unique domain of Lck [23]. We here examined whether TSAd may also serve as a molecular scaffold

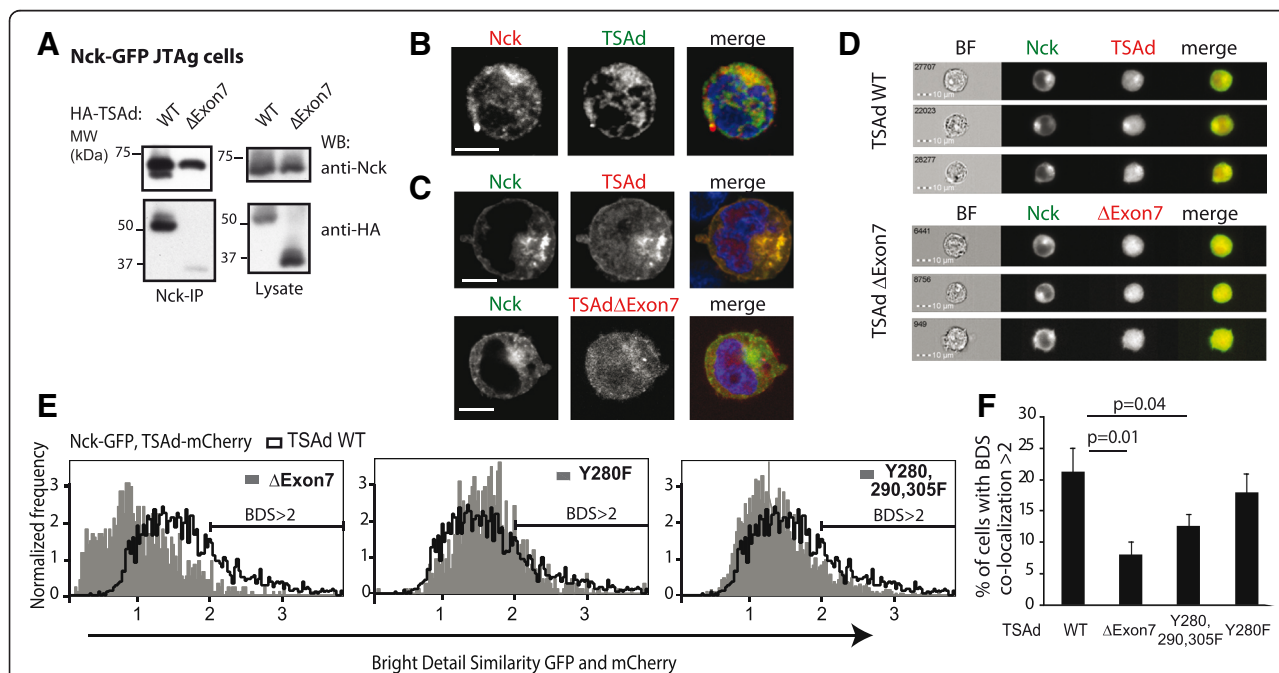


Fig. 3 Co-localization of TSAd and Nck in T cells. **a** Nck co-immunoprecipitation of TSAd expressed in JTAG cells. Immunoblot of Nck-IP and lysate from JTAG cells transfected with Nck-GFP and HA-TSAd constructs. One representative of three experiments is shown. **b** Confocal microscopy images (60x magnification) of one CD3/28 activated human CD3+ T cell, stained with anti-Nck (red), anti-TSAd (green) and Hoechst 33342 (blue nuclear staining in the merged image). Scale bar = 5 μ m. Images are representative of two independent experiments. **c** Confocal microscopy images (100x magnification) of transfected, fixed and permeabilized JTAG cells showing Nck-GFP (green), TSAd-mCherry (red) and merged images with nuclei staining Hoechst 33342 (blue). Images are representative of three independent experiments. Scale bar = 5 μ m. **d** and **e** Co-localization of Nck-GFP and the indicated TSAd-mCherry molecules analyzed with the ImageStream cytometer, gated on double positive GFP and mCherry cells. **d** Gallery of representative IFC images. **e** Co-localization analyzed with the "Bright Detail Similarity" (BDS) algorithm in the IDEAS software. The BDS score is the log transformed Pearson's correlation coefficient of the localized bright spots with a radius of 3 pixels or less within the masked area in the two input images. Overlay of BDS histograms comparing co-localization of various TSAd molecules to Nck are shown. Data are representative of four independent experiments. **f** Percentage of cells that shows high degree of co-localization (BDS score > 2) in (e) for Nck-GFP with each of the TSAd constructs. Graph shows mean \pm standard deviation from four experiments. P-values are indicated (2-tailed paired t-test)

bringing Lck into the vicinity of Nck. We initially used 293T cells as these cells lack expression of T-cell specific proteins and thus provide a “pure” system to investigate T cell protein interactions. Cells were transiently transfected with Nck, Lck and/or TSAAd cDNA constructs, and subjected to Nck-IP (Fig. 4a). No interaction of Nck with Lck was observed (Fig. 4a, lane 2). However, in cells co-expressing TSAAd, Lck co-immunoprecipitated with Nck (Fig. 4a, lane 3). Moreover, while Nck was tyrosine phosphorylated in the presence but not in the absence of Lck, the level of Nck tyrosine phosphorylation was strongly increased in the presence of TSAAd (Fig. 4a, pTyr-blot, compare lane 2 and 3).

To assess whether TSAAd promotes Nck-Lck interaction also in T cells, we performed Nck co-immunoprecipitation in lysates from control JTAG cells or JTAG cells where endogenous TSAAd expression had been suppressed by siRNA. Briefly, cells were stimulated with PMA and ionomycin over night to increase the amount of endogenous TSAAd, washed, and restimulated with anti-CD3 antibodies. Nck co-precipitated with TSAAd in these cells (Fig. 4b) but when restimulated with anti-CD3, less interaction of Nck with TSAAd was observed (Fig. 4b and c). Similarly, TSAAd promoted the interaction of Nck with Lck in JTAG cells. However less Lck was associated with Nck in anti-CD3 stimulated cells (Fig. 4d and e).

Finally we examined the influence of TSAAd on Nck-Lck association in primary T cells, using CD4⁺ murine *SH2D2A*^{-/+} or *SH2D2A*^{-/-} blast T cells. The *SH2D2A* gene encodes TSAAd. In line with the increased Lck co-immunoprecipitation with Nck in 293T and JTAG cells, more Nck associated Lck molecules was observed in *SH2D2A*^{-/+} blast T cells compared to the TSAAd deficient *SH2D2A*^{-/-} blast T cells (Fig. 4f and g). Taken together, these data suggest that TSAAd promotes interaction between Nck and Lck. In the presence of all three molecules, Nck may become phosphorylated by Lck.

TSAAd promotes Nck-SLP-76 interaction

Nck is known to bind to the cytosolic adaptor SLP-76 [13] as well as the guanine nucleotide exchange factor Vav1 [11]. Since we had established TSAAd as an Nck interaction partner, and since TSAAd influenced the interaction of Nck with Lck, we explored whether TSAAd also influences interaction of SLP-76 or Vav1 with Nck. Western blotting of Nck immunoprecipitated from PMA/ionomycin treated JTAG cells, showed a significant reduction in Nck and SLP-76 interaction in the absence of TSAAd, both in resting and CD3 restimulated cells (Fig. 5a and b). In contrast, similar amount of Vav1 was co-immunoprecipitated with Nck in the presence or absence of TSAAd (Fig. 5a and c). These data suggests that TSAAd not only promotes Nck interaction with Lck, but

also influences Nck's interaction with some but not all of its partners.

TSAAd promotes F-actin polymerization in T cells

Nck is a known regulator of the actin cytoskeleton [9, 15, 17]. The interaction between TSAAd and Nck thus led us to examine the effect of TSAAd on polymerized actin (F-actin) in JTAG cells. JTAG cells were transfected with cDNA constructs encoding TSAAd WT-mCherry or TSAAd-mCherry lacking both the Nck SH2 and the Nck SH3 interaction sites (TSAAd Δ Exon7), and amount of F-actin was quantified with IFC (Fig. 6). As an internal reference in each sample, we calculated the relative increase in F-actin in cells that were transfected (mCherry +) compared to non-transfected cells (mCherry-) (Fig. 6a). TSAAd WT-mCherry transfected cells displayed significantly more F-actin than cells expressing either TSAAd- Δ Exon7-mCherry or mCherry alone (Fig. 6a and b). Following anti-CD3 stimulation, an increase in F-actin content was observed both in the mock transfected and the TSAAd transfected cells (Fig. 6c). However, the relative increase in F-actin amount in cells expressing TSAAd WT was significantly less pronounced than for cells expressing mCherry alone (mock) or TSAAd- Δ Exon7-mCherry (Fig. 6c and d). Taken together, these data show that TSAAd influence F-actin content in both resting and CD3 stimulated cells, and that this effect is dependent on the Nck and Lck interaction sites on TSAAd.

Discussion

In this work we identified Nck as a novel interaction partner for TSAAd. Through peptide array analysis, precipitation assays and co-localization studies, we found that Nck and TSAAd interact both through the Nck SH2 domain and the Nck SH3 domains. Moreover, TSAAd promoted interaction of Nck with Lck and SLP-76. Cells expressing TSAAd displayed increased amount of polymerized F-actin, while TSAAd lacking the Nck and Lck interaction sites, failed to do so. Interaction of TSAAd with Nck may represent one pathway whereby TSAAd modulates actin polymerization events.

We used scoring matrix-assisted ligand identification (or SMALI), a web-based program for predicting binding partners for SH2-containing proteins [20] to identify potential novel interaction partners for TSAAd. The SH2 domain binding motives were identified by SMALI based on data from an oriented peptide array library (OPAL) screening. OPAL allows systematically testing the residues in the phosphopeptide one by one. The SMALI analysis covers 76 of the 120 SH2 domains in the genome. Nck1 SH2 binding to TSAAd pTyr²⁸⁰ displayed the best SMALI score for that particular position in TSAAd. A limitation of using OPAL screening or phage display techniques to identify SH2 domain binding motifs is that

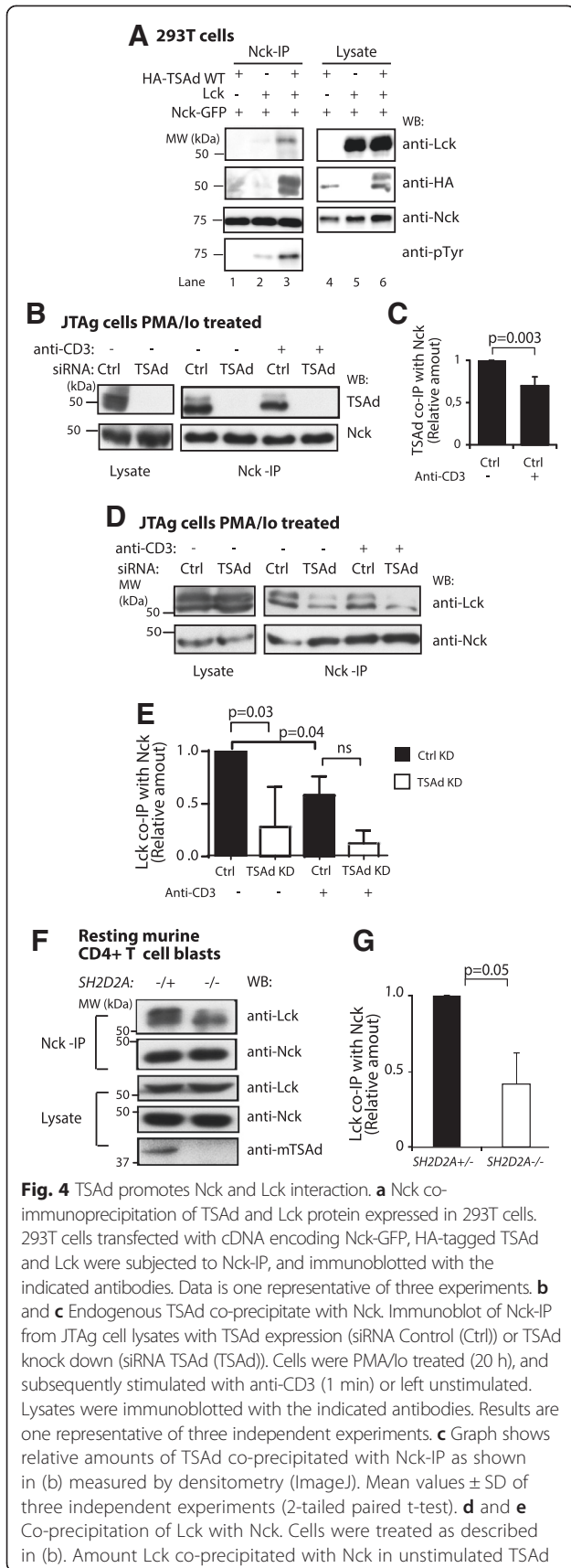


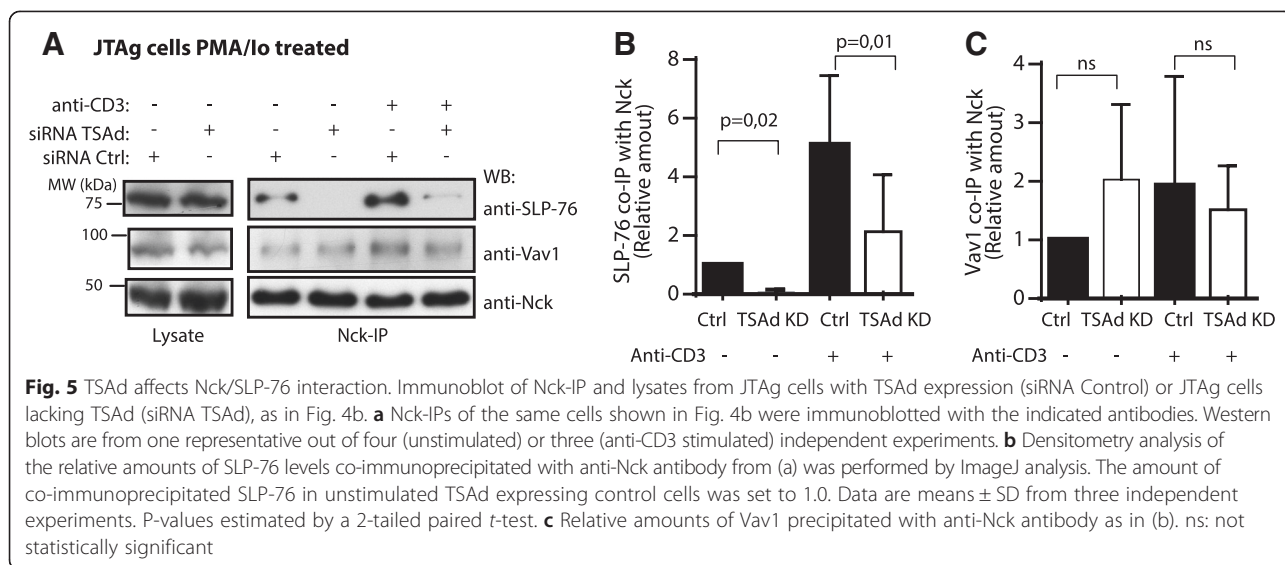
Fig. 4 TSAd promotes Nck and Lck interaction. **a** Nck co-immunoprecipitation of TSAd and Lck protein expressed in 293T cells. 293T cells transfected with cDNA encoding Nck-GFP, HA-tagged TSAd and Lck were subjected to Nck-IP, and immunoblotted with the indicated antibodies. Data is one representative of three experiments. **b** and **c** Endogenous TSAd co-precipitate with Nck. Immunoblot of Nck-IP from JTAG cell lysates with TSAd expression (siRNA Control (Ctrl)) or TSAd knock down (siRNA TSAd (TSAd)). Cells were PMA/Io treated (20 h), and subsequently stimulated with anti-CD3 (1 min) or left unstimulated. Lysates were immunoblotted with the indicated antibodies. Results are one representative of three independent experiments. **c** Graph shows relative amounts of TSAd co-precipitated with Nck-IP as shown in (b) measured by densitometry (ImageJ). Mean values \pm SD of three independent experiments (2-tailed paired *t*-test). **d** and **e** Co-precipitation of Lck with Nck. Cells were treated as described in (b). Amount Lck co-precipitated with Nck in unstimulated TSAd

expressing cells (siRNA Control) was set to 1. KD: knock down. Mean values \pm SD of three independent experiments (2-tailed paired *t*-test). ns: not statistically significant. **f** Nck-IP and lysates from resting CD4+ murine *SH2D2A*^{-/+} or *SH2D2A*^{-/-} blast T cells. Results shown are one representative out of three independent experiments. **g** Graph shows relative amount of Lck co-precipitated with Nck IP as shown in (f) as measured by ImageJ. Amount of Lck co-precipitated with Nck in *SH2D2A*^{-/+} CD4+ T cells was set to 1. Mean values \pm SD of three experiments. P-values are indicated (2-tailed paired *t*-test)

amino acids that are prohibitive for SH2 domain interaction in a given position may go undetected [25]. Also, the SMALI analysis only covers 60 % of the existing SH2 domain in the human genome. Thus it is possible that there are other SH2 domain containing proteins that also may be predicted to interact with TSAd pTyr²⁸⁰ with a similar probability.

Nck was found to interact with both TSAd pTyr²⁸⁰ and pTyr³⁰⁵, with pTyr²⁸⁰ as the preferred binding site. We have previously shown that TSAd pTyr²⁸⁰, pTyr²⁹⁰ and pTyr³⁰⁵ provide interactions sites for the Lck SH2 domain [5, 21]. Moreover, we previously showed that Lck interacts with the same TSAd PRR [18] as Nck (this paper). It thus appears that Nck and Lck may compete for the same binding sites on TSAd. However, since TSAd promoted the association of Nck with Lck, it is possible that these two molecules may also bind to TSAd simultaneously. Our current and previous data suggest that Nck- and Lck-SH2 domains have different preferences regarding the TSAd phosphotyrosines, with the pTyr²⁹⁰ [21] or pTyr³⁰⁵ as the preferred Lck SH2 binding sites. Of note is that the three C-terminal tyrosines of TSAd are highly conserved [5]. This points to a critical role for these tyrosines in TSAd's function. Similar to Lck, Nck has multiple binding sites on TSAd. This may increase the avidity between Nck and TSAd, and allow for simultaneous interaction of TSAd with both Nck and Lck. We did not directly examine whether Nck and Lck may dock onto the three TSAd pTyr simultaneously. However, one intriguing possibility is that one important function of the C-terminal TSAd tyrosines is to serve as docking sites for both the Nck and Lck SH2 domains, bringing Lck into the vicinity of Nck and its binding partners.

In pull-down assays from 293T cells, the Nck SH3 domains 1 and 3 were found to interact, presumably in a cooperative manner with the PRR of TSAd, while in Jurkat cells none of the single SH3 domains were able to pull down TSAd. This could be due to competition of individual Nck SH3 domains with other molecules present in Jurkat cells but not 293T cells, where Lck itself is a good candidate. The Nck SH3.2 domain did not show interaction with TSAd. This is in agreement with previously reported Nck SH3 interactions. Different Nck

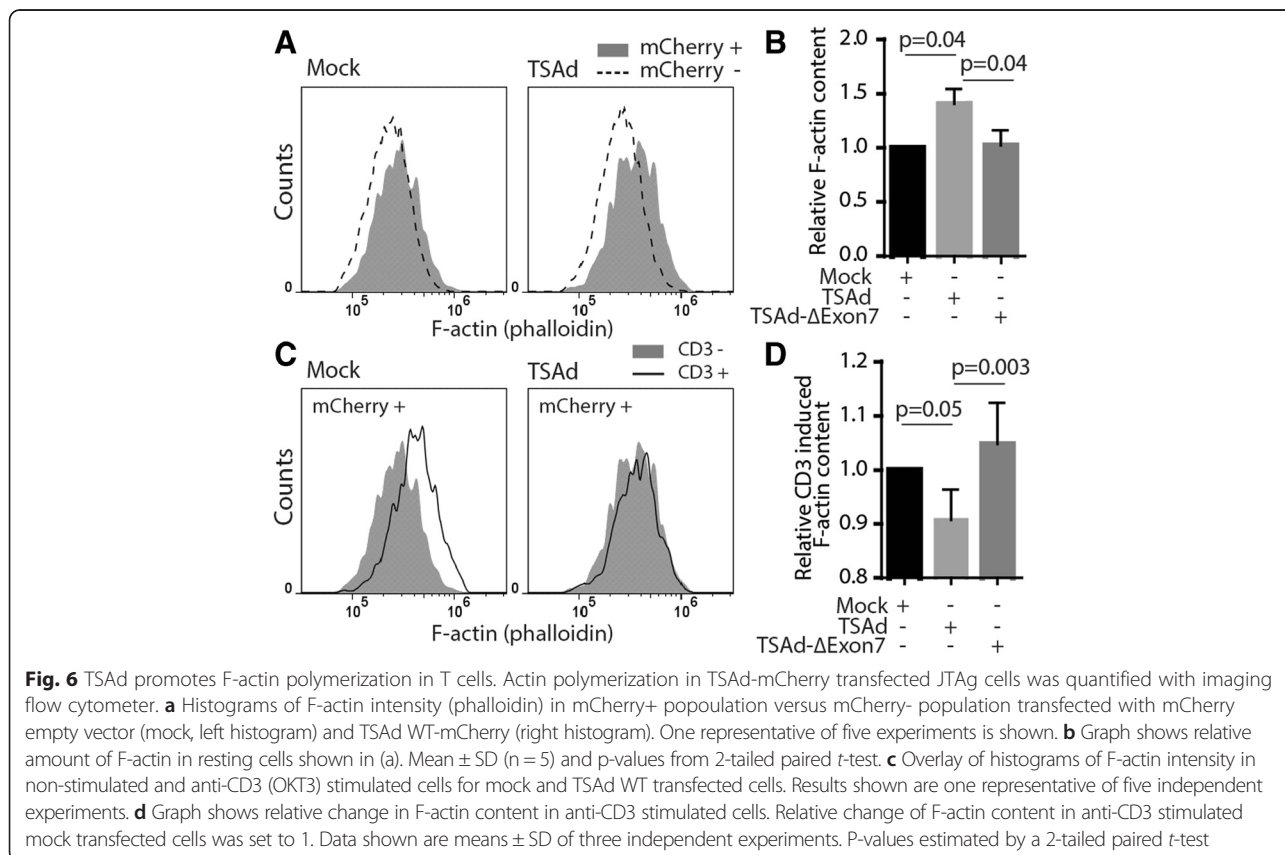


interaction partners show a clear preference to individual SH3 domains, and the interaction between Nck ligands and the different Nck SH3 domains are often found to be cooperative [10].

TSAd is expressed only at low levels in naïve T cells, thus to examine the effect of TSAd we here either used overexpression of exogenous TSAd or stimulated T cells

with PMA/ionmycin or anti-CD3 to ensure expression of TSAd. In these cells, Nck and TSAd were found to constitutively interact without additional stimulation.

Both confocal microscopy and IFC supported the notion that Nck and TSAd are co-localized in the cytoplasm of unstimulated Jurkat cells. The intracellular localization of TSAd has previously been assessed [3, 18, 26]. Although



both we [18] and others [26] previously noted that TSAd was also located in the nucleus, we observe mainly cytoplasmic localization of TSAd in unstimulated cells. We thus favor the notion that TSAd is primarily a cytosolic adaptor, linking Lck with some of its substrates, including also Nck.

In the presence of both TSAd and Lck, Nck phosphorylation was increased. Phosphorylation of Nck has also been reported previously. However, the physiological significance of this phosphorylation event is not clear. Nck phosphorylation is increased following TCR engagement in Jurkat cells [27]. c-Abl has been found to phosphorylate Tyr¹⁰⁵ on Nck1, providing a negative feedback loop on p38 activation upon VEGF-A signaling [28]. Furthermore, Fyn has been reported to phosphorylate Nck. A model for VEGFR-2 signaling was thus proposed, whereby the phosphorylated VEGFR-2 recruits Nck via its SH2 domain, leading to phosphorylation of Nck and subsequent recruitment of Fyn and phosphorylation of Nck by Fyn [29]. We have previously reported that TSAd is recruited to the activated VEGFR-2 receptor and that in its absence, VEGF-A stimulation fails to induce actin stress fiber formation [7]. Moreover, TSAd interacts with Src in endothelial cells, and is required for activation of Src upon VEGF signaling [6]. Although it needs to be formally demonstrated, our current study suggests that TSAd may also recruit Nck to the activated VEGFR-2, providing a possible link between VEGFR-2 signaling and the actin cytoskeleton.

The role of Nck in T cell activation has been strongly debated, and reports have been contradictory. Elimination of Nck with siRNA knock down in Jurkat cells leads to cells that are hyporesponsive to TCR stimulation [30]. Recruitment of Nck via its SH3.1 domain to a PRR in the CD3 ϵ , has been proposed to be the earliest event in TCR triggering [15, 31–34]. Still, the role of CD3 ϵ -Nck interaction in T cell activation remains controversial [32, 35, 36]. It is possible that Nck recruits a priming tyrosine kinase to CD3 ϵ , most likely the Src kinase Lck to allow for the phosphorylation of adjacent CD3 ζ subunits [37]. It has previously been reported that Nck binds to the unique domain of Lck via its SH3 domains, and that phosphorylation of Ser⁵⁹ disrupts the interaction [23]. Our current work suggests that TSAd may link Nck with Lck, providing a scaffold to which both molecules may bind simultaneously. Alternatively TSAd may alter Nck or Lck or both thus promoting subsequent interaction of these molecules with each other. We observed a TSAd independent interaction of Lck with Nck both in murine T cell blasts lacking TSAd, and in TSAd suppressed JTAG cells. Since we failed to observe any interaction between Lck and Nck-SH3.1-3 nor Nck-SH2 domains in pull-downs in 293T cells, a possible explanation is that a different adapter molecule exist in T cells that bridges Lck and Nck.

Whether TSAd may be a missing link between CD3 ϵ , Nck and a kinase, is unclear. It has previously been reported that TSAd is essential for activation of Lck [38]. In contrast we have previously shown that in the absence of TSAd, CD3 ζ is more highly phosphorylated upon TCR stimulation [21], and vice versa, that when TSAd is overexpressed, CD3 ζ is less phosphorylated [39]. Our interpretation of this apparent paradox is that TSAd both serves to activate Lck by interacting with the Lck SH3 and SH2 domains, and at the same time provides tyrosines that are substrates for Lck [5]. TSAd may thus serve as a competitive substrate “sink” allowing temporary and spatial control of Lck activity, allowing particular substrates, such as Nck bound to TSAd, to become phosphorylated, while other substrates may be protected from Lck. With respect to the possible role of Nck-TSAd-Lck interaction for initiation of TCR signaling, one caveat is also that TSAd is expressed only at low levels in naïve resting cells. We thus favor a model where TSAd promoted interaction between Nck and Lck play a role preferentially in experienced human T cells, where TSAd expression is already induced. In experienced cells TSAd may contribute to actin dependent processes, such as migration [8, 40] or stabilization of the immunological synapse [21].

TSAd also promoted the interaction of Nck with SLP-76. Nck interacts via its SH2 domain with SLP-76 pTyr¹¹³ and pTyr¹²⁸ upon TCR engagement [11, 13, 17]. Accordingly, we also observed more Nck associated with SLP-76 in our restimulated JTAG cells. In comparison Nck-Vav1 dimers are proline dependent and are constitutively formed [11, 12] which fits with our observation that the interaction between Nck and Vav1 remained unaffected by TSAd.

How TSAd may affect SLP-76 interaction with Nck remains to be studied. An intriguing observation was that upon CD3 stimulation, less TSAd and more SLP-76 was associated with Nck. Our observation that Lck is recruited to Nck in the presence of TSAd, indicates that the Lck and TSAd together primes Nck for interaction with SLP-76. Additionally, the kinase Itk is also a binding partner for both SLP-76 [12] and TSAd [8, 21, 41]. One possibility is that TSAd, Lck, Itk, Nck and SLP-76 participate in a multimolecular complex, where each of the molecules may interact with several of the other molecules in a cooperative manner.

Conclusions

We report a novel association between TSAd and Nck. In the presence of TSAd, association of Nck with Lck as well as SLP-76 is increased. This may allow recruitment of Lck into the vicinity of Nck binding partners, and thus provide a mechanism whereby TSAd influence the actin cytoskeleton.

Methods

Plasmids and constructs

TSAd cDNA was cloned into the pEF-HA expression vector, and constructs encoding full-length TSAd [39], TSAd Δ 239-274, TSAd Δ Exon 7 (aa 239–334) [18], single, double and triple TSAd tyrosine (Y) to phenylalanine (F) mutants [5] were cloned as previously described. Point mutations in TSAd cDNA were generated by QuickChange™ Site-Directed Mutagenesis (Stratagene). mCherry tagged TSAd constructs were cloned into the pcDNA3 vector. Nck1-GFP and GST-Nck1 constructs were provided by L. Buday [42]. The GST-Lck SH3 and GST-Itk SH3 constructs were generated as described [4, 8, 18]. All constructs were verified by sequencing.

Antibodies

The following monoclonal antibodies were used: anti-Lck (3A5), anti-SLP-76 (F7), and anti-GST (B12) (Santa Cruz Biotechnology, Inc.), anti-HA (HA.11, Bio Site), anti-Nck (108; BD Transduction Laboratories™), anti-human CD3 ϵ (OKT3; American Type Culture Collection), anti-phosphotyrosine (4G10; Upstate Biotechnology), and unconjugated and DyLight 488 conjugated anti-TSAd (anti-SH2D2A) (3C7, Origene). The following polyclonal antibodies were used: anti-Nck (Millipore), anti-Vav (C-14; Santa Cruz Biotechnology, Inc.) and anti-TSAd (1715 T, provided by V. Shapiro). Horse radish peroxidase (HRP)-conjugated goat anti-mouse IgG, goat anti-mouse IgG light chain specific and goat anti-rabbit IgG (all from Jackson ImmunoResearch Laboratories), and goat-anti rabbit IgG Alexa Fluor® 647 conjugate (Molecular Probes) were used as secondary antibodies.

Expression and purification of recombinant GST-fusion proteins

GST-fusion proteins of Nck SH2, Nck SH3, Lck SH3 and Itk SH3 domains were produced in *E. coli* BL21-Codon Plus® (Stratagene) in M9 minimal salt media at 15 °C, and purified on Glutathione 4B Sepharose™ beads (GE Healthcare), according to the manufacturer's instructions. Protein concentration and purity were analyzed by SDS-PAGE and CommaSsie Brilliant Blue Staining.

Cell cultures and transfections

Human embryonic kidney (293T) cells (American Type Culture collection) and Jurkat SV40 T antigen (JTAG) cells [43] were cultured in complete RPMI (cRPMI) medium [RPMI 1640 supplemented with 5 % (293T cells) or 10 % FBS (JTAG cells), 1 mM sodium pyruvate, 10 mM HEPES, 1 % MEM non-essential amino acids, 100 units/ml penicillin, 100 μ g/ml streptomycin (all from GIBCOBRL®, Life Technologies™) and 0.5 μ M 2-mercaptoethanol (Sigma). Transfections of $1.5 \cdot 10^7$ JTAG

cells in RPMI 1640 with 0.5–10 μ g of plasmid DNA were performed using a BTX electroporator (Genetronix) at 240 V and 25 ms. Transient transfectants were cultured for 16–24 h. 293T cells ($2.2 \cdot 10^6$) were plated in 5 ml complete media on 10 cm cell culture plates, and allowed to adhere for 24 h. The transfection solution containing 0.04–6 μ g DNA and 42.4 μ g/ml polyethylenimine (PolyScience) in PBS was added drop-wise to the 293T cells. Transfected cells were further propagated for 16–24 h.

siRNA-mediated knock down of TSAd expression

JTAG cells ($1.5 \cdot 10^7$) were transiently transfected with 1 μ M solution of TSAd siRNA [7] to inhibit TSAd expression or control siRNA (24 h in total), as previously described [18]. After transfection, cells were stimulated with 50 ng/ml phorbol 12-myristate 13-acetate (PMA) and 500 ng/ml Ionomycin (Io) (both from Sigma) for 20 h to induce TSAd expression.

Mice and propagation of murine CD4+ T blast cells

Mice used in this study were bred under conventional conditions, and approved by The National Animal Research Authority, via their local representative at the University of Oslo. The mice were regularly screened for common pathogens and housed in compliance with guidelines set by the Experimental Animal Board under the Ministry of Agriculture of Norway. TSAd deficient (*SH2D2A*^{-/-} or RIPB-KO) C57BL/6 mice were kindly provided by Professor J. A. Bluestone [41] and the *SH2D2A*^{-/-} mice were backcrossed >10 generations into C57BL/6 mice and maintained on a C57BL/6 background as described [7]. Murine splenocytes and lymph node cells were obtained by crushing the organs through a cell strainer (70 μ m nylon, BD Biosciences). CD4+ T cells were purified from splenocytes or lymph node by negative selection (DynaL® Mouse CD4 Negative Isolation Kit (114.15), Life Technologies). The recovered population was more than 93 % CD4+ T cells, as analyzed by flow cytometry (FACS Calibur, BD Biosciences). For propagation of blast cells from primary murine CD4+ T cells, 30U mouse IL-2/ml and Dynabeads® Mouse T-Activator CD3/CD28 (Life Technologies) (following the manufacturer's protocol) was added to the cRPMI medium.

Isolation and activation of human CD3+ T cells

Peripheral blood mononuclear cells (PBMCs) from healthy donors were isolated from buffy coats using Lymphoprep™ as previously described [44]. CD4+ T cells were isolated by negative selection (Dynabeads® Untouched Human T cells (113.44D), Life Technologies). The recovered population was more than 95 % CD3+ T cells, as analyzed by flow cytometry (FACS Calibur, BD Biosciences). Cells were activated with Dynabeads®

Human T-Activator CD3/CD28 (one bead per cell), following the manufacturer's protocol (Life Technologies) in cRPMI medium for 48 h.

Cell stimulation and lysis

Prior to use in experiments, the anti-CD3/CD28 beads were removed from long term activated murine CD4⁺ T cells and cells were rested for 24–48 h before harvesting. For short time activation of JTAG cells, the cells were washed with PBS, resuspended to $5 \cdot 10^7$ cells/ml and stimulated with 5 μ g/ml anti-CD3 (OKT3) antibody for the indicated time points. In some experiments pervanadate treatment for 5 min (0.01 % H₂O₂, 100 μ M Na₃VO₄) was used. Stimulation was terminated by adding ice cold PBS and centrifugation. Cells were lysed in lysis buffer containing; 1 % Nonidet-P40, 50 mM n-octyl- β -D-glucoside, 25 mM Tris (pH 7.5), 100 mM NaCl, 20 mM NaF, 1 mM Na₃VO₄ and 1 \times protease inhibitor cocktail (Roche) for 30 min on ice. For co-immunoprecipitation, reduced amounts of detergents were used; 0.25 % Nonidet-P40 and 12.5 mM n-octyl- β -D-glucoside. LDS lysis were used for immunoprecipitation in murine CD4⁺ T cells (final concentration) 0.1 % LDS, 1 % Triton X-100, 0.1 M LiCl, 1 mM PMSE, 5 mM EDTA, 50 mM Hepes, 1 mM Na₃VO₄ and 1 \times protease inhibitor cocktail.

Immunoprecipitation and pull-down assays

For immunoprecipitation (IP), lysates from 293T and JTAG cells were pre-cleared three times for 30–45 min with Dynabeads Protein G (Life Technologies), and incubated with the relevant antibodies conjugated to Dynabeads Protein G (Life Technologies) for 1 h at 4 °C. For pull-down assays, cell lysates from $2.2 \cdot 10^6$ – $1.5 \cdot 10^7$ cells were pre-cleared three times for 30 min with a 2:1 mixture of GST/Glutathione SepharoseTM 4B beads (GE Healthcare), and added to aliquots of Glutathione SepharoseTM 4 Fast Flow (GE Healthcare) coupled GST-fusion proteins. The mixture was rotated for 1 h at 4 °C. Beads from IP or pull-down were washed three times in 1 \times lysis buffer, and precipitated proteins were separated by SDS-PAGE and detected by immunoblotting. Quantification was performed on ECL autoradiography films using ImageJ software [45].

SDS-PAGE and Western blot

Proteins were denatured in SDS loading buffer, separated by SDS-PAGE, and transferred to a PVDF membrane (Bio-Rad Laboratories) using a Hoefer Semi-Phor Semi-Dry Transfer Unit (Amersham Biosciences), or a Trans-Blot[®] Turbo[™] Transfer System (BioRad). Blots were blocked in PBS-T (pH 7.4, 0.1 % Tween) with 3 % BSA (Biotest) or 3 % skimmed milk (Sigma), and subsequently incubated with the indicated antibodies in the

appropriate blocking buffer. Signals were detected by horseradish peroxidase (HRP)-labeled secondary antibodies and Super Signal[®] west Pico Stable Peroxide Solution (Pierce), and developed on Amersham Hyperfilm[™] ECL (GE Healthcare).

SMALI predictions of SH2 interactions

The web based scoring matrix-assisted ligand identification (SMALI) algorithm [19, 20], was used to predict SH2 domains interacting with phosphotyrosine peptides of TSAAd [NP_003966.2]. A domain scan was carried out. A relative SMALI score greater than 1.0 indicated a strong potential binding.

Peptide spot array analysis

Peptide arrays were synthesized on nitrocellulose membranes using a MultiPep automated peptide synthesizer (INTAVIS Bioanalytical Instruments AG) as described [46]. Membranes were spotted with relevant TSAAd peptides and probed with GST-tagged Nck SH2 and SH3 fusion proteins (50–150 μ g/ml), followed by anti-GST antibody and anti-mouse HRP secondary antibody. Signals were detected by chemiluminescent detection by Super Signal[®] West Pico stable peroxide solution (Pierce).

Confocal microscopy

JTAG transfectants were harvested 16–24 h post transfection for confocal microscopy analysis. For human CD3⁺ T cells activated with anti-CD3/CD28 Dynabeads, beads were removed. $1 \cdot 10^6$ cells were washed in PBS, adhered to polylysine microscope slides (VWR) for 10 min, fixed with 4 % paraformaldehyde (Fluka Chemika) for 10 min, and rinsed twice with washing buffer (PBS, 2 % FCS). Subsequently, the cells were permeabilized with 0.1–0.3 % Triton X-100 in PBS for 5 min, stained with 2 μ g/ml Hoechst 33342 (Molecular Probes) for 20 min, rinsed twice with washing buffer. Human CD3⁺ T cells were further stained with 5 μ g/ml anti-Nck (Millipore) and anti-TSAAd DyLight 488 (2.5 μ g/ml) for 1 h, rinsed three times and stained with goat-anti rabbit IgG Alexa Fluor[®] 647 (10 μ g/ml) for 1 h, rinsed three times with washing buffer. Both Jurkat cells and human CD3⁺ T cells were rinsed once with dH₂O before mounted with SlowFade[®] Gold (Life Technologies) and sealed with nail polish. Confocal images were acquired on an Olympus Fluoview FV1000 BX61WI upright microscope equipped with a PlanApo 100 \times NA 1.40 oil objective.

Imaging flow cytometry

JTAG cells co-transfected with Nck-GFP and TSAAd-mCherry constructs were harvested 16–24 h post transfection. Cells were incubated with 2 μ g/ml Hoechst 33342 (Molecular Probes) for 20 min, washed in PBS,

fixed with 4 % paraformaldehyde (Fluka Chemika) for 10 min, and rinsed twice with PBS. Data of $1-3 \cdot 10^4$ cells per sample were acquired on a 12-channel ISX Imaging Flow Cytometer with 40× objective (Amnis Corporation). Samples were acquired with a bright-field (BF) area lower limit of $50 \mu\text{m}^2$ to eliminate debris. Single stained controls were collected with no illumination and with excitation lasers switched on. The data was analyzed using IDEAS 4.0 software (Amnis). The IDEAS compensation wizard was used to create a compensation matrix for single stained image files. The matrix was used to correct for spectral overlap in raw sample files.

F-actin polymerization analysis

JTAG cells transfected with empty vector mCherry (mock), TSAd-mCherry or mutated TSAd-mCherry constructs were harvested 20 h post transfection. The cells were incubated with 2 $\mu\text{g}/\text{ml}$ Hoechst 33342 in RPMI for 20 min for nuclear staining prior to harvesting. $1 \cdot 10^6$ cells were washed once in PBS and subsequently resuspended in 100 μl PBS and stimulated with anti-CD3 (OKT3, 10 $\mu\text{g}/\text{ml}$) for 30 s at 37 °C or PBS as unstimulated control. Stimulation of cells was terminated by addition of 100 μl fixation/permeabilization/staining solution consisting of 8 % paraformaldehyde, 0.25 mg/ml L- α -lysophosphatidylcholin and 1,25U Alexa Fluor® 647 phalloidin (Life Technologies). Cells were washed twice with 2 % FBS in PBS (pH 7.4), and resuspended in 50 μl PBS for analysis of phalloidin intensity on a 12-channel ISX Imaging Flow Cytometer with 40× objective (Amnis Corporation). The data was analysed using IDEAS 6.1 software (Amnis) and FlowJo.

Statistical analysis

Quantitative data are shown as mean \pm standard deviation. Statistical significance was defined as P-values < 0.05 and was estimated by a 2-tailed paired Student's *t* test.

Abbreviations

TSAd: T cell specific adaptor protein; Nck: Non-catalytic region of tyrosine kinase adaptor protein; PRR: Proline rich region; SH2: Src homology 2 domain; Lck: Lymphocyte specific protein tyrosine kinase; Itk: Interleukin 2 inducible protein tyrosine kinase; SLP-76: SH2 domain containing leukocyte protein of 76 kDa; LAT: Linker of activated T cells; APC: Antigen presenting cell; VEGF: Vascular endothelial growth factor; TCR: T cell receptor; IS: Immunological synapse; GST: Glutathione-S-transferase; FBS: Fetal bovine serum; PMA: Phorbol 12-myristate 13-acetate; IO: Ionomycin.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

CDH, VGS and SG performed the experiments, analyzed the data, performed statistical analysis and wrote the paper. LK and GA carried out the experiments and analyzed the data. LB participated in the design of the study and helped to draft the manuscript. AS conceived the study, participated in its design, analyzed the data and wrote the paper. All authors read and approved the final manuscript.

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