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Published in:
The Journal of biological chemistry

Document Version:
Publisher's PDF, also known as Version of record

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The cytoplasmic tail of L-selectin interacts with the adaptor-protein complex AP-1 subunit μ1A via a novel basic binding motif

Received for publication, November 18, 2016, and in revised form, January 30, 2017. Published, Papers in Press, February 24, 2017. DOI 10.1074/jbc.M116.768598

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Edited by Luke O’Neill

L-selectin regulates leukocyte adhesion and rolling along the endothelium. Proteins binding to the cytoplasmic tail of L-selectin regulate L-selectin functions. We used L-selectin cytoplasmic tail peptide pulldown assays combined with high sensitivity liquid chromatography/mass spectrometry to identify novel L-selectin tail-binding proteins. Incubation of the L-selectin tail with cell extracts from phorbol 12-myristate 13-acetate-stimulated Raw 264.7 macrophages resulted in the binding of μ1A of the clathrin-coated vesicle AP-1 complex. Furthermore, full-length GST-μ1A and the GST-μ1A C-terminal domain, but not the GST-μ1A N-terminal domain, bind to L-selectin tail peptide, and the intracellular pool of L-selectin colocalizes with AP-1 at the trans-Golgi network. We identified a novel basic protein motif consisting of a cluster of three dibasic residues (356RR357, 359KK360, and 362KK363) in the membrane-proximal domain of the L-selectin tail as well as a doublet of aspartic acid residues (366DD367) in the membrane-distal end of the L-selectin tail involved in μ1A binding. Stimulation of Raw 264.7 macrophages with PMA augmented the amount of μ1A associated with anti-L-selectin immunoprecipitates. However, full-length GST-μ1A did not bind to the phospho-L-selectin tail or phospho-mimetic S364D L-selectin tail. Accordingly, we propose that phosphorylation of μ1A is required for interaction with the L-selectin tail and that L-selectin tail phosphorylation may regulate this interaction in vivo. Molecular docking of the L-selectin tail to μ1A was used to identify the μ1A surface domain binding the L-selectin tail and to explain how phosphorylation of the L-selectin tail abrogates μ1A interaction. Our findings indicate that L-selectin is transported constitutively by the AP-1 complex, leading to the formation of a trans-Golgi network reserve pool and that phosphorylation of the L-selectin tail blocks AP-1-dependent retrograde transport of L-selectin.

Leukocyte trafficking is an essential mechanism of immune surveillance. The migration to and from peripheral tissues enables leukocytes to patrol the entire body, providing immune responses as needed. The exit of leukocytes from the circulation and migration into tissues is regulated at the level of cell interactions with the vascular endothelium. At least two families of adhesion receptors participate in this interaction: the integrins and the selectins (1). Selectins mediate the initial phase of leukocyte recognition of endothelium, and this takes the form of leukocyte rolling along the vessel wall (1). There are only three types of selectins: L, E, and P. L-selectin is expressed exclusively in leukocytes and acts as a major contributor of leukocyte tethering and rolling (2). Indeed, mice deficient in L-selectin display defects in lymphocytes homing to lymphoid organs and in neutrophil accumulation in inflamed tissues (3). Unlike E- and P-selectins, L-selectin is constitutively expressed on the membrane and possesses a membrane-proximal cleavage site that is hydrolyzed after leukocyte activation, leading to shedding of its extracellular domain (4). L-selectin shedding plays a key role in the regulation of leukocyte functions. It impairs the homing of activated lymphocytes to lymph nodes (5), augments neutrophil’s rolling velocity (4), controls transendothelial migration of monocytes (6), and confers lytic activity to T lymphocytes (7).

L-selectin is composed of an N-terminal lectin domain, binding endothelial ligands, followed by an epidermal growth factor-like domain, a short consensus repeat domain, a transmembrane domain, and a short cytoplasmic tail of 17 amino acids (8). The cytoplasmic tail of L-selectin is essential for L-selectin function and localization on the tips of microvilli. Indeed, L-selectin mutants lacking the membrane-distal 11 amino acid residues cannot support leukocyte rolling along endothelial venules and adhesion (9). This defect in L-selectin function cannot be explained by changes in ligand binding and is instead due to a defect in the recruitment of binding partners to L-selectin cytoplasmic tail.

This work was supported by the 7th Framework Programme for Research of the European Community through the award of an individual Intra-European Marie Curie Fellowship (to K. D.) (project 326812). This work was also supported by Deutsche Forschungsgemeinschaft (DFG) Grant Schu802/2–4 (to P. S) and grants from the British Heart Foundation (BHF) (to A. I.). The authors declare that they have no conflicts of interest with the contents of this article.

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Interaction of L-selectin with μ1A-adaptin

Figure 1. Identification of μ1A as a novel protein interacting with L-selectin cytoplasmic tail. A, peptide sequences used to identify proteins interacting with the L-selectin cytoplasmic tail. B, Raw 264.7 macrophages were stimulated for 1 min with PMA (1 μg/ml), after which cells were lysed. Lysates were incubated with synthetic desthiobiotinylated peptides (scrambled control or L-selectin tail peptides) immobilized on Dynabeads Myone streptavidin. Proteins bound to the peptides were eluted, acetone-precipitated, and digested with trypsin and Lys-C. Quantitative mass spectrometric analysis was performed on a LTQ Orbitrap mass spectrometer and analyzed using the MaxQuant Software. Peptides deriving from the bait (Sell for L-selectin) and L-selectin-binding proteins are indicated on the graph. The AP1m1 gene encodes for the μ1A subunit of the AP-1 complex.

To date, only a limited number of L-selectin tail binding partners are known. These include calmodulin (10), the ERM (ezrin-radixin-moesin) family of proteins (11), α-actinin (12), Lck (13), and protein kinase C (PKC) family members (14). Regions of the L-selectin tail involved in the interaction with these proteins have been mapped. For instance, calmodulin binds the unusual highly positively charged membrane-proximal N-terminal motif of dibasic residues, \(356^{RRLKKGKKS}364\). Moesin binds arginine 357 and lysine 362 of this domain (15). On the other hand, the more acidic C-terminal sequence of 11 amino acids of L-selectin cytoplasmic tail binds the cytoskeletal protein \(\alpha\)-actinin (12) and the tyrosine kinase Lck (13), whereas the binding sites for PKC family members have not yet been identified. Binding of moesin to the RRLKK motif facilitates the anchor of L-selectin to microvilli, which is critical for leukocyte adhesion and rolling (15). Calmodulin binding is thought to induce a conformational change in the extracellular lectin domain of L-selectin that renders the cleavage site resistant to proteolysis (10). Interaction of \(\alpha\)-actinin with the C-terminal region of the tail is important for leukocyte tethering and rolling but is not essential for ligand recognition (12). Finally, the recruitment of Lck to the C-terminal tyrosine residue (13) and binding of PKC family members to the L-selectin tail (14) are essential for L-selectin outside-in signaling.

Despite this gained knowledge, several aspects of L-selectin functions remain poorly understood. For example, the mechanism controlling the replenishment of L-selectin on the membrane surface and the clustering of L-selectin on the tips of microvilli are not well known. Identification of novel L-selectin cytoplasmic tail-interacting proteins may shed light on the mechanisms regulating L-selectin functions.

The murine L-selectin tail contains only one serine residue at position 364, which is conserved in human L-selectin. L-selectin is phosphorylated in response to T lymphocyte receptor cross-linking or treatment with PMA (14) (an activator of PKC) or engagement of chemokine receptors (16). It was proposed, but not proven, that such phosphorylation may be essential for the dynamic association/dissociation of proteins to the L-selectin tail relevant to the regulation of L-selectin functions. Indeed, moesin (11) and PKC family members (14) bind to the L-selectin tail in lymphocytes stimulated with PMA, but not in resting cells, whereas calmodulin binds L-selectin tail only in resting cells (10).

In this study, we aimed at identifying novel proteins interacting with L-selectin cytoplasmic tail domain by using sophisticated mass-spectrometry-based proteomics. We report the binding of the μ1A-adaptin subunit of the AP-1 complex (17–19) to the L-selectin tail and the regulation of this binding by phosphorylation of L-selectin cytoplasmic tail and μ1A. We propose that the AP-1-dependent transport of L-selectin may not be constitutive but regulated by the dynamic interaction between L-selectin tail and μ1A in specific cellular compartments.

Results

Identification of novel proteins interacting with the L-selectin cytoplasmic domain

To identify novel proteins binding to L-selectin cytoplasmic tail, we synthesized a desthiobiotinylated peptide comprising the entire 17-amino acid-long cytoplasmic tail of murine L-selectin. As a control, a desthiobiotinylated scrambled peptide of similar size was designed. A linker between the desthiobiotin group and the first amino acid of the two peptides was added (Fig. 1A). This was to ensure that the desthiobiotin group, when coupled to Dynabeads Myone streptavidin C1 beads, does not...
interfere with the binding of proteins to the N-terminal membrane-proximal residues of L-selectin tail peptides.

To perform peptide pulldown assays, Raw 264.7 macrophages were stimulated with PMA (1 min, 1 µg/ml) and lysed in mammalian protein extraction reagent (MPER)² buffer. Cleared lysates were incubated with either the L-selectin tail peptide (L-sel peptide) or a scrambled peptide (sc peptide) pre-coupled to Dynabeads MyOne streptavidin C1 beads. Cell lysis and experimental conditions (incubation time, washing conditions) were optimized to allow identification of novel L-selectin tail-interacting proteins. Our liquid chromatography (LC)/mass spectrometry (MS) analysis was validated by the identification of L-selectin peptides (Sell in Fig. 1), which derived from the bait used. We identified four novel proteins interacting specifically with L-selectin tail peptide. These include μ1A-adaptin (AP1m1 of the adaptor-protein complex 1 AP-1, ID: 11767), Cfh (complement component factor h, ID: 12628), Grn (granulin, ID: 14824), and Lrrc48 (leucine-rich repeat containing 48, ID: 74665) (Fig. 1). The highest MS signal intensity associated with L-selectin tail peptide was obtained for μ1A; therefore, we decided to analyze this interaction in more detail.

μ1A is one of the four adaptin subunits of the adaptor-protein complex 1 (AP-1). AP-1 controls the sorting of transmembrane proteins into clathrin-coated transport vesicles, mediating transport between the trans-Golgi network (TGN) and endosomes (17, 18, 20). This way, this vesicular carrier regulates the transport of cargo proteins (including receptors) to the plasma membrane (21).

The C-terminal domain of μ1A interacts with L-selectin

To confirm the μ1A/L-selectin tail interaction, we investigated the binding of full-length μ1A as well as of μ1A sub-domains, expressed in E. coli as GST-fusion proteins, to L-selectin tail peptide. μ1A consists of a N-terminal domain (aa 1–153), positioned close to the membrane, and a C-terminal domain (aa 154–423), which binds the canonical YXXO sorting motif (where Ø is a bulky hydrophobic residue) (17, 18, 22, 23). Expression of cDNAs encoding for full-length μ1A (GST-FL-μ1A), the C- (GST-C-μ1A) or the N-terminal domain (GST-N-μ1A) of μ1A in E. coli resulted in the synthesis of GST-fusion proteins with predicted molecular weights as shown on the polyacrylamide gel stained with Coomassie Blue (Fig. 2A).

We next incubated the L-sel peptide, or the sc peptide with purified GST-FL-μ1A. Thereafter, the beads were collected, washed, and resuspended in Laemmli buffer. Proteins were resolved on 10% SDS-PAGE, transferred onto a PVDF membrane, and subjected to Western blotting analysis using an anti-GST Ab (Fig. 2, B and C) or an anti-μ1A Ab (Fig. 2D). As shown in Fig. 2B, GST-FL-μ1A binds to the L-sel peptide but not to the sc peptide. GST (added in excess) did not bind to the L-sel peptide, confirming binding specificity. In addition, we found that GST-C-μ1A, but not GST-N-μ1A, binds the L-sel peptide (Fig. 2, C and D). Thus, the C-terminal domain of μ1A binds L-selectin cytoplasmic tail, as it has been demonstrated for proteins with canonical YXXO sorting motifs, whereas, as expected, the N-terminal domain, which binds the β1 subunit of the AP-1 complex (23), did not bind to the L-selectin tail.

μ1A is associated with anti-L-selectin receptor immunoprecipitates

We next sought to characterize the interaction between μ1A, as part of the AP-1 complex, and L-selectin tail in cells in culture. To this end, Raw 264.7 macrophages were stimulated or not for different time periods with PMA (1 µg/ml), before being lysed. Cell lysates were incubated with either an anti-L-selectin Ab or a control isotype-matched IgG2a. μ1A (AP-1) association with anti-L-selectin immunoprecipitates was investigated by Western blotting analysis using an anti-μ1A Ab.

We found little association between L-selectin and μ1A (AP-1) at steady state in resting cells. However, we observed a time-dependent increase in the amount of μ1A (AP-1) associated with anti-L-selectin immunoprecipitates upon stimulation of the cells with PMA (1 µg/ml) (Fig. 3A, top). As a control, we showed that μ1A (AP-1) is not associated with IgG2a immunoprecipitates (Fig. 3B, top). We also verified that anti-L-selectin Abs precipitated L-selectin (Fig. 3, A and B, bottom panels), as evidenced by the detection of proteins with molecular masses of 75–120 kDa representing the different glycosylated forms of L-selectin (24). However, due to its low level of expression in Raw 264.7 macrophages, the presence of L-selectin in crude cell lysates could not be detected under these experimental conditions.

L-selectin and AP-1 in vivo colocalization

To test for in vivo L-selectin and AP-1 colocalization by confocal microscopy, we used an established THP-1 cell line expressing wild type, full-length L-selectin green fluorescent protein (GFP) (6). Endogenous AP-1 was labeled with an antibody directed against the γ1 adaptin AP-1 subunit (20). This antibody recognizes the C-terminal flexible region of γ1, which is even accessible when AP-1 is part of a membrane coat. Actin staining (red) served as control. Images show two pools of L-selectin (green), one at the plasma membrane and one in intracellular vesicles (Fig. 4, A and B). The intracellular pool is clustered around large membrane domains that have AP-1 bound (blue) and therefore represent TGN membranes.

Mander’s colocalization coefficient was measured to prove L-selectin/μ1A (AP-1) colocalization in intracellular vesicles (overlap coefficient between blue and green signals). We compared the colocalization coefficient measured in an area including intracellular vesicles with the colocalization coefficient measured in an area containing the plasma membrane (Fig. 4B). Because there is no AP-1 at the plasma membrane, the colocalization coefficient measured in this area is not a real colocalization signal and is used as a background reference.

We found that the colocalization coefficients were 0.80 ± 0.09 and 0.15 ± 0.01 for intracellular vesicles and plasma membrane areas, respectively (Fig. 4C) (p < 0.0001). This result

²The abbreviations used are: MPER, mammalian protein extraction reagent; Ab, antibody; CCV, clathrin-coated vesicles; EE, early endosomes; PMA, phorbol 12-myristate 13-acetate; TBS, Tris-buffered saline; TGN, trans-Golgi network; MPR, mannose 6-phosphate receptor(s); Ahx, hexanoic acid; TRITC, tetramethylrhodamine isothiocyanate.
demonstrates that L-selectin and AP-1 (μ1A) colocalize in intracellular vesicles. The limited L-selectin/AP-1 colocalization is in line with the transient AP-1/cargo protein interactions during protein sorting and transport via clathrin-coated vesicles (CCV). Thus, TGN-associated L-selectin may serve as a reserve pool. These proteins will be transported to the plasma membrane when this pool needs to be replenished. One function of AP-1 is the sorting of mannose 6-phosphate receptors (MPR), which facilitate the transport of soluble lysosomal enzymes from the TGN to endosomes. AP-1-dependent MPR sorting ensures that 50% of the cellular MPR pool resides in the TGN (awaiting newly synthesized lysosomal enzymes), whereas only 10% of the cellular pool is at the plasma membrane due to their highly efficient AP-2-mediated clathrin-dependent endocytosis (20, 21). Thus, AP-1-dependent L-selectin sorting might be responsible for the formation of a localized L-selectin reserve pool in the TGN.

Phosphorylation of Ser-364 of L-selectin tail peptide prevents binding of μ1A

To investigate whether phosphorylation of serine 364 of the L-selectin tail affects μ1A binding, we synthesized an L-selectin tail peptide, in which serine 364 was replaced by a phosphoserine (L-sel-p-peptide). We found that GST-FL-μ1A binds L-selectin cytoplasmic tail peptide in vitro. A, BL21 bacteria were transformed with pGEX-5X3 plasmids encoding for the different GST forms of μ1A proteins, including full-length μ1A (GST-FL-μ1A) or the N-terminal part (GST-N-μ1A) or C-terminal part (GST-C-μ1A) of μ1A. After induction with isopropyl-β-D-thiogalactopyranoside, bacteria were collected and lysed, and lysates were incubated with glutathione-Sepharose beads. After 1 h, the beads were collected and washed, and 2× Laemmli buffer containing 1 mM DTT was added. Samples were run on 10% SDS-PAGE, and proteins were stained with Coomassie Brilliant Blue. The positions of GST-FL-μ1A, GST-N-μ1A, GST-C-μ1A, and GST are indicated on the left by arrows. B, purified GST-FL-μ1A or GST were incubated with a scrambled peptide (sc peptide) or L-selectin cytoplasmic tail peptide (L-sel tail peptide) coupled to streptavidin-coated beads. After 1 h, the collected beads were washed and resuspended in Laemmli buffer. The proteins were separated on 10% SDS-PAGE and transferred onto a PVDF membrane, which was subjected to Western blotting analysis using an anti-GST Ab (top). The positions of GST-FL-μ1A and GST are indicated on the left by arrows. Loading controls (which correspond to the amount of GST-FL-μ1A or GST used for the peptide pulldown assay) are shown below. Note that an excess of GST versus GST-FL-μ1A has been used. C, binding of GST-N-μ1A or GST-C-μ1A to a scrambled peptide (sc peptide) or L-selectin tail peptide (L-sel tail peptide) was carried out as described in B. D, binding of GST-FL-μ1A or GST-C-μ1A to a scrambled peptide (sc peptide) or L-selectin tail peptide (L-sel tail peptide) was carried out as described in B and C with the exception that Western blotting analysis was carried out with an anti-μ1A Ab.
These peptides were incubated with GST-FL-μ1A or GST-C-μ1A. In one experiment, residues Asp-369 and Asp-370 of the L-selectin tail were mutated to alanine (L-sel S364A). The C-terminal, negatively charged aspartic acid residues are involved in the interaction with L-selectin. We also investigated whether the C-terminal, negatively charged aspartic acid residues are involved in the interaction with L-selectin.

We found that replacing a single arginine or a single lysine residue by asparagine augmented the binding of GST-C-μ1A to these L-selectin mutated tails (Fig. 6D). Thus, the highly positively charged membrane-proximal region of the L-selectin tail is essential for μ1A binding, whereas its membrane-distal part, with its doublet of negatively charged amino acids, weakens the interaction with μ1A.

**Clusters of positively charged amino acids are essential for μ1A interaction with the L-selectin tail**

The membrane-proximal domain of the L-selectin tail contains an unusual cluster of three dibasic (positively charged) amino acids, 356RR357, 359KK360, and 362KK363. To investigate whether this RRLKKGGK sequence of the L-selectin tail is involved in the interaction with μ1A, we designed peptides in which one arginine or one lysine residue within any of the three dibasic motifs was exchanged by an alanine residue (Fig. 5C). These peptides were incubated with GST-FL-μ1A or GST-C-μ1A, and an interaction between the proteins and peptides was analyzed as described above.

We found that replacing a single arginine or a single lysine residue within any of the three dibasic clusters was sufficient to prevent GST-FL-μ1A (Fig. 6B, left) or GST-C-μ1A (Fig. 6B, right) from interacting with the L-selectin tail. We also investigated whether the C-terminal, negatively charged aspartic acid residues Asp-369 and Asp-370 of the L-selectin tail have a function in μ1A binding. We replaced separately each aspartic acid residue by an asparagine residue (D369N/D370N), or both were substituted with asparagine residues (D369N/D370N) (Fig. 7C). We found that replacement of one or two aspartic acid residues by asparagine augmented the binding of GST-C-μ1A to these L-selectin mutated tails (Fig. 6D). Thus, the highly positively charged membrane-proximal region of the L-selectin tail is essential for μ1A binding, whereas its membrane-distal part, with its doublet of negatively charged amino acids, weakens the interaction with μ1A.

**Molecular docking of the L-selectin tail to μ1A**

To identify the μ1A surface domain binding L-selectin cytoplasmic tail and to interpret the results of μ1A binding to the L-selectin tail with substituted amino acids, we explored docking of the L-selectin tail to the crystal structure of μ1A, as determined in the tetrameric AP-1 complex. To do this, we took into consideration the fact that the L-selectin tail binding surface domain of μ1A has to be accessible and not sterically blocked by the interactions of μ1A with the other subunits of the AP-1 complex.
complex. Fig. 7A shows the favorable docking pose, which explains the binding specificities of μ1A to L-selectin tail peptides. The L-selectin tail/μ1A interaction is predicted to have a strong electrostatic nature generated by two complementary charged surface areas at the binding interface. The positively charged cluster of arginine and lysine residues of the N-terminal region of the L-selectin tail are matched by several negatively charged residues, namely the aspartate and glutamate residues Glu-311, Glu-337, Glu-340, and Asp-417 of μ1A. In addition, the positively charged residues of the L-selectin tail form hydrogen bonds with residues Asn-318, Asn-338, and Gln-419 of μ1A. The interactions cited above are disrupted when the positively charged residues of the RRLKKGKK motif of the L-selectin tail are substituted to alanine, and this explains the loss of μ1A binding. Ser-364 of the L-selectin tail does not form a specific interaction with μ1A, in line with the unaltered binding of the S364A tail to μ1A, but it is in close proximity to the negatively charged residues Glu-311 and Glu-340 of μ1A. Therefore, introducing a negatively charged group at position 364 of the L-selectin tail form polar interactions with Lys-199 and Arg-410 of μ1A, thereby enhancing the binding of the two proteins. This explains the augmented binding of μ1A to the L-selectin tail peptides in which the aspartic acid residues 369 and 370 were replaced by asparagine residues. Our predicted binding mode is in good agreement with our biochemical binding experiments.

Since μ1A (AP-1) and μ2 (AP-2) are highly homologous and because both bind YXXØ-based sorting motifs and have overlapping cargo specificities, we also modeled the L-selectin tail onto the C-terminal μ2 domain to examine whether a binding mode could permit an interaction, although μ2 was not identified in our L-selectin tail pulldown experiments (Fig. 1). Docking predicts an interaction energy of 150–275 kcal/mol for L-selectin/μ2 binding poses, contrasting with the 400–500 kcal/mol for L-selectin/μ1A binding poses. Thus interactions of the L-selectin tail with μ2 are much less favorable, confirming the pulldown experiment. It is likely that L-selectin is not endocytosed by the AP-1-homologous plasma membrane AP-2 complex.
Interaction of L-selectin with \( \mu 1A \)-adaptin

L-selectin is present as an intracellular pool, which colocalizes with AP-1, and 4) the C-terminal \( \mu 1A \) domain binds selectively the non-phosphorylated L-selectin tail.

\( \mu 1A \) is one of four adaptin subunits of the AP-1 complex. AP-1 selects the cargo proteins for CCV, and it coordinates CCV formation as well as CCV uncoating, a prerequisite for vesicle-organelle membrane fusion. Besides \( \mu 1A \), the AP-1 complex contains the adaptin subunits \( \gamma \), \( \beta 1 \), and \( \sigma 1A \) (18, 19). To incorporate cargo proteins into transport vesicles, \( \mu 1A \) binds to specific sequences (called sorting motifs or cargo-binding domains) within the cytoplasmic domains of such proteins. \( \mu 1A \) is composed of two domains. The N-terminal domain (amino acids 1–153) binds the \( \beta 1 \) subunit of the AP-1 complex and thus contributes to the assembly of the AP-1 complex. The C-terminal \( \mu 1A \) domain (amino acids 153–422) binds sorting motifs of cargo proteins incorporated into CCV (23). In keeping with these structure/function studies of the AP-1 complex, we discovered that GST-C-\( \mu 1A \) but not GST-N-\( \mu 1A \) binds to the L-selectin tail.

Two canonical sorting motifs in cytosolic tails of transmembrane proteins have been described for AP-1: the tyrosine-based motif YXX\( \Phi \), which is bound by the \( \mu 1A \) C-terminal domain, and the dileucine-based motifs (e.g., (D/E)XXL(I/L)), which are bound by \( \sigma 1A \) and/or \( \sigma 1B \) adaptins (25, 26). L-selectin cytoplasmic tail does not contain either of these two sorting motifs. This means that other sequences must be involved in the interaction with \( \mu 1A \).

In addition to these two canonical motifs, non-canonical binding motifs have been identified that interact with AP-1 as well. For example, yeast AP-1 binds a 12-residue domain of the pheromone receptor Ste13p, which contains a tribasic sequence, KRK (27). In addition, AP-1 binds to the highly basic RARHRRNVDGR sequence of the polymeric immunoglobulin receptor (28). Based on these examples, the three dibasic clusters of amino acids in L-selectin tail most likely represent a novel, basic, non-canonical binding motif sequence for \( \mu 1A \).

To investigate this possibility, we designed L-selectin tail peptides in which one arginine or lysine residue within any of the three dibasic motifs was exchanged by an alanine residue (R357A, K360A, or K363A). We showed that GST-FL-\( \mu 1A \) or GST-C-\( \mu 1A \) did not bind R357A, K360A, or K363A L-selectin tail peptide. This experiment demonstrated that each dibasic residue within the N-terminal membrane-proximal region of the L-selectin tail is essential for binding \( \mu 1A \). To understand the role of residues Arg-357, Lys-360, and Lys-363 for \( \mu 1A \) interaction, we used molecular docking to predict the predominant binding mode of the L-selectin tail with the 3D structure of \( \mu 1A \). We found that the positively charged cluster of arginine and lysine residues of the N-terminal region of the L-selectin tail is bound by the \( \mu 1A \) domain. The match of this binding mode with the N-terminal region of the L-selectin tail is consistent with the observations that \( \mu 1A \) binding is mediated by arginine and lysine residues within the N-terminal region.
is influenced by the membrane bilayer. Thus, in a phospholipid bilayer (which mimics the inner leaflet of the plasma membrane), the positively charged N-terminal part of the L-selectin tail interacts with anionic phosphatidylserine lipids at the membrane interface through electrostatic interactions. This prevents calmodulin to interact with the N-terminal part of the L-selectin tail (29) and promotes L-selectin shedding (15). Disruption of the interaction between the positively charged N-terminal part of the L-selectin tail with the phospholipid bilayer is a prerequisite for the binding of calmodulin to the L-selectin tail. Such disruption is brought about by the interaction of the FERM domain of moesin with L-selectin to induce a separation of the cationic N-terminal part of the L-selectin tail from the phospholipids (30). We do not know whether the phospholipid bilayer plays a similar role in intact cells for at least two reasons. First, the lipid bilayer is itself a complex mixture of lipids. Its structure in the plane of the bilayer is that of a microscopic mosaic of regions that differ in composition (i.e. domains or clusters) (31). In particular, receptors are found in domains, such as the caveola, that are rich in cholesterol. Second, the cited authors (29, 30) have not taken into account in their model that calcium interacts with phospholipid bilayers. Such interactions induce changes in lipid dynamics, structure, and affinity of phospholipid-binding proteins (32).

AP complexes are targeted to their respective membrane via binding to specific phosphoinositides, AP-1 to phosphatidylinositol 4-phosphate and AP-2 to phosphatidylinositol 4,5-bisphosphate. There are no data in the literature describing inhibition of a sorting motif/AP complex binding by basic residues located in the vicinity of the sorting motif (23, 25, 26). In fact, phosphatidylinositol 4-phosphate incorporated into liposomes binds AP-1, and the interaction between adaptins and sorting motif-containing proteins is further stabilized by phospholipids during the polymerization of the vesicle coat (33). These results indicate a role of phospholipids and phosphoinositides in the interaction of AP-1 with its binding partners and vesicular assembly.

The μ2 adaptin of the plasma membrane AP-2 complex collects proteins for clathrin-mediated endocytosis, and many proteins are transported by AP-1 as well as AP-2 CCV (17, 18). μ2 was not isolated in our screen, despite the fact that it is highly homologous to μ1A. Therefore, we asked why μ2 could
not bind L-selectin tail. This is probably explained by the fact that the key amino acids in μ1A involved in L-selectin tail binding are not found in μ2. Indeed, only the first glutamic acid residue of the L-selectin binding motif in μ1A is conserved in μ2 (Glu-311 > Glu-321, Glu-337 > Ser-347, Glu-340 > Ala-350, Asp-417 > Ile-430). Also, the asparagine residues 318 and 338 and glutamine 419 of μ1A are not conserved in μ2, where they are replaced by leucine and glutamic acid residues. These data strongly indicate that the AP-2 complex, which is highly homologous to the AP-1 complex, does not play a role in the regulation of L-selectin cell surface expression.

We tested for L-selectin and AP-1 colocalization in vivo by confocal microscopy. Besides the plasma membrane pool of L-selectin, we detected an intracellular pool of L-selectin localized next to AP-1 decorated domains of the TGN. This indicated that AP-1-dependent L-selectin sorting might be responsible for the formation of this putative L-selectin reserve pool.

Murine L-selectin tail contains one unique serine residue at position 364, which is conserved in human L-selectin. It was shown that upon stimulation of lymphoblastoid cell lines with PMA or chemoattractants, the L-selectin tail becomes rapidly phosphorylated (16), and this is probably caused by recruitment of PKC family members to the L-selectin tail (14). However, little is known of the relationship between L-selectin tail phosphorylation and its interaction with binding partners in vivo. We found low amounts of μ1A associated with anti-L-selectin immunoprecipitates in resting Raw 264.7 macrophages. However, this association increased when the cells were stimulated with PMA. This result could be explained in at least two ways, involving phosphorylation of μ1A and/or L-selectin tail. One phosphorylation step would be the enhanced phosphorylation of μ1A in response to PMA, because such phosphorylation would augment its interaction with the L-selectin tail. Indeed, the cytoplasmic AP-1 complex is present in a closed conformation, in which the cargo binding domains of μ1A and σ1A are blocked by γ1 and β1 adaptins. The complex has to undergo a conformational change to an “open” state. This causes the release of the μ1A C-terminal domain from the large adaptins, enabling its movement toward the membrane and the cytoplasmic sorting domains of membrane proteins. The open conformation is favored by threonine phosphorylation of μ1A in the flexible linker region connecting the N- and C-terminal domains (34–36). Such a mechanism of activation has also been described for the L-selectin tail-interacting protein moesin. Phosphorylation of moesin by PKCα on a threonine residue located in the C-terminal part unfolds the protein, thus allowing it to interact with the L-selectin tail (37).

The second phosphorylation reaction to explain a PKC-stimulated association between μ1A and L-selectin tail would be the L-selectin tail phosphorylation. However, μ1A does not bind the phospho-L-selectin tail peptide, and this is due to the fact that the phosphate group on serine 364 is in close proximity to the negatively charged residues Glu-311 and Glu-340 of μ1A. In keeping with our results, other investigators showed inhibition of calmodulin binding to L-selectin-p-tail in vitro and in intact THP-1 cells (6).

The AP-1 coat is found on the TGN and on early endosomes (EE), and it mediates constitutive protein transport from the EE to the TGN as well as from the TGN to EE, as exemplified by the mannose 6-phosphate receptors, furin and sortilin (20, 21, 26). Interestingly, besides its role as cargo binding and constitutive regulator of protein transport, the role of AP-1 in regulated secretory pathways has also been shown (38–40). Our data showing that phosphorylation of both μ1A and the L-selectin tail controls the interaction between these two proteins indicate a regulated, rather than a constitutive, AP-1-dependent secreted pathway for L-selectin. We developed a model of AP-1-dependent L-selectin sorting based on our data and the known functions of AP-1 (Fig. 7B). We propose that stimulation of μ1A phosphorylation activates AP-1 and enhances AP-1-dependent protein transport from the TGN to EE. Phosphorylation of the L-selectin tail in activated leukocytes would prevent its binding to μ1A (AP-1) in EE, leading to less incorporation of L-selectin into AP-1 CCV for retrograde EE to TGN transport. Consequently, L-selectin would transiently accumulate in EE, from which it can be readily transported to the plasma membrane through exocytosis (21). Such an indirect function of AP-1 in the regulation of plasma membrane protein transport has been demonstrated by us for both mannose 6-phosphate receptors in mouse fibroblasts (20, 21). Thus large amounts of L-selectin would be exported to the plasma membrane and could be readily clustered on microvilli relevant for leukocyte rolling and adhesion. We are currently investigating this working hypothesis of stimulation-dependent compartment-specific L-selectin tail and μ1A phosphorylation in leukocytes.

In summary, we have made several novel and important observations regarding L-selectin functions. First, we identified μ1A as a novel protein interacting with the L-selectin tail. Second, we demonstrated colocalization of an intracellular pool of L-selectin with AP-1. Third, a novel binding motif formed by a triplet of dibasic amino acid motifs in L-selectin (RRXXKKXXK) mediates this interaction. Fourth, L-selectin tail phosphorylation inhibits μ1A binding. From these data we can conclude that AP-1, but not AP-2, is involved in the transport pathway regulating L-selectin secretion to the plasma membrane upon stimulation.

**Experimental procedures**

**Antibodies**

The antibodies and their sources were as follows. Anti-L-selectin Ab lam 1–116 (sc-13505) was purchased from Santa Cruz Biotechnology (Heidelberg, Germany); the mouse IgG2a isotype control Ab (UPC-10), was purchased from Sigma-Aldrich (Munich, Germany); the anti-μ1A Ab (ab 170277) was bought from Abcam (Cambridge, UK), the anti-GST mouse monoclonal Ab (catalog no. 71097-3) was from Novagen/Merck (Darmstadt, Germany).

**Chemicals**

Protease inhibitor tablets were from Roche Applied Science (Penzberg, Germany), and phosphatase inhibitor mixtures 1 and 2 were purchased from Sigma-Aldrich. Dynabeads Myone streptavidin C1 and MPER buffer were from Invitrogen/Thermo Fisher Scientific (Karlsruhe, Germany), and Protein A-Sepharose TM CL-4B and streptavidin-Sepharose beads.
were purchased from GE Healthcare Life Sciences (Munich, Germany). RPMI 1600 medium was purchased from Gibco/Thermo Fisher Scientific. Fetal bovine serum and octyl-β-d-glucoside were from Sigma-Aldrich. TAPI-O was from Peptides International (Louisville, KY).

**Peptides**

Automated solid phase synthesis of peptides using Fmoc (N-(9-fluorenyl)methoxycarbonyl)/t-Bu chemistry was carried out at the Biochemistry core facility of the Max Planck Institute for Biochemistry ( Martinsried, Germany). Characterization and quality control (purity) was done by RP-HPLC. All peptides were desthiobiotinylated, and a spacer (hexanoic acid (Ahx)) between the desthiobiotin group and the first amino acid of the peptides was introduced. The following peptides were used asusin this study: desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin D370N tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin D369N tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin K363A tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin K360A tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin R357A tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin K360A tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin K363A tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin D369N tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin D370N tail); desthiobiotin-GAhxRRLKGKKSQERMDDPY-OH (L-selectin D369N, D370N tail).

**Cell culture**

Raw 264.7 macrophages were grown in 75-cm² flasks at 37 °C in an atmosphere of 5% CO₂ in RPMI 1600 medium supplemented with 10% FCS, antibiotics, and L-glutamine (20 mM).

**Immunoprecipitation and Western blotting**

Raw 264.7 macrophages were pretreated with the sheddase inhibitor TAPI-O (10 min, 10 μg/ml) and then incubated in the absence or presence of PMA (1 μg/ml) for different time periods. Thereafter, cells were collected by centrifugation (200×g, 2 min) and washed with PBS. The tubes were put on ice for 5 min, after which 1 ml of lysis buffer was added. The lysis buffer consisted of 50 mM Tris-HCl, pH 7.4, 30 mM octyl-β-glucoside, 1 mM EDTA, 110 mM NaCl, protease and phosphatase inhibitors. Cell lysates were clarified by centrifugation, and the supernatants were subjected to immunoprecipitation. This was performed by exposure to the anti-L-selectin Ab (1 μg/ml) or the isotype-matched IgG2a Ab (1 μg/ml) for 1 h and then to 40 μl of a 50% slurry of protein A-Sepharose for 1 h. The beads were subsequently collected by centrifugation (200×g, 1 min) and washed three times with lysis buffer. The beads were then resuspended in 2× concentrated Laemmli sample buffer and boiled under reducing conditions (1 mM DTT) for 5 min. The immunoprecipitated proteins were subjected to electrophoresis on 10% SDS-PAGE and transferred to polyscreen PVDF transfer membranes. The proteins were subjected to electrophoresis on 10% SDS-PAGE and transferred to polyscreen PVDF transfer membranes, and Western blotting analysis was carried out as described above using a rabbit anti-μ1A Ab or an anti-GST Ab.

L-selectin peptides coupled to streptavidin-Sepharose beads were also incubated with lysate extracts of Raw 264.7 macrophages. Briefly, the cells were lysed in buffer composed of 50 mM Tris-HCl, pH 7.4, 0.5% CHAPS, 110 mM NaCl, 50 ng of purified μ1A expressed as GST-fusion proteins was added. The Eppendorf tubes were rotated for 1 h in the cold room, after which pellets were collected by centrifugation (200×g, 1 min). The pellets were washed three times with washing buffer (50 mM Tris-HCl, pH 7.4, 0.5% CHAPS, 300 mM NaCl). The beads were subsequently resuspended in 2× concentrated Laemmli sample buffer and boiled under reducing conditions for 5 min. The proteins were subjected to electrophoresis on 10% SDS-PAGE and transferred to polycrystalline PVDF transfer membranes, and Western blotting analysis was carried out as described above using a rabbit anti-μ1A Ab or an anti-GST Ab.

**Proteomics analysis**

Raw 264.7 macrophages were stimulated for 1 min with PMA (1 μg/ml), after which the cells were pelleted and washed with ice-cold PBS. The cells were subsequently lysed with Miper buffer. Clarified lysates were then incubated with Dynabeads Myone streptavidin C1 that had been coupled to either L-selectin tail peptide or scrambled peptide (see above). After a 2-h incubation in the cold room under rotation, the beads were collected with a magnet and washed gently three times with Miper buffer. The beads were then resuspended in 50 mM Tris-HCl, pH 8.0, containing 10 mM biotin to dissociate the biotin analogue desthiobiotin from streptavidin beads. After 1 h in the cold room, supernatants were collected, and proteins were precipitated overnight at -20 °C with 100% acetone. Proteins were pelleted by centrifugation (15,000×g, 15 min) in a cold centrifuge. Subsequently, the pellet was air-dried under a fume hood.
20 μl of 6 M guanidium HCl containing a 10 mM concentration of the reducing agent tris(2-carboxyethyl)phosphine and 40 mM of the alklylation agent 2-chloroacetamide were added to each pellet, and the samples were boiled for 10 min. 180 μl of LT-Digestion buffer (25 mM Tris-HCl, pH 8.5, containing 10% acetonitrile) was added, followed by 1 μg of each trypsin and Lys-C endopeptidase. The tubes were left overnight at 37 °C under shaking. Thereafter, 8 μl of 25% TFA (1% final) was added to stop the reaction. The peptides were bound to triple filter surface (catalog no. SDB-RPS 3X), and after washing the filters three times with 0.2% TFA, proteins were eluted with 60 μl of buffer containing 80% acetonitrile and 5% ammonium. Peptides were analyzed on an LTQ-Orbitrap analyzer, identified, and quantified using the MAXQUant software (41).

**Purification of GST-fusion proteins**

Expression of μ1A as GST-fusion proteins has been described in detail (42). Briefly, pGEX-5X3 plasmids were transformed into E. coli BL21. 1 ml isopropyl β-D-thiogalactopyranoside was added overnight to bacterial cultures grown at 26 °C. Bacteria were harvested, put on ice, washed once with ice-cold PBS, and lysed by the addition of a buffer made up of 26 °C. Bacteria were harvested, put on ice, washed once with ice-cold PBS, and lysed by the addition of a buffer made up of 50 mM Tris-HCl, pH 7.4, 1% Triton X-100, 110 mM NaCl, and protease inhibitors. This was followed by sonication. The lysates were clarified by centrifugation (10,000 × g, 15 min), supernatants were collected, and 10% glycerol was added. These supernatants were incubated in the cold room with glutathione-Sepharose pre-equilibrated in PBS buffer. After 1 h, the beads were collected by centrifugation (200 × g, 1 min) and washed three times with ice-cold PBS. The GST-fusion proteins were eluted by adding 200 μl of buffer containing 50 mM Tris-HCl, pH 8.0, and 10 mM reduced glutathione. Eluted solutions were passed through a G-25 Sephadex column, which was subjected to centrifugation (500 × g, 1 min), and GST-fusion proteins free of glutathione were collected. Protein concentrations were estimated by spectrometry.

**Immunofluorescence confocal microscopy**

THP-1 cells stably expressing wild type L-selectin tagged to green fluorescent protein (WT L-selectin-GFP) were generated and cultured as described previously (6). Cultured THP-1 cells were harvested and resuspended in warmed undiluted RPMI at a density of 0.5 × 10⁶ cells/ml. Cells were subsequently seeded onto poly-L-lysine-coated coverslips and allowed to settle for 10 min at 37 °C. RPMI was aspirated, and coverslip-bound cells were fixed in 3% paraformaldehyde for 20 min at room temperature. Cells were washed three times in PBS and then permeabilized in 0.6% saponin for 10 min at room temperature. After washing fixed cells with PBS, cells were blocked in 5% fetal calf serum containing Fc receptor block (Miltenyi Biotec Ltd.) for 1 h at room temperature before staining with 1:300 (v/v) of 0.1 mg/ml TRITC-phalloidin and anti-γ-1 AP-1 mouse monoclonal antibody (250 μg/ml; BD Biosciences) diluted at 1:75 (v/v) overnight at 4 °C. Cells were washed free from excess phalloidin and antibody, blocked as before, and then stained with secondary goat anti-mouse (Life Technologies) Alexa Fluor 633 (1:400 dilution). Specimens were mounted onto glass slides using mounting medium (Dako). Images were acquired using Leica SP5 confocal microscope, using a ×63 objective lens. Scale bar, 7.5 μm.

**Molecular modeling**

Molecular docking of the L-selectin tail to μ1A was conducted using the HEX version 8.0 protein-protein docking program (43). The crystal structure of μ1A (Protein Data Bank code 4P6Z) and the structural model of 17-residue long L-selectin tail was modeled using the crystal structures of the 8-residue-long L-selectin tail (Protein Data Bank code 2LGF). The remaining 8 amino acid residues of the L-selectin tail were modeled using the loop prediction option of the Prime module within Schrödinger software (44). The L-selectin tail in a wild type as well as mutated forms (S364pS, S364D, D369N, and D370N) were used for docking to facilitate a choice of docking solutions. The μ1A-L-selectin binding surface interactions were mapped based on shape and electrostatics complementarities as implemented in HEX version 8.0. Default settings for other docking parameters were used. HEX version 8.0 has generated over 100 solutions for each docking run. Besides the HEX energy function, the obtained docking pose of the L-selectin tail in the wild type and mutated forms was evaluated based on its ability to explain the mutagenesis data. In addition, only docking poses on the surface of μ1A, which is not in an interaction with other subunits within, AP-1 were considered.

**Author contributions**—K.D. designed the project, performed the research, analyzed the data, and wrote the manuscript; I. G. T. performed the modeling of μ1A-L-selectin tail interaction, analyzed the data, and wrote part of the manuscript; A. I. performed the colocalization studies; P. S. provided scientific expertise, analyzed the data, and wrote part of the manuscript. All authors read and approved the final manuscript.

**Acknowledgments**—We thank Yosuke Yoneyama and Shin-Ichiro Takahashi (University of Tokyo) for providing the pGEX-5X3 plasmids encoding for full-length μ1A or the N-terminal or C-terminal region of μ1A. Ralph Böttcher, Markus Moser, and Reinhard Fässler (Max Planck Institute for Biochemistry, Martinsried, Germany) are acknowledged for fruitful discussions. Herbert Schiller (Max Planck Institute for Biochemistry, Martinsried, Germany) is acknowledged for help in the identification and quantification of L-selectin tail-interacting proteins using the MAXQUant software. We thank Stephan Uehbel (core facility of the Max Planck Institute for Biochemistry, Martinsried, Germany) for the synthesis of L-selectin tail peptides.

**References**

Interaction of L-selectin with μ1A-adaptin


The cytoplasmic tail of L-selectin interacts with the adaptor-protein complex AP-1 subunit μ1A via a novel basic binding motif
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doi: 10.1074/jbc.M116.768598 originally published online February 24, 2017

Access the most updated version of this article at doi: 10.1074/jbc.M116.768598

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