Primary Structure of the Leukocyte Function–associated Molecule-1 α Subunit: an Integrin with an Embedded Domain Defining a Protein Superfamily

Richard S. Larson, Angel L. Corbi, Lisa Berman, and Timothy Springer

Center for Blood Research, Department of Pathology, Harvard Medical School, Boston, Massachusetts 02115

Abstract. The leukocyte function-associated molecule 1 (LFA-1, CD11a/CD18) is a membrane glycoprotein which functions in cell-cell adhesion by heterophilic interaction with intercellular adhesion molecule 1 (ICAM-1). LFA-1 consists of an α subunit (M_r = 180,000) and a β subunit ($M_r = 95,000$). We report the molecular biology and protein sequence of the α subunit. Overlapping cDNAs containing 5,139 nucleotides were isolated using an oligonucleotide specified by tryptic peptide sequence. The mRNA of 5.5 kb is expressed in lymphoid and myeloid cells but not in a bladder carcinoma cell line. The protein has a 1,063-amino acid extracellular domain, a 29-amino acid transmembrane region, and a 53-amino acid cytoplasmic tail. The extracellular domain contains seven repeats. Repeats V-VII are in tandem and contain puta-

THE leukocyte function-associated 1 (LFA-1)¹ molecule is a member of a family of three leukocyte glycoproteins involved in cell-cell adhesion. This family of proteins, LFA-1, Mac-1, and p150,95, are heterodimers consisting of distinct α subunits ($M_r = 180,000, 170,000,$ and 150,000, respectively) and a common β subunit ($M_r =$ 95,000) (25, 47). NH₂-terminal sequencing of the α subunits in the mouse and human has suggested that they are structurally related (33, 48); however, the cell surface expression and function of these molecules differs. LFA-1 is expressed on virtually all leukocytes and is involved in a large number of adhesion-dependent phenomena. mAb directed against LFA-1 inhibit antigen-specific T helper cell function and cytolytic functions such as cytotoxic T lymphocyte-mediated killing, antibody-dependent cytoxicity by granulocytes, and natural killer activity (47). LFA-1 is also involved tive divalent cation binding sites. LFA-1 has significant homology to the members of the integrin superfamily, having 36% identity with the Mac-1 and p150,95 α subunits and 28% identity with other integrin α subunits. An insertion of ~ 200 amino acids is present in the NH₂-terminal region of LFA-1. This "inserted/ interactive" or I domain is also present in the p150,95 and Mac-1 α subunits but is absent from other integrin α subunits sequenced to date. The I domain has striking homology to three repeats in human von Willebrand factor, two repeats in chicken cartilage matrix protein, and a region of complement factor B. These structural features indicate a bipartite evolution from the integrin family and from an I domain family. These features may also correspond to relevant functional domains.

in antigen-independent interactions that mediate cell localization to sites of inflammation such as leukocyte adhesion to endothelial cells, fibroblasts, epidermal keratinocytes, and synovial cells (12–14, 17, 30, 53). Mac-1 and p150,95 are expressed on monocytes, granulocytes, and some activated lymphocytes, and function as adhesion molecules in cellcell and cell-substrate interactions as well as complement receptors for C3bi (1).

The importance of these three glycoproteins is signified by the clinical syndrome known as leukocyte adhesion deficiency (LAD) (1). The primary defect in LAD occurs in the β subunit common to LFA-1, Mac-1, and p150,95 (22) resulting in deficient surface expression of the $\alpha\beta$ complexes. LAD patients have recurrent bacterial infections which are sometimes fatal and their leukocytes are deficient in a wide range of adhesion-dependent functions.

The structure of the β subunit common to LFA-1, Mac-1, and pl50,95 has been determined (23, 27) and has revealed homology to extracellular matrix (ECM) receptors. These similarities led to the concept of a family of α/β heterodimers designated the integrins (21, 41). The term "integrin" emphasizes the role of these proteins as transmembrane links between the extracellular environment and the cytoskeleton.

^{1.} Abbreviations used in this paper: CMP, cartilage matrix protein; ECM, extracellular matrix; FNR, fibronectin receptor; ICAM-1, intracellular adhesion molecule-1; LFA, leukocyte function-associated 1; vWF, von Willebrand factor.

Three subfamilies of integrins are defined by their distinct β subunits. The β_1 subunit is common to the fibronectin receptor (FNR) and some antigens appearing very late in leukocyte activation whereas β_3 is common to platelet glycoprotein IIb/IIIa (gpIIb/IIIa) and the vitronectin receptor. β_1 and β_3 integrins are all ECM receptors and are involved in cell-substrate adhesion, matrix assembly, regulation of cell growth, differentiation, and localization during morphogenesis, and wound healing. We will refer to the β_1 and β_3 family as the extracellular matrix receptor integrins. The β_2 subunit is common to LFA-1, Mac-1, and pl50,95, whose expression is limited to leukocytes, and we designate these as the leukocyte integrins.

The ligand of LFA-1 is an inducible cell surface glycoprotein, intercellular adhesion molecule-1 (ICAM-1) (M_r = 90,000), which is found on cells of many lineages including leukocytes, endothelial cells, fibroblasts, and epidermal keratinocytes (14, 29, 40, 46, 49). ICAM-1 mRNA and surface expression is induced by inflammatory mediators including interferon-gamma, interleukin-1, tumor necrosis factor, and lipopolysaccharide (36, 46, 49); thus, it may regulate cell interaction and localization in inflammation. LFA-1-dependent adhesion of cells to planar lipid membranes containing ICAM-1 requires metabolic energy, a functional cytoskeleton, and divalent cations (29). Cell activation can enhance LFA-1-dependent adhesion without any effect on LFA-1 or ICAM-1 surface expression as shown by phorbol ester-induced homotypic adhesion of B, T, and monocytic cells (39). ICAM-1 is a member of the immunoglobulin superfamily and consists of five immunoglobulin constant region-like domains. In contrast to most ligands of the ECM receptor integrins which contain the critical recognition sequence arginine-glycine-aspartic acid (RGD), ICAM-1 does not contain an RGD sequence (46, 49). The LFA-1-ICAM-1 receptorligand pair is thus far the only known example of a member of the integrin superfamily interacting with a member of the immunoglobulin superfamily.

We have been interested in the structural basis for the important function of LFA-1 in inflammation and the immune response. Furthermore, we wished to define the relationship of LFA-1 to other leukocyte integrins and to the ECM receptor integrins. The lack of an RGD sequence in ICAM-1 raised the question of whether the LFA-1 α subunit has structural features typical of ECM integrins or whether novel features consistent with the lack of RGD recognition exist. Therefore, we have characterized the LFA-1 α subunit at the protein and mRNA level and have determined its complete sequence. The amino acid sequence demonstrates that LFA-1 is an integral membrane protein containing an extracellular domain, a single hydrophobic transmembrane domain, and a cytoplasmic tail. The extracellular domain has seven repeats; the three repeats located most COOH-terminal contain putative divalent cation binding sites. LFA-1 has significant homology with the other members of the integrin superfamily. A region near the NH₂ terminus of the molecule contains an insertion of \sim 200 amino acids similar to p150,95 (9) and Mac-1 (4, 7, 38). This domain has significant homology to the type A domains of von Willebrand factor (vWF), complement factor B, and the repeats of chicken cartilage matrix protein (CMP). These similarities suggest relevant functional domains within the LFA-1 α subunit as well as novel evolutionary relationships.

Materials and Methods

Protein Purification

The mAb, TS1/22, which is directed against the LFA-1 α subunit, was purified and coupled using cyanogen bromide to CL-4B Sepharose at 2 mg mAb per ml of packed bed. SKW3 cells (42.2 g) were lysed in 300 ml of lysis buffer (25) and the lysate was spun at 5,000 g, the pellet discarded, and then spun at 16,000 g for 2 h. The supernatant was then sequentially passed through a precolumn of activated and quenched CL-4B Sepharose and then a TS1/22 mAb Sepharose column. The TS1/22 column was washed sequentially (25), and the LFA-1 molecule was eluted with 0.5 M NaCl, 0.1% Triton X-100, 1 mM iodoacetamide, 10 U/ml aprotinin, and 0.025% NaN₃, 50 mM triethylamine, pH 11.5, and the pH immediately neutralized. The fractions containing LFA-1 were pooled, lyophilized, and precipitated in 5 vol ethanol at -20° C overnight.

Purified protein was reduced and alkylated (23) and subjected to preparative SDS-PAGE. The band corresponding to the α subunit was visualized with 1 M KCl, excised, and electroeluted (20). The purified α subunit was lyophilized and precipitated in 4 vol ethanol at -20°C overnight. The pellet was resuspended and digested with 1% (wt/wt) trypsin (23). The tryptic fragments were then isolated by HPLC (Beckman Instruments Inc., Palo Alto, CA) on a C4 reverse phase column (Vydac, Hesperia, CA). The peptides were eluted on a 0-60% acetonitrile gradient in 1% trifluoroacetic acid. Several peaks were rechromatographed isocratically in a concentration of acetonitrile determined by the equation, F = 0.9E-2, where F is the volume percentage of acetonitrile under isocratic conditions for a peptide that eluted at E percent during the linear gradient (57). Peaks were collected in 1.5-ml polypropylene tubes and concentrated to $<50 \ \mu$ l. Eight peaks were subjected to microsequencing. The sequence of one peptide, L64, was used to synthesize a single sequence oligonucleotide (5'-GGGATGTTGTGGT-CATGGATGGTGGGCTCAAT-3') according to suggestions of Lathe (26).

cDNA Cloning, Restriction Mapping, and Nucleotide Sequencing

The production and screening of the cDNA library was performed as previously described (9). Restriction maps of the selected clones were determined by double and partial digests (28). Restriction fragments were subcloned into either M13mp18 and mp19 (31) or pGEM-3Z, 4Z, or 7Z (Promega Biotec, Madison, WI). Deletions of the fragments in pGEM were made using Exonuclease III and S1 nuclease (18). Sequencing was by the dideoxy termination method (43). The coding, 5'-untranslated and 3'untranslated regions were determined 100, 100, and 36.7% in both orientations, respectively.

Southern and Northern Blot

Southern blots were performed as described elsewhere (8). For Northern blots, 10 μ g of poly (A+) RNA isolated from SKW3, U937, IB4, or EJ cells was subjected to electrophoresis on a 1.0% formaldehyde gel and transferred to nitrocellulose (5). The nitrocellulose (Bio-Rad Laboratories, Richmond, CA) was prehybridized and hybridized in 2× SSC, 1× Denhardt's solution, 0.1% SDS, and 10 μ g/ml of herring sperm DNA. A 1.8-kb Eco RI probe from the 5' end of the cDNA clone, λ 3R1 (Fig. 2), was labeled by nick translation and used as probe.

Computer Analysis

Homology searches and alignment of sequences used the Microgenie DNA program (Beckman Instruments Inc.), FASTP (56) on the NBRF and NEW databases (National Biomedical Research Foundation, Washington, DC), and FASTP using the SWISS-PROT data base (Bionet Intelligenetics, Mountainview, CA). These alignments were then optimized, and the percent identity and statistical significance were determined using ALIGN (NBRF) (11).

Hydrophobicity was determined according to Hopp and Woods using the Microgenie DNA program (19).

Results and Discussion

Protein Purification and Peptide Sequence

LFA-1 was solubilized from SKW3, a T lymphoma cell line,



Table I. Sequences of Tryptic Peptides

Residues	Amino acid sequence
95-107	X X D Q (N) T Y L S G L (E) Y L F
199-210	HMLLLTNTFGAI
254-260	Y I I G I G K
405-413	V L L F Q E X Q G
494-503	GEAITALTXI
541-554	IEGTQVLSGIQXFG
564-576	X (L) E (G) D (V/G) L A D V A V G A E
803-817	K V E M L K P H S E I X V S
929-946	<u>I (E/Q) P S I H D H N I P</u> X L E A V X G

Parentheses indicate ambiguity in the sequence. The underlined residues were used to generate an oligonucleotide probe.

(32). The insert size was determined and the longest clone, λ 5L5, was restriction mapped, and sequenced (Fig. 2). This clone contained the nucleotide sequence corresponding to the oligonucleotide probe (85% identity to the "best-guess" probe) and agreed perfectly with the tryptic peptide sequence (16 of 16 residues). However, this clone did not encode the entire protein since not all of the tryptic peptide sequences were present in the open reading frame. The 5' 1.0-kb Eco RI fragment of λ 5L5 was used to select 14 additional clones. The clone λ 3R1 had an identical restriction map in the overlapping regions and contained an additional 1.0-kb 5' fragment (Fig. 2).

The composite sequence of λ 5L5 and λ 3R1 contains 5,139 nucleotides (Fig. 3). There is an open reading frame of 3,510 nucleotides, a 5' untranslated region of 94 nucleotides, and a 3' untranslated region of 1,535 nucleotides which contains a polyadenylation signal site 15 nucleotides before the poly(A+) tail. Within the 3' untranslated region there is a typical Alul repeat consisting of two tandem related sequences each terminated by an A-rich segment (16).

Northern blots demonstrate an LFA-1 α subunit mRNA of 5.5 kb in SKW3 T lymphoma cells, U937 myelomonocytic cells, and IB4 B lymphoblastoid cells (Fig. 4 *A*, lanes *l*-3); however, no signal was detected in EJ bladder carcinoma cells (Fig. 4 *A*, lane 4). The mRNA expression is in agreement with the restriction of cell surface expression of LFA-1 to hematopoietic cells. The β subunit showed the same pattern of expression (Fig. 4 *B*). In Southern blots, the 1.2-kb 5' Eco RI-Bam HI fragment from clone λ 2L2 hybridized to two fragments of 10 and 8 kb (8). A genomic clone isolated from a cosmid library possesses the same two fragments



the sequence possessing the lowest codon redundancy was used to specify a single oligonucleotide sequence which used the most commonly occurring human codons (26).

cDNA Cloning and Characterization

The 32-mer oligonucleotide probe was used to isolate 20 clones from a size-selected λ gt10 cDNA library constructed from PMA-stimulated myeloid cells (9). These cells have been previously shown to synthesize the LFA-1 α subunit



Figure 2. Restriction map of the LFA-1 α subunit cDNA clones. The open reading frame and untranslated regions are indicated with a thick and thin line, respectively. Arrows indicating the sequencing strategy are shown. The relevant restriction sites are Bal I (*Bl*), Bam HI (*Ba*), Bgl II (*B*), Cla I (*C*), Eco RI (*R*), Nru I (*N*), Pst I (*P*), Sca I (*Sc*), Sma (*S*), and Sph I (*Sp*).

	ANG ANG GAT TOC TOC ANC ACT GTG ATG GOC ATG GOC GTG CTG CTG CTG TOC TTT TTC TTC GOC COC GOC TOC ANC CTG GAC GTG CGG GOC GOG GOG GOC GOC GOC GOC GOC GO	: 205
13	CCA CCG CCC CCC CCG AGG CAC TITT GGA THC CCC CFIC CTG CAG GTC GGA AAC GCG GTC ATC CTG GGA GCT CCA GCG GAG GCG AAC AGC ACA GGA NGC CFIC TAT CAG TAC CAG P P R A G R H F G Y R V L Q V G N G V I V G A P G E G N S T G S L Y Q C Q	316
50	TO GO CACA GOA CAC TO C CTIG OCA GTC ACC CTIG AGA GET TO C A <u>AC TAT ACC</u> TO C AAG TAC TIG GCA ATG ACC ACA GAC GOA ACG AGA AGC ATT THE GO TOT S G T G H C L P V T L R G S <u>N Y T</u> S K Y L G M T L A T D P T D G S I L A C	427
87	GAC OCT GGE CTE TET CGA AGE TET GAC CHE AAC MAT CTE AST GGE CTE TET THE CTE THE CEE MA GGE GGE GGE GGE GGE GGE GGE GGE GGE GG	538
124	CAR TGT ATC ARC GOC ARC GTA GAC CTG GTA TTT CTG TTT GAT GGT TGG ATG ARC TTG CAG GAA TTT CAG ARA ATT CTG GAC TTC ATG ARG GAT GTG ATG ARG ARG ARG ARG ARG ARG ARG ARG ARG AR	. 649
161	CTC ACC ANC ACT TOG TAC CAG TIT GCT GCT GTT CAG TIT TOC ACA AGC TAC AAA ACA GAA TIT GAT TIC TCA GAT TAT GTT AAA TOG AAG GAC CCT GAT GCT CTG CTG AAG L S N T S Y Q F A A V Q F S T S Y K T E F D F S D Y V K W K D P D A L L K	760
198	CAT GTA AAG CAC ATG TTG CTG TTG ACC AAT ACC TTT GGT GOC ATC AAT TAT GTC GOG ACA GAG GTG TTC COG GAG CTG GOG GOC COG GCA GAT GOC ACA AAA GTG CTT H V K <u>H M L L L T N T F G A I</u> N Y V A T E V F R E E L G A R P D A T K V L	871
235	ATC ATC ATC AGG GAT GGG GAG GAC ACT GAC AGT GGC AAC ATC GAT GGG GAT AGG ATT GGG ATT GGG AAG GAT TTT CAG AGC AAG GAG AGT CAG I I I T D G E A T D S G N I D A A K D I I R <u>Y I I G I G K</u> H F Q T K E S Q	982
272	GREACC CTC CAC ARA TITE OCA TCA ARA COC GOC ACC GAG TITE GTG ARA ATT CTG GAC ACA TITE GAC ARA GAT CTA TTC ACT GAG CTG CAG ARA GAT CTAT GTC E T L H K F A S K P A S E F V K I L D T F E K L K D L F T E L Q K K I Y V	1093
309	ATT GAG GGC ACA AGC AMA CAS GAC CTG ACT TOC TTC AAC ATG GAG CTG TOC TOC ACC GGC ATC AGT GCT GAC GGC GAT GCA GTC GTG GGG GCA GTA GGA GGC I E G T S K Q D L T S F N M E L S S S G I S A D L S R G H A V V G A V G A	1204
346	AAG GAC TGG GCT TGG GGC TTT CIT GAC CTC GAG GGC GAC CTG CAG GAC ACA TTT ATT GGC AAT GAA CCA TTG ACA GGC AGA GTG AGA GGC TAT TTG GGT TAC ACA K D W A G G F L D L K A D L Q D D T F I G N E P L T P E V R A G Y L G Y T	: 1315
383	GTG ACC TGG CTG CCC CCG CAA ANG ACT TGG TTG CTG GCC TGG GGA GCC CCT CGA TAC CAG CAC ATG GCC CGA GTG CTG CTG TTC CAA GAG CCA CAG GCC GGA GGA CAC V T W L P S R Q K T S L L A S G A P R Y Q H M G R <u>V L L F Q E P Q G</u> G G B	1426
420	TO CAC CAS STE CAS ACA ATE CAT GOS ACC CAS ATT GOS TET TAT TTE GET GOS GAS CTE TOTE GOS GTE GAS CAS ACA GAS CTE CTE CTE ATT GOS WSQVQTIHGTQIGSYYFGGELCGVDVDQDGETELLLIG	1537
457	CCC CCA CTG TTC TAT GOG GAS CAG AGA GGA GGC CGG GTG TTT ATC TAC CAG AGA AGA CAS TTG GGG TTT GAA GAA GTC TCA GAG CTG CAG GGC GAC COC GGC TAC CCA CTG A P L F Y G E Q R G G R V F I Y Q R R Q L G F E E V S E L Q G D P G Y P L	: 1648
494	GOG GGG TTT GGA GAA GOC ATC ACT GCT GTG ACA GAC ATC ANG GGG GAT GGG GGG GTG GTG GGG GGC GCT GTG GAG GAG GGG GGT GTG TAG ATC TTC AA G R <u>F G E A I T A L T D I</u> N G D G L V D V A V G A P L E E Q G A V Y I F N	: 1759
531	GOG AGG CAC GOG GOG CTT AGT COC CAS GCA AGT CAG GOG ATA GAG GOG AGC CAA GTG CTC TCA GGA ATT CAG TGG TTT GGA GGC TTC GAT GGA GTG GAG GTG AAG GAC CTT GA G R H G G L S P Q P S Q R I E G T Q V L S G I Q W F G R S I H G V K D L E	1872
568	GOG GAT GOC CTG GCA GAT GTG GCT GTG GOG GCT GAG ACC CAG ATG ATG GTG CTG ACC TCG GTG GTG GAT ATG GTC ACC CTG ATG TCC TTC TCT OCA GCT GAG ATG <u>G D G L A D V A V G A E</u> S Q M I V L S S R P V V D M V T L M S F S P A E I	; 1981
605	CCA GTG CAT GAA GTG CAG TAC TAC TAC ACC AST AAC ANG ATG ANA GAA GGA GTT ANT ATC ACA ATC TAT TTC CAG ATC ANG TCT CTC TAC COC CAG TTC CAA GGC CAG P V H E V E C S Y S T S N K M K E G V N I T I C F Q I K S L Y P Q F Q G R	: 2092
642	CTG GTT GCC AAT CTC ACT TAC ACT CTG CAG CTG GAT GGC CAC GGG ACC AGA AGA GGG GGG TTG TTC GCA GGG AGG AAT GAA CTC AGA AGG AAT AGA GCT GTC ACC AGG L V A [N] L] Y T L Q L D G H R T R R R G L F P G G R H E L R R N I A V T T	: 2203
679	AGC ATG TCA TCC ACT TAC TTC TCA TTC COLG GAT TGT GTT GAG GAC CTC ATC TCC COLC AND GAT GTT TCC CTG GAT TTC TCT TGG GAG GAG GAG GAG ACA CGC SMSCTDFSFHFPVCVQDLISPIN <u>VS</u> LM <u>FS</u> LWEEEGTP	; 2314
716	AGG GAC CAA AGG GOC GAG GAC GAG ATA COG GOC ATC CTG AGA COC TOC CTG CAC TOG GAA ACC TOG GAG ATC COT TITT GAG AAG AAC TIGT GOG GAC GAC AAG AAG AG TIGT R D Q R A Q G K D I P P I L R P S L H S E T W E I P F E K N C G E D K K C	2425
753	GAG GCA AAC THE AGA GHE TOUT TOUT TOUT GOLA AGA TOUR AGA GOU CHE GOT CHA ACT GOT TOT GOLAGOUTTE GOLAGOUTTE GAA GAA GAA GAA GAA GAA GAA GAA GAA GA	2536
790	TAC TGG GTC CAG CTG GAC CTG GAC TTC COC GGG GAC CTC TCC TTC COC AAG GTG GAG ATG CTG AAG CCC CAT AGC CAG ATA OCT GTG AGC TGC GAG GAC CTT OCT GAA GAC Y W V Q L D L H F P P G L S F R <u>K V E M L K P H S Q I P V S</u> C E E L P E E	; 2647
827	TO CAGE CTT CTG TO CAGE ACTA TAT TO CAAT GTG AOC TO CO CATO TTC ANA ACA GOA GOC CAG TOG CTT GOT CTG CAG ATG ATG TTT ANT ACA CTG GTA ANC AOC TO TO S R L L S R A L S C N V S S P I F K A G H S V A L Q M M F N T L V N S S W	; 2758
864	GOG GAC TOG GTT GAA TTG CAC GOC AAT GTG AAC TGT AAC AAT GAG GAC TCA GAC CTC CTG GAG GAC AAC TCA GOC ACT ACC ATC ATC COC ATC CTG TAC COC ATC AAC ATC G D S V E L H A N V T C N N E D S D L L E D N S A T T I I P I L Y P I N I	2869
901	CTC ATC CAG GAC CAA GAA GAC TCC ACA CTC TAT GTC AGT TTC ACC COC AAA GGC COC AAG ATC CAG GAC ATG GAC ATG GAC ATG GAC ATG CAG GCT TCC ATC CAC L I Q D Q E D S T L Y V S F T P K G P K I H Q V K H M Y Q V R <u>I Q P S I H</u>	2980
938	CAC CAC ANC ADA COC HOC CTE GAE GCT GTE GTT GGE GTE CCA CAE GCT COC AGE GAE GGE GGE COC ATC ACA CAC CAE TES ACC GTE CAE ATE GAE GCT COC GTE COC TEC CAE D H N I P T L E A V V G V P Q P P S E G P I T H Q W S V Q M E P P V P C H	3091
975	TAT GAG GAT CTG GAG AGG CTC COG GAT GCA GCT GTT GTC TCTC COC GGA GDC CTG TTC COC TCC TCT GTC TTC AGG CAG GAG ATC CTC GTC CAA GTG ATC GGG ACT Y E D L E R L P D A A E P C L P G A L F R C P V V F R Q E I L V Q V I G T	3202
1012	CTG GAG CTG GTG GGA GAG ATC GAG GOC TCT TOC ATG TTC AOC CTC TGC AOC TTC CTC TOC ATC TOC ATG CAC TTC CAC CTC TAT GOC AOC AOC GOC TOC L E L V G E I E A S S M F S L C S S L S I S F <u>N S S</u> K H F H L Y G S <u>N A S</u>	3313
1049	CTG GCC CAG GTT GTC ATG AAG GTT GAC GTG GTG GTG TAT GAG AAG CAG ATG CTG CTG CTG AGG GTG GTG GTG GTG GTG GTG GTG GTG G	3424
1086	CTG TAC ANG GTT TAC TAC ANA COG ANC CTG ANG CAG ANG ANG ANG ANG CAG GOCT GOC AGA GOT GTC COG ANA TOCA ANA GAC TCT GAG CAG CTG GOCA TCT GAG CAG L Y K V G F F K R N L K E K M E A G R G V P N G I P A E D ③ E Q L A ⑤ C Q	, 3535
1023	CAS OCTOBE CARL COLOR CTOC CTIC CARL GAG ANG CAC TOT CARL AFT GET GET GET GET GET GET GET GET GET GE	; 3660
	CCK_TIGGCTICIGCATICIGCCJGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG	C 3808 G 3956 T 4104 C 4252 G 4400 T 4548 G 4696 <u>T</u> 4844 C 4992 A 5139

GAATIOUCICTTICACOCIGICIAGGTIGOCAGCAAAIOOCAGGGOCICCTGAGGCGCCCTGGGGGOCACAGGIOOCTCCAGTGCIGGAAGG



Figure 4. (A) Northern blot analysis of 10 μ g of poly (A+) selected RNA from SKW3 (lane 1), U937 (lane 2), IB4 (lane 3), and EJ cells (lane 4). (B) Identical blot reprobed with the β subunit cDNA probe (23). The position of the 28 and 18 S rRNA is indicated.

which are contiguous and hybridize with different regions of the cDNA (unpublished data) showing that the LFA-1 α subunit is a single copy gene.

Protein Sequence

All 105 amino acids determined by microsequencing of tryptic peptides were found in the translated open reading frame, confirming the authenticity of the cDNA clones (Fig. 3). Hydrophobicity analysis shows that the LFA-1 α subunit is a typical transmembrane protein with a 25-residue hydrophobic signal sequence, an extracellular domain of 1,063 residues, a single hydrophobic transmembrane region of 29 residues, and a short cytoplasmic tail of 53 residues. The NH₂-terminal residue of human LFA-1 was identified by homology to the NH₂-terminal sequence of murine LFA-1 (48) (Fig. 6, see below). Human LFA-1 has 55% identity with murine LFA-1 over the first 20 amino acids. A classical signal peptide with a consensus sequence (Ala-X-Ser/Pro) for the cleavage peptidase precedes the NH₂-terminal sequence. There are three putative upstream transcription initiation sites (ATG) in frame. The use of the first initiation site

(nucleotide position 89) is generally favored (24) and gives a 25-residue signal sequence with several NH_2 -terminal polar groups as is typically found in signal sequences (54).

The mature protein is $M_r = 126,193$. 12 N-linked glycosylation sites (Asn-X-Thr/Ser) are present in the extracellular domain. These findings are consistent with the size previously determined by SDS-PAGE for the in vitro translated murine LFA-1 α subunit ($M_r = 140,000$), the murine and human LFA-1 α subunit glycoproteins ($M_r = 180,000$ and 177,000) (44), and with previous studies on the glycosylation of LFA-1 (10, 32).

Within the extracellular domain there are seven internal repeats (Fig. 5). The degree of identity is highest among the three repeats (14.5-33%) located toward the COOH terminus which show a statistically significant relationship ($P < 10^{-2} - < 10^{-6}$). The relationship among the four repeats located toward the NH₂ terminus is weaker and is discernible by conservation of flanking sequences (Fig. 5); the homology between repeat IV and V is significant ($P < 10^{-4}$). The central regions of the three COOH-terminal repeats is similar with the EF hand divalent cation binding site motif, perhaps due to convergent evolution (Fig. 5). Previous studies have shown that Mg²⁺ alone or at lower concentration in conjunction with Ca²⁺ is necessary for ligand binding func-

^{2.} These data sequence have been submitted to the EMBL/GenBank Data Libraries under the accession number Y00796.

Figure 3. Nucleotide sequence and derived protein sequence of the α subunit of LFA-1.² The potential glycosylation sites are boxed. The sequences of the tryptic peptides and the transmembrane region are underlined with a thick and shaded line, respectively. The putative serine phosporylation sites are circled. The nucleotides in the 3' untranslated region that are underlined correspond to an Alul sequence and the polyadenylation site is boxed.



Figure 5. Alignment of the internal repeats of the LFA-1 α subunit. The consensus flanking sequence is shown above the aligned repeats. Consensus residues were defined as those appearing in at least 30% of the integrin α subunits analyzed (Mac-1 [4, 7], p150,95 [9], vitronectin receptor [50], fibronectin receptor [3], and platelet glycoprotein IIb [37]). The calcium and magnesium binding sites of parvalbumin, troponin C (52), and galactose binding protein (55) are aligned as shown. The asterisk indicates that more than one oxygen is involved in cation binding.

tion (29) (Dustin, M. L., and T. Springer unpublished results), suggesting that these sites bind divalent cations.

Distinct Subfamilies of Integrin α Subunits

We compared the LFA-1 α subunit to other integrin α subunits (Fig. 6). The LFA-1 α subunit has striking and consistently higher homology to other leukocyte integrin α subunits (Mac-1, 35.7%; p150,95, 37.4%) than to ECM receptor integrin α subunits (vitronectin receptor, 26.4%; FNR, 27.8%; gpIIb 30.2%). The ECM receptor integrin α subunits are more related to one another ($\bar{x} = 41.9\%$ SD = 3.7%) than to the leukocyte integrins (Fig. 8, see below). Further structural features distinguish the leukocyte and ECM receptor integrins. The leukocyte integrins contain an insertion of ~ 200 amino acids near the NH₂-terminal region of the protein that is not present in the three sequenced ECM receptor integrins (Fig. 6). Furthermore, a region containing dibasic amino acid protease cleavage sites in vitronectin receptor, FNR, and gpIIb/IIIa (residues 853-871) is absent from LFA-1 as well as p150,95 and Mac-1 α subunits, correlating with the lack of proteolytic processing of the leukocyte integrin α subunits (42). As a whole, these structural features define two subfamilies of the α subunit integrins, the leukocyte integrins and the ECM receptor integrins.

All the integrin α subunits thus far sequenced have tandem repeats similar to those of LFA-1 (3, 9, 37, 41, 51) (Fig. 5). Putative divalent cation binding sites are present in repeats V-VII in all the integrin α subunits and in repeat IV only in the ECM receptor integrins. These sites are similar to the EF hand divalent cation binding sites located in the turn between two α helices in troponin C, parvalbulmin (52), and galactose binding protein (55) among others. In the EF hand loop the metal is chelated to six oxygen-containing side groups spaced every second or third residue. The putative divalent cation binding sites on LFA-1 and the other integrins have a similar primary structure with the putative chelating residues and intervening glycine residues being conserved (Fig. 5). However, the residue in the -Z position is replaced by a hydrophobic residue.

An Inserted Domain in the Leukocyte Integrins

Searches of the NBRF and SWISS-PROT protein sequence data banks revealed that the LFA-1 domain of ~ 200 amino acids, which is not present in the sequenced ECM integrins, is homologous to the domains of the same size in vWF and a cartilage matrix protein (Figs. 7 and 8). These alignments have 20.4-32.1% identity and are statistically significant (P $< 10^{-9} - < 10^{-23}$). Of the integrins sequenced to date, this 200-amino acid domain is unique to the leukocyte integrins. Homologous domains are inserted in several proteins and in well-studied examples have been documented to mediate interaction with ligand (see below). For this reason we will refer to the 200-residue region of the LFA-1, Mac-1, and p150,95 α subunits as the "I" (inserted/interactive) domain. The homology unit is present in three tandem repeats in vWF (A domains) (45) ($P < 10^{-15} - < 10^{-23}$ for comparison of the three repeats to one another) and two repeats separated by an EGF domain in CMP (2) ($P < 10^{-23}$). With the exception of the NH₂-terminal region of the CMP repeat 1 which has not been sequenced, the repeating units in both vWF and CMP correspond precisely to the region homologous to the I domain, supporting the concept that this homology unit is a domain. Similar structural homologies have been noted for the murine and human Mac-1 α subunit (7, 38) and the degree of homology of Mac-1 and p150,95 with vWF and CMP is similar to that found with LFA-1 (17.8-31.6% identity, P $< 10^{-9} - < 10^{-23}$). Factor B has previously been found to be homologous with the vWF A repeats ($P < 10^{-4} - < 10^{-7}$) (45); factor B in turn is homologous but at a lower level with the LFA-1, Mac-1, and p150,95 I domains ($P < 10^{-2} - < 10^{-4}$) (7, 38). In factor B a single homology unit is bounded on one side by the site for the cleavage which activates the B zymogen to the active Bb fragment, and on the other side by the serine protease domain (6).

Based on these homologies, we propose the following pathway for leukocyte integrin evolution. A primordial gene duplicated and gave rise to at least two branches of integrin α subunits, the leukocyte integrins and the ECM integrin receptors. The percent identity among the I domains is very

mLFA1 LFA1 MAC1 p150 IIb VNR FNR	YNLDYRGARS FIS PRAGRHFI YNLDYRGARS FIS PRAGRHF GYRVILOVGRGVIVGAPGEGNSTGSLYQQDSGT-GHCLPYTERGE-MYTSKYLGMT FNLDTENAMTFDENARG-FGDSVVDLQGSRVVVGAPORIVAMDRISLYQCDYST-GSGEFIRLQVF-VEAVNMSLGLS FNLDTEELTAFRVDSAG-FGDSVVDLAGSBVVVGAPORITAMDRIGLYQGIYST-GSGEFIRLQVF-VEAVNMSLGLS INLDTVQLT-FYAGPNGSGFGFSLDFHKDSHGRVA-IVVGAPRTLG-PSQEEFGVFTICPWRAEGGGC-FBLLFDLRDEFRNVGSQT FNLDTVSP-AEYSGPEGSYFGFAVDFFVBSASSRMFLLVGAPANTTQPGIVEGGVVLCPWGASTARGGGC-FBLLFDLRDEFRNVGSQT FNLDTSP-AEYSGPEGSYFGFAVDFFVBSASSRMFLLVGAPANTTQPGVEGGVVLCPWGASTARGGGC-FBLLFDLRDEFRNVGS	73 76 83 84 91
LFA1	LATDPTDGS-LLACDPGLG-ETCDONTYLSGLCYLFRONTQGPMLQGRPGFDECIKGN	129
MAC1	LATTTSPFQ-DLACGPTVH-OTCSENTYVKGLCTLFGSNLRQOPQKFPEALRGCPQED	132
p150	LASTTSPSQ-LLACGPTVH-HECGRMYLIGSLCTLGPTQLTQRLPVSRDEOPRQE	130
IIb	LQTFKARQGLGASVVSWSDVTDACAPHQHNVLEKTEEAEKTPVGSCTLAQPBSGRRAESVSCRENTLSRIYVENDFSWDKRYCEA	169
VNR	PLEFKSHQWFGASVRSKQDKLLACAPLYHWRTEMKQEREPVGTCFLQDGTKTVEYAPCRSQDIDADGQGFCDG	157
FNR	PVEYKSLQWFGASVRSKQDKLLACAPLYHWRTEKEPLSDPVGTCFLSTDNFTRILEYAPCRSDFSWARGQGYCDG	166
LFA1 MAC1 p150 IIb VNR FNR	GISADLS GESAGIT GESAGIT GESAGIT GESEV GESEVT GESEVT GESEVT GESEVT GESEVT GESEVT GESEVT GESEVT	334 344 342 176 164 173
LFA1	R-CHAQVGAVGAKOWAGG-FLDLKADLDDDTFIGNEPLTPEVHAGYLGYLVTWLPSRQXDSLLASGAPRYQHMGRVLLF-O	412
MAC1	SNG-PLLSTVGSYDWAGGVFLVTSKEKSFFINMTRVDSDMNDAYLGYAAAIILKNRVQSLVLAAPRYQHHGRVLLF-O	421
p150	PDD-FVLJAVGSFTNSBGAFLYFPNMSPTFINMSQENVDMRDSYLGYSTELALWKGVQSLVLAAPRYQHHGKAVIFTO	419
IIb	QADELVUGAPGGYTYFLGLLAQAPVADIFSSYRPGILLMUVSQSLSFDSSNPDFTPGYMGYSDAVGEFDGDLNDTEVVDAPHTWSWTLGAVFIL	271
VNR	KTGRVVLGUPGSYGMOCQILSATQEQIASKYDPNVYSIKYNNQ-LATRTAQAIFDGYKGYSDAVGGFSGD-DDEDFVDGSPRAARTLGMVYIYD	257
FNR	KADRVLGGPGSFYMQGQLISDQVAEIVSYYPEYLINDVQGQ-LQTRQASSIYDDSYLGYSDAVGEFSGD-DDEDFVAGVFKGNLTYGYVTILN	266
LFA1	EPQGGGHWSQVQTIHGTQIGSYFGGELCGVDVDQDGENELLLIGAPLFYGEQRGGRVFIYQRROLGFE-EVSELQGDPGYPLGRFGE	498
MAC1	NTGFWESNANVKGTQIGAYFGASLCSVDVDSNGSNDLVLIGAPHYMEQTRGGQVSVCPLPRGQRARWQCDAVLXGEQGDPGYPLGRFGA	507
p150	VSRQWRKAEVTGTQIGSYFGASLCSVDVDTDGSNDLVLIGAPHYMEQTRGGQVSVCPLPRGQRARWQCDAVLXGEQGADPGRFGA	504
IIb	SYYQRLHRLRGEQMAAYFGFSVAATDUNGDGRHDL-LVGAPHYMESADRKLAEVGRVYVLFLQFRGARARW-CDAVLXGEQGADPGRFGA	358
VNR	GKNMSSLYNFGEQMAAYFGFSVAATDUNGDGRHDL-LVGAPLFMDRGSDGKIDEVGQVSVSLQRASGDFQTTKLNGFEVFARFGS	342
FNR	GSDIRSLYNFSGEQMASYFGYAVAATDVNGDGLDDG-IVGAPLFMDRGSDGKIDEVGRVYGYLQRPAGIEPTPTLTLTSHDEF	353
LFA1 MAC1 p150 IIb VNR FNR	ATTALINDINGDGIVDVAVGAP-LEE-DGAVYIFNGRHG-GLSPPSORIEGTOVLSGIONFGRSIHGVKDLGDGIADVAVGAESONIVL-S ALINALGDVNGDRUNDVATGAP-GEEDNRGAVYTIFNGRHG-GLSPGNSORIEGTOVLSGIONFGRSIHGVKDLGDGIADVAVGAESONIVL-S ALINALGDVNGDRUNDVVIGAP-GEEDNRGAVYTIFNG-VGPSISSENSORIAGSOLSSRUDTGOALSGGODLTODGIVDLTVGA-DGHVLLLR ALIANLGDVNGDRUNDVVIGAP-GEEDNRGAVYTIFNG-VGPSISSENSORIAGSOLSSRUDTGOALSGGODLTODGIVDLTVGA-ACGVLLLR AIANLGDUNDGRUNDTAVAN AVGGPSGGOUVUNGLSSEGLSSRUDTGOALSGAVDTDDNGVPGLTVGAYGA AIANLGDUNDGRUNDTAVAN AVGGPSGGOUVUNGLSSEGLSSRUDTGOALSGAVDTDDNGVPGLTVGAYGANGVAVI AIANLGDUNDGRUNDTAVAN AVGGPSGGOUVUNGLSSEGLSSRUDSPFYSSAFGFSLNGAVDTDDNGVPGLTVGAYGANGVAVIA AIANLGDUNDGRUNDTAVAN AVGGPSGGOUVUNGLSSEGLSSRUDSPFYSSASGPSLNGAVDTDDNGVPGLTVGAYGANGVAVI AIANLGDUNDGRUNDTAVGAP-GGETGQGOVFVPFGBYGGILGSKPSQILEGQWA-ARSMPPSFGYSMKGANDTDDKNGVPGLTVGAFGVDRAILYR SUTPLGDLDDGGUNDVAIGAPFGGETGQGOVFVPFGBYGGILGSKPSQVLQPKWA-ASHTPDFFGSALRGBRDTDGNGVPDLIVGAFGVDRAILYR	586 598 595 449 436 447
LFA1	SRFVVDMVTLASFSPAEIPVHEVECSYSTSNAMKEGVNILLEFOLKSLYPOFDGRLVANLTY-FIDEL-DGHRTHRRGLF-PGGRHELRRNIAVT	677
MAC1	SEPVLRVKAINSFNAREVARNVFECND-QVVKGKEAGEVRVCLHVQKSTRDRLREGQIQSV-VTPDLALDSGRPHSRAVFNETKNSTRROT-OVL	690
p150	TREVLWVGVSMDFILPAEIPRSAFECRE-QVVSEQTLODSNICLYTDINRSKNLLGSRDLQSS-VFIDLALDSGRPHSRAVFNETKNSTRROT-OVL	687
IIb	AQPVVKASVQLVQDSLN-PAVKSCVLPQTKTPVSCFNTDFCVGATGHN-IPOK-DSLNAE-LOLDRQK-PRQGR-RVLLGSQQAGTTINLDLG	538
VNR	ARPVITVNAGLEVYPSILNQDNKTCSLPGTALKVSCFNTPFCLKADGKGVLPRK-LNFQVE-LLDLKQKGAI-RRALFUSRSPSHSMMTIS	528
FNR	GRPLVAASASLTIFPAMFNPEERSCSLEGNPVACINLSFCLNASGKHVADSIGFTVE-LQLDWQKQKGGV-RRALFLASTQARDTQTLLIQ	536
LFA1 MAC1 p150 IIb VNR FNR	FERSCTOFSFHFPVCVODLISPINVSLNFSLWEEEGTPROQRAOCKDIPPILRPSLHSETWEIPFEKNCGEDKKCEANLRVSFS-FARSRA GLE-QTCETLKLQLPNCLEDPVSPIVLRLNFSLVGTP-LSAFGNLRPVLAEDAQRLFTALFPFEKNCGNDNICQDDLSTTFSFHSLDCL GLK-AHCENFNLLLP-SCVEDSVTPITLRLNFTLVGRP-LLAFRNLRPVLAEDAQRLFTALFPFEKNCGADHICQDDLSTTFSFBGLKSL GKHEPICHTTNAFLRDEADFRDKLSPIVLSLNVSLVGRP-LLAFRNLRPMLAALAQRYTTASLPFEKNCGADHICQDDLSTTFSFBGLKSL RGGUNGCEELLAYLRDESEFRDKLSPITTFTFTRTL	767 777 774 623 617 625
LFA1	LALTAF-ASLSVELSLSNLEED AYWVDDDHFFPFGLSFRKV-EMLKOPHSOIPVSCEELPESS-RLLSRALSCNVSSE-IFKAGHSVALQFMF	855
MAC1	VVGGPREFNVTVVRNDGED-SYRTOVTFFFPLDLSKRKVSTLRTSGSORSWRLACESASSTEVSGALKSTSC-SINHPIFPENSEVIFNITF	868
p150	LVGSN-LELNADWHVNDGED-SYRTTITSHPAGLSVRTVABGOROGOLSLHLTDSAPVGSQGTWSTSCR-INHLIFRGAQTFLAFF	863
IIb	LVGADNVLELQMDAANEGEG-AYEAELAVHLPOGAHYWRALSNVEGFERLI-O-NQKKENETRVVLCELGNPMKKATQLLAGLAFL	705
VNR	YIGDDNPLTLLWKAQNQGEG-AYEAELAVTAPPEAEYSGLVRHPGNFSSLSQ-DYFAVNQSRLEVCDLGNPMKATQLLAGLRF	699
FNR	YLGDKNALNLTFHAQNVGEGAYEAELAVTAPPEAEYSGLVRHPGNFSSLSQ-DYFAVNQSRLEVCDLGNPMKATQLLAGLRF	708
LFA1	NTLVNESING DEVEDHANVIG NNEDSOLLEDNSATT-IIFILINPINILIQOODO-STLVVSFTFROPKIHOVNEMYOVFIQPSII	936
MAC1	DVDSKASLG-NKQLLKANVISENNMPRTNKTEFQLELEVKYAVYMVVTSHGV-STKVLNFTASENTSRVMOHOVQVSNLGORSLP-I	952
p150	DVSFKAVLG-DRLLLIANVESENNTPRTSKTTFQLELEVKYAVYTVVSSHDD-FIKVLNFTSEEKESHVANHRVQVINLGOROLP-V	948
IIb	SVGNLEEAGESVSFQLQIRSKNSQNPNSKIVLLDVPVRAEAQVELKG-VSFPASLVVAAEEGEREQNSLDSWGFNVEHTVELKNNGPGTVNGL	797
VNR	SVKQOSEMDTSVKFDLQIQSSMGFDKVSFVVSHKVDLAVLETRG-VSSPENIFLPIPNNEMENDEPETEEDVGPVVOHTMELKNNGPSTSKA	793
FNR	TVPHLKDTKKTIQFDFQILSKNLNNSQSDVVSFRLSVEAQAQVTLNG-VSKPEAVLFPVSDWHERDQPQKFEDLGPAVHHVVELINQGPSSISQG	802
LFA1 MAC1 p150 IIb VNR FNR	HDH-NIFT-LEAVWGVPQPPSEGFITHQWSVQMEPPVPCHWEDLERLFDAAEPC SLVFLVPVRLNQTVIWDRPQVTFSENLSSTCHTKE-RLPSHSDFLAELRKAPVUNC SINFWVPVELNQEAVWMDVEVSHPQNPSLRCSSEK-TAPPASDFLAHIQKN	988 1007 1003 880 874 874
LFA1	-LPGAG-FRCPM-VF-ROEIG-VQVICADEGVGEDEASSMFSLCSSLSIFFNSSRHFHLYGSNAS-LAQVVMKVDVVYEKQMDYDV	1069
MAC1	SIAVCQRIGGDIPFFGIQEEFNATLKGNLSFDWYIKTSHNHL-LIVGTAEITFNSSVFTILPGQGAVVRSOTETKVEPFEVPNPLPLIV	1095
p150	SIAGCDAFRCDDISFSVQSEDDFLKGNLSFGWVRQULQKKV-FDVGVA-EITFDTSVYSQLPGQEAFMRAQTTTVLEKYKVHNPTPLIV	1091
IIb	DSAFCTVVGCDLQEMARQRAMMIVLAFLWLPSLYQRPLOFVLQSHAWFNYSSLPXAVPVJSLPAGEAQVWTQLLRALEERATPIWVL	970
VNR	GVAQCDKIVCQDGRLDRGKSAILYVKSLWTETFNKKENQNHSYSLSSAFFNIEFVIFFYLAMPIEDITNSTLVTTNTWGIQPAPMPVPVWV	966
FNR	PEAECFRLRCELGPLHQDESQSLDCHFRVWAKTFLQREHQPFS-LQCEAVYKALKMPYRILPRQLPQKERQVATAVQWTKAEGSYSVPLWI	960
LFA1 MAC1 p150 IIb VNR FNR	LSGU-GGLLLULIFIVLYKVGFFKRNLKEKHEAGRGVPNGIPAGDSEGLASGQ-BAGDPDCLKPLHEKDSESGGGKD GSSV-GGLLLLALITAALYKUGFFKRQYKDMESTGGPPGAEPQ GSSU-GGLLLLALITAAVYKVGFFKRQYKDMESTGGPPGAEPQ V-GVLGGLLLLATITAVYKVGFFKRQ-YKDMESTGGPPGAEPG V-GVLGGLLLLATITAVLKVGFFKRQFPRCBEDDEEGE IILAVLAGLLLLAVVVFMERGFFKRQFPCBEDEFEQLQPH-ENGEGNSET IILAILFGLLLGULIYILYRGFFKRSLPYGTAMEKAQLK-PPATSDA	1145 1137 1144 1008 1018 1008

Figure 6. Alignment and comparison of the human LFA-1 α subunit with the other members of the integrin superfamily (Mac-1 [4, 7], p150,95 [9], vitronectin receptor [50], fibronectin receptor [3], and platelet glycoprotein IIb [37] and the NH₂ terminus of the murine LFA-1 α subunit [48]. The residues common to LFA-1 and at least one other integrin are boxed. The area of the I domain is shown in Fig. 7. The protease cleavage site in the ECM receptor α subunits are indicated with black dots.

LFA1 MAC1 P150 VWFA1 VWFA2 VWFA3 FACTOR CMP2 CMP1	в	130 133 131 513 734 927 244 195 1	VDLVFLFDGSMSLQPDEFQKILDFMKDVMKKLSNTSNQFAAVQFSTSYKTEFDFSDYV SDIAFLIDGSGSIIPHDFRRMKEFVSTVMEQLKKSKTDFSLMQISEEFRIHFIFKEFQ QDLVFLLDGSGSISSISSRNFATMMNFVRAVISQFQRPSTDFSLMQFSNKFQDHFIFEEFR LDLVFLLDGSSRLSEAEFEVLKAFVVDMFERLRI-SQKWVRVAVVEYHDGSHAYIGLK LDVAFVLEGSDKIGEADFNRSKEFMEVLQRMDV-GQDSIHVTVLQYSYMVTVEYPFS LDLVILLDGSSSFPASYFDEMKSFAKAFISKANIGPRLTQVSVLQYGSITTIDVPWN MNIYLVLDGSDSIGASNFTGAKKCLVNLIEKVASYGVKPRYGLVTYATYPKIWVKVSEAD LDLVFLIDGSKSVRENFELVKKFINQ-IVESLEV-SEKQAQVGLVQYSSV-RQEFPL	
LFA1 MAC1 P150 VWFA1 VWFA2 VWFA3 FACTOR CMP2 CMP1	в		KMKDPDALLKHVKHVILLTNTFGAI-NYVATEVFREELGARPDATKVII NNPNPRSLVKPITQLLGRTHTATGIT-KVVRELFNITNGARKNAFKILV RTSNPLSLIASVHQLQGFIYTATAIQ-NVVHRLFHASYGARRDATKILI DRIGRESELRRIASQVKYAGSQVQSTSEVLKYILFQIFSKIDRPGASRIALL EAQSKGDILQRRVREIRYQGGNRINTGLAL-RYLSDHSFLVSOGDREQAPNLVY VVPEKAHLLSLVDVVQREGGPSQIGDALGFAV-RYLTSEGARFGASKAVVI SS-NADWVTKQLNEINYEDHKLKSGTNTKKALQAVYSMMSWPDDVPPEGWNRTRHVII GQFKNKKDIKAA-VKKMAYMEGINTGLAL-KYLVDSSFSIANGARFGVPKVGI KTHQTKAE-LLQAVARIEPLSTGIMTGLAI-QFAISRAFSDTEGARLRSPNINKVAI-	
LFA1 MAC1 P150 VWFA1 VWFA2 VWFA3 FACTOR CMP2 CMP1	в		IITDGE-ATDS-GNIDAAKD-IIRYIIIGIGKHFQIKESQET-LHKFASKPASEFVK VITDGEKFGDPLGYEDVIPEADREGVIRYVIGVGDAFRSEKSRQE-LNTIASKPPRDHVF VITDGKKEGDSLDYKDVIPMADAAGIIRYAIGVGLAFONRNSWKE-LNDIASKPSQEHTF LMASQEPQRMSRNFVRYVQEKKKKULVIPVGIGPHANLKQIRLIEKQA-PENKAFV MVTGNPASDETKRL-PGDIQVVPIGVGPNANVQELIERIGW-PNAP LVTDV-SVDSVDAAADAARSNRVTVFPIGIGDRYDAAQLRILAG-PAGDSNV LMTDGLHNMGGDPITVIDEIRDLLYIGKDRKNPR-EDYLDVYVFGVGP-LVN VFTDGRSQDYITDAAKKAKDLGFRMFAVGVGNAVEDELREIASEPVAEHYF VVTDGRPQDGVQ-VSARARQAGIEIFAIGVGRVDMHTLRQIASEPLDDHVD	
LFA1 MAC1 P150 VWFA1 VWFA2 VWFA3 FACTOR CMP2 CMP1	В		ILDT-FEKLKDLFTELOKKINVIEGTSKODLTSFNMELSSS QVNN-FEALKTIQNOLREKIFAIEGTQTGSSSSFEHEMSQE KVED-FDALKDIQNOLKEKIFAIEGTETTSSSSFELEMAQE LSS-VDE-LEQQRDEIVSYLCDLAPEAPPPTLPPDMAQVTVGP ILIQDFETLPREAPDUVLQRC-CSEGLQIPTLSPAPD VKLQRTEDUPTMUT-LGNSFLHKLCSGFVRI QVNINALASK-KDNEQHVFK-VKDMENLEDVFYQMIDESOSL YTA-DFRTISNIGKLOMKICVEEDPCECKSIVKFQTKV YVESYSVIEKLTHKGFQEAFCVVSDLCATGDHDCEQICI	1

Figure 7. Alignment and comparison of the LFA-1 α subunit I domain with the homologous domains in Mac-1 (4, 7), p150,95 (9), vWF (45), factor B (34), and CMP (3).

similar to the overall percent identity among the leukocyte integrins, implying that a single I domain incorporated into a primordial leukocyte integrin only once rather than independently. Then, the gene duplicated and gave rise to LFA-1 and a Mac-1/p150,95 primordial gene. Further duplication of the Mac-1/p150,95 primordial gene gave rise to Mac-1 and p150,95. This scheme is consistent with the observation that Mac-1 and p150,95 are more closely related to each other (61.0%) than to LFA-1 (35.7, 37.4%), respectively). Since LFA-1, Mac-1, and p150,95 α subunits are located on chromosome 16 band pl1 (8), these α subunits genes remained in close proximity. ECM α subunits appear to be a separate branch since they are more closely related to each other than to the leukocyte integrins even ignoring the I domain (Fig. 8). Although three sequenced α subunits of ECM integrins bind to two different β subunits, the two α subunits associated with the β_3 subunit are not more closely related to one another than to the α subunit associated with the β_1 subunit. Thus, although there are three distinct β subunits, only two subfamilies of α subunits can be distinguished by sequence homology considerations. The structural properties of the leukocyte integrin α subunits may not be unique to their subfamily since some VLA integrin α subunits may have a homologous I domain and lack a proteolytic cleavage site (Hemler, M. personal communication).

The I domain appears to be incorporated in proteins which are otherwise structurally distinct. These structurally similar domains form an important protein superfamily which we will designate the I domain superfamily. Repeats within vWF and within CMP are internally more related than to the domains in other proteins (Fig. 8). Thus, it appears a single domain was incorporated and then duplicated or triplicated. Members of the I domain superfamily serve important recognition functions in several proteins. The A1 domain of vWF binds to glycoprotein Ib and heparin (15) while both the Al and A3 domains are involved in binding to collagen. The domain in factor B is located in a region of the molecule available for interaction with its ligand C3b (6, 34). The domain in CMP may also have an important role in interaction with collagen (2) and cartilage proteoglycan (35). These domains lack glycosylation and cysteines for the most part. Similarly, the I domain and the repeats with divalent cation binding sites in LFA-1 contain only one N-glycosylation site and cysteine (Fig. 8). Similar to the Mac-1 (7, 38) and p150,95 (9) α subunits, the N-glycosylation sites and cysteines are located primarily NH2-terminal and COOH-terminal to this region. Therefore, the I domain would be accessible to ligand and capable of conformational changes which may be important in regulation of ligand binding (39).

337 335 717

 $115 \\ 450 \\ 393 \\ 172$

Structural differences between the ECM receptor integrin α subunits and LFA-1 correlate with differences in recognition specificity; RGD containing peptides block the binding of the VNR, FNR, and gpIIb/IIIa to their ligands (41) but not binding of LFA-1 to ICAM-1 (29). Furthermore, ICAM-1



Figure 8. Schematic representation of the evolutionary relationships of the domains homologous to the I domain. Black lollipops (above schematic) and circles (below schematic) indicate the respective sites of potential glycosylation and cysteines within LFA-1. The percent identity and the standard deviations among the different integrin α subunits and members of the I domain superfamily are shown. Sequences were compared pairwise using the ALIGN program. The I domain sequences were as follows: LFA-1, Mac-1, and p150,95 α subunits (130-327; 133-337; 131-335, respectively), the A repeats of von Willebrand factor (Al: 513-717; A2: 734-922; A3: 927-1115), factor B (245-450), and collagen matrix protein (repeat I: 1-167; repeat 2: 195-393). Relevant probability values are shown in the text.

does not contain an RGD sequence and unlike other known integrin ligands, is a member of the immunoglobulin gene superfamily. It will be of interest to determine whether the I domain confers specificity for non-RGD containing ligands.

This study demonstrates that the LFA-1 α subunit belongs to the integrin superfamily but possesses an additional domain. This I domain and homologous domains constitute a protein "domain" family that is of functional importance. Since LFA-1 is involved in a large number of leukocyte functions and may have more than one ligand (39) (Dustin, M. L., and T. Springer, manuscript in preparation), it is possible the more than one functional domain exists in the LFA-1 α subunit. The availability of cDNA clones for the α and β subunits will allow these and other structure-function relationships to be examined.

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