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Hematologically important mutations: Leukocyte Adhesion Deficiency (first update)

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Abstract

Leukocyte adhesion deficiency (LAD) is an immunodeficiency caused by defects in the adhesion of leukocytes (especially neutrophils) to the blood vessel wall. As a result, patients with LAD suffer from severe bacterial infections and impaired wound healing, accompanied by neutrophilia. In LAD-I, mutations are found in *ITGB2*, the gene that encodes the β subunit of the β_2 integrins. This syndrome is characterized directly after birth by delayed separation of the umbilical cord. In the rare LAD-II disease, the fucosylation of selectin ligands is disturbed, caused by mutations in *SLC35C1*, the gene that encodes a GDP-fucose transporter of the Golgi system. LAD-II patients lack the H and Lewis Le^a and Le^b blood group antigens. Finally, in LAD-III (also called LAD-I/variant) the conformational activation of the hematopoietically expressed β integrins is disturbed,

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leading to leukocyte and platelet dysfunction. This last syndrome is caused by mutations in *FERMT3*, encoding the kindlin-3 protein in all blood cells that is involved in the regulation of β integrin conformation.

Keywords

LAD-I; LAD-II; LAD-III; *ITGB2*; *SLC35C1*; *FERMT3*; β integrins; GDP-fucose transporter; kindlin-3

Introduction

Leukocyte adhesion deficiency (LAD) is an autosomal recessive disorder caused by decreased expression or functioning of CD18, the β_2 subunit of the leukocyte β_2 integrins [1]. This deficiency leads to severe impairment of leukocyte adhesion to the vascular wall and leukocyte migration to sites of infection and inflammation. The patients suffer from recurrent, life-threatening bacterial and fungal infections and from impaired wound healing. Characteristic features are delayed separation of the umbilical cord and strong leukocytosis, especially neutrophilia, during periods of infection. Many LAD patients die at young age despite intensive antibiotic therapy. Bone-marrow transplantation is the treatment of choice. LAD is a rare immunodeficiency, but the exact incidence is not known.

The integrins are transmembrane receptors composed of α and β subunits that mediate cellular adhesive interactions throughout the body. At present 18 α and 8 β subunits have been identified that are loosely organized into integrin families. The β_2 integrins form a family of four heterodimeric proteins, only expressed on leukocytes, with one of four α subunits coupled to a common β_2 subunit: $\alpha_L\beta_2$ (LFA-1, CD11a/CD18), $\alpha_M\beta_2$ (Mac-1 or CR3, CD11b/CD18), $\alpha_X\beta_2$ (p150,95, CD11c/CD18) and $\alpha_D\beta_2$ (CR4, CD11d/CD18), the latter only being expressed on macrophages. Decreased expression of the common β_2 subunit leads to a similar decrease in the expression of all four α subunits on the leukocyte surface. The four β_2 integrins act as adhesion proteins, mediating adhesion of leukocytes to other cells and to extracellular matrix proteins. The α subunits and the β_2 subunit are transmembrane proteins, intracellularly connected to the leukocyte cytoskeleton. Binding to extracellular ligands leads to a conformational change of the β_2 integrins, increased binding of intracellular target proteins and downstream signal transduction to cell spreading and altered gene expression, cell proliferation, differentiation and apoptosis (“outside-in” signaling). Leukocyte activation, e.g. as a result of chemokine binding to chemokine receptors, antigen binding to the T-cell receptor or ligand binding to selectins, induces conformational changes in the extracellular regions of the β_2 integrins, leading to a higher affinity for their ligands (“inside-out” signaling) [2].

Classical LAD-I

In the most common form of LAD, called LAD-I (OMIM #116920), mutations are found in *ITGB2* (integrin beta-2), the gene located at 21q22.3 (OMIM *600065) that encodes the β_2 integrin protein. Usually, this leads to the absence or decreased expression of the β_2 integrins on the leukocyte surface, but sometimes a normal expression of nonfunctional β_2 integrins is

found. In a previous publication we listed 34 mutations found in *ITGB2* of LAD-I patients [3]. In the present publication, 53 newly identified mutations have been added to Table 1 (marked with * in the last column). Mutations that have not been previously published elsewhere are marked as “Unpubl.”. Most of the point mutations are found in a ~240-residue domain that is highly conserved in all β integrin subunits and coded for by exons 5 – 9 of *ITGB2* (figure 1). This “ β I domain”, together with its α I counterpart, constitutes the major ligand-binding site of the β_2 integrins. Both I domains also contain a metal ion-dependent adhesion site (MIDAS motif) consisting of an Asp-X-Ser-X-Ser sequence.

Table 2 contains information on apparently benign polymorphisms that have been identified in *ITGB2*. The mutation c.1756C>T (p.Arg586Trp) in *ITGB2* was found in two non-related patients on the same allele with the mutation c.742-14C>A at the 3' end of intron 6 (leading to the incorporation of four additional amino acids in exon 7) [4,5]. By itself, the 586Trp CD18 molecule, expressed in COS cells, supports about 75% of normal β_2 integrin expression [5], indicating that this mutation may be regarded as a polymorphism. Therefore, the c.1756G/T alternatives are shown as SNPs in Table 2. Other polymorphisms shown in Table 2 have not yet been defined at the functional level.

LAD-II

Two other, extremely rare forms of LAD exist. Patients with LAD-II (OMIM #266265) have a defect in the fucosylation of various cell surface glycoproteins, some of which function as ligands for L-selectin [6]. As a result, the initial “rolling” of leukocytes over the endothelial vessel wall in areas of inflammation, which is mediated by reversible contact between L-selectins on the leukocytes and E- or P-selectins on the endothelial cells with their respective sialated fucosyl ligands on the opposite cells, is disturbed. Both via intracellular signaling and by slowing down the leukocytes, this rolling allows integrins to bind their ligand on endothelial cells, which is needed for stable adhesion. Thus, in LAD-II, one mechanism of β_2 -integrin activation is lacking, leading to decreased leukocyte adhesion to the vessel wall and decreased transendothelial migration into the tissues. However, the chemokine and T-cell receptor pathways of β_2 -integrin activation are still operative, and the infectious episodes in LAD-II patients are therefore in general less severe than those seen in LAD-I patients. On the other hand, the fucosylation defect affects not only selectin ligands but also other essential glycoproteins, leading to severe mental and growth retardation. The molecular defect in LAD-II has been identified as a deficiency in a GDP-fucose transport protein in the Golgi system [7,8]. This protein is encoded by *SLC35C1* (Solute carrier family 35 member C1) at 11p11.2 (OMIM *605881). Table 3 lists the mutations found in this gene in six families with LAD-II patients. Two of these mutations concern Arg147 and Thr308, both highly conserved amino acids in the family of nucleotide-sugar transporters group 2 and suggested to be involved in substrate recognition [9]. Supplementation of fucose led to a substantial clinical improvement and correction of hypofucosylation in the patient homozygous for the Arg147Cys mutation, whereas it was of no benefit to the patients homozygous for the Thr308Arg mutation [7,8]. Possible polymorphisms in *SLC35C1* are also listed in Table 3.

LAD-III (LAD-I/variant)

Finally, patients have been described with a defect in the “inside-out” signaling of leukocytes required for activation of β_2 integrins into structures that bind their ligands with high affinity [10]. These patients, in addition to infections, also present with a bleeding disorder, indicating that the signaling defect also affects the β_3 integrin fibrinogen receptor $\alpha_{IIb}\beta_3$ on blood platelets. The molecular defect of this variant form of LAD (LAD-III or LAD-I/variant, OMIM #612840) has recently been assigned to mutations in *FERMT3* (fermitin family homolog 3) at 11q13.1 (OMIM *607901), the gene encoding kindlin-3, a protein involved in inside-out signaling to all blood cell-expressed β integrins (β_1 , β_2 and β_3) [11-14]. Kindlin-3 is apparently expressed not only in hematopoietic cells, but also in endothelial cells [15]; the biological significance of its deficiency in endothelial cells in LAD-III is not yet clear. A discussion has raged in the literature about the importance of a genetic variation in the gene encoding CalDAG-GEF1 (a guanine nucleotide exchange factor for Rap1, involved in integrin activation) in some patients with LAD-III, in addition to mutations in *FERMT3* found in these patients [11,12,16]. However, since the functional defect in such patients can be corrected by reconstitution with kindlin-3 but not by reconstitution with CalDAG-GEF1 [14], this variation in CalDAG-GEF1 is of no importance for the functional defect in LAD-III patients. Table 4 and figure 1 list the mutations found in *FERMT3* in 21 families with LAD-III patients. A hotspot of p.Arg509X mutations in kindlin-3 points to a founder effect, since these mutations are all found in Turkish families originating from Anatolia.

Final remarks

Additional information about the tabulated mutations and about LAD in general can be found in a recent review [16] and in the cited literature. In Table 1 we have used the notation LAD-I⁰, LAD-I⁻ and LAD-I⁺ for differentiating the various phenotypes of LAD-I. In this nomenclature the superscript symbol indicates whether the protein is present at < 5% of normal expression (⁰), diminished in expression (⁻), i.e. between 5% and 20% of normal expression, or normally present but nonfunctional (⁺). This information is based on immune reactivity of the patients' leukocytes with monoclonal antibodies analyzed by flow cytometry and sometimes on similar analyses of COS cells cotransfected with mutant CD18 molecules and wild-type CD11 molecules. In case this information is not known, this is indicated as (?). In a number of cases functionality of the mutant CD18 proteins was tested in cellular adherence assays to β_2 ligands.

The nucleotide numbering system we have used is based on the cDNA sequence and follows the convention that +1 is the A of the ATG initiation codon. This differs from the numbering of the GenBank sequences; for *ITGB2* (GenBank Accession Number M15395) subtract 72 from the GenBank sequence number to make the initiator A +1. The notation of the mutations follows the recommendations of the Human Genome Variation Society [17] (see also www.hgvs.org/mutnomen). The consequences of the mutations for protein composition have been checked with the Mutalyzer program (www.lovd.nl/mutalyzer) [18]. Where possible we have cross-referenced the mutations indicated in the present article with those in an LAD-I, LAD-II and LAD-III database that lists these patients by accession number.

These databases contain additional biochemical, genetic and clinical information and are available at www.uta.fi/imt/bioinfo/ITGB2base, www.uta.fi/imt/bioinfo/SLC35C1base and www.uta.fi/imt/bioinfo/FERMT3base, respectively. Moreover, information can also be found in the HGMD database at www.hgmd.cf.ac.uk/ac/search.php.

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REFERENCES

1. Anderson, DC.; Smith, CW. Leukocyte adhesion deficiencies. In: Scriver, CR.; Beaudet, AL.; Sly, S.; Valle, D., editors. *The Metabolic and Molecular Basis of Inherited Disease*. McGraw-Hill; New York: 2001. p. 4829-4856.
2. Hynes RO. Integrins: versatility, modulation and signaling in cell adhesion. *Cell*. 1992; 69:11–25. [PubMed: 1555235]
3. Roos D, Law SKA. Hematologically important mutations: Leukocyte Adhesion Deficiency. *Blood Cells Mol. Dis.* 2001; 27:1000–1004. [PubMed: 11831866]
4. Nelson C, Rabb H, Arnaout MA. Genetic cause of leukocyte adhesion deficiency. Abnormal splicing and a missense mutation in a conserved region of CD18 impair cell surface expression of β_2 integrins. *J. Biol. Chem.* 1992; 267:3351–3357. [PubMed: 1346613]
5. Wright AH, Douglass WA, Taylor GM, Lau Y-L, Higgins, Davies KA, Law SKA. Molecular characterization of leukocyte adhesion deficiency in six patients. *Eur. J. Immunol.* 1995; 25:717–722. [PubMed: 7705401]
6. Etzioni A, Frydman M, Pollack S, Avidor I, Phillips ML, Paulson JC, Gershoni-Baruch R. Recurrent severe infections caused by a novel leukocyte adhesion deficiency. *N. Engl. J. Med.* 1992; 327:1789–1792. [PubMed: 1279426]
7. Luhn K, Wild MK, Eckhardt M, Gerardy-Schahn R, Vestweber D. The gene defective in leukocyte adhesion deficiency II encodes a putative GDP-fucose transporter. *Nat. Genet.* 2001; 28:69–72. [PubMed: 11326279]
8. Lubke T, Marquardt T, Etzioni A, Hartmann E, von Figura K, Korner C. Complementation cloning identifies CDG-IIc, a new type of congenital disorders of glycosylation, as a GDP-fucose transporter deficiency. *Nat. Genet.* 2001; 28:73–76. [PubMed: 11326280]
9. Handford M, Rodriguez-Furlán C, Orellana A. Nucleotide-sugar transporters: structure, function and roles in vivo. *Braz. J. Med. Biol. Res.* 2006; 39:1149–1158. [PubMed: 16981043]
10. Kuijpers TW, van Lier RAW, Hamann D, de Boer M, Thung LY, Weening RS, Verhoeven AJ, Roos D. Leukocyte adhesion deficiency type 1 (LAD-1)/variant. A novel immunodeficiency syndrome characterized by dysfunctional β_2 integrins. *J. Clin. Invest.* 1997; 100:1725–1733. [PubMed: 9312170]
11. Mory A, Feigelson SW, Yarali N, Kilic SS, Bayhan GI, Gershoni-Baruch R, Etzioni A, Alon R. Kindlin-3: a new gene involved in the pathogenesis of LAD-III. *Blood*. 2008; 112:2591. [PubMed: 18779414]
12. Kuijpers TW, van de Vijver E, Weterman MA, de Boer M, Tool AT, van den Berg TK, Moser M, Jakobs ME, Seeger K, Sanal O, Unal S, Cetin M, Roos D, Verhoeven AJ, Baas F. LAD-1/variant syndrome is caused by mutations in FERMT3. *Blood*. 2009; 113:4740–4746. [PubMed: 19064721]
13. Moser M, Bauer M, Schmid S, Ruppert R, Schmidt S, Sixt M, Wang HV, Sperandio M, Fässler R. Kindlin-3 is required for beta2 integrin-mediated leukocyte adhesion to endothelial cells. *Nat. Med.* 2009; 15:300–305. [PubMed: 19234461]
14. Svensson L, Howarth K, McDowall A, Patzak I, Evans R, Ussar S, Moser M, Metin A, Fried M, Tomlinson I, Hogg N. Leukocyte adhesion deficiency-III is caused by mutations in KINDLIN3 affecting integrin activation. *Nat. Med.* 2009; 15:306–312. [PubMed: 19234463]

15. Bialkowska K, Ma YQ, Bledzka K, Sossey-Alaoui K, Izem L, Zhang X, Malinin N, Qin J, Byzova T, Plow EF. *J. Biol. Chem.* 2010; 285:18640–18649. [PubMed: 20378539]
16. Etzioni A. Genetic etiologies of leukocyte adhesion defects. *Curr. Opin. Immunol.* 2009; 21:481–486. [PubMed: 19647987]
17. Den Dunnen JT, Antonarakis SE. Mutation nomenclature extensions and suggestions to describe complex mutations: a discussion. *Hum. Mutat.* 2000; 15:7–12. [PubMed: 10612815]
18. Wildeman M, van Ophuizen E, den Dunnen JT, Taschner PE. Improving sequence variant descriptions in mutation databases and literature using the Mutalyzer sequence variation nomenclature checker. *Hum. Mutat.* 2008; 29:6–13. [PubMed: 18000842]
19. Rivera-Matos IR, Rakita RM, Mariscalco MM, Elder FFB, Dreyer SA, Cleary TG. Leukocyte adhesion deficiency mimicking Hirschsprung disease. *J. Pediatr.* 1995; 127:755–757. [PubMed: 7472832]
20. Castriconi R, Dondero A, Cantoni C, Della Chiesa M, Prato C, Nanni M, Fiorini M, Notarangelo L, Parolini S, Moretta L, Notarangelo L, Moretta A, Bottino C. *Blood.* 2007; 109:4873–4881. [PubMed: 17272509]
21. Fiorini M, Piovani G, Schumacher RF, Magri C, Bertini V, Mazzolari E, Notarangelo L, Notarangelo LD, Barlati S. ITGB2 mutation combined with deleted ring 21 chromosome in a child with leukocyte adhesion deficiency. *J. Allergy Clin. Immunol.* 2009; 124:1356–1358. [PubMed: 19864007]
22. Roos D, Meischl C, de Boer M, Simsek S, Weening RS, Sanal O, Tezcan I, Güngör T, Law SKA. Genetic analysis of patients with leukocyte adhesion deficiency. Genomic sequencing reveals otherwise undetectable mutations. *Exp. Hematol.* 2002; 30:252–261. [PubMed: 11882363]
23. Sligh JE Jr, Hurwitz MY, Zhu C, Anderson DC, Beaudet AL. An initiation codon mutation in CD18 in association with the moderate type of leukocyte adhesion deficiency. *J. Biol. Chem.* 1992; 267:714–718. [PubMed: 1346132]
24. Cher THB, Chan HS, Klein GF, Jabkowski J, Schadenböck-Kranzl G, Zach O, Law SKA. A novel 3'splice-site mutation and a novel gross deletion in leukocyte adhesion deficiency (LAD)-1. *Biochem. Biophys. Res. Commun.* 2011; 404:1099–1104. [PubMed: 21195692]
25. Lopez Rodriguez C, Nueda A, GrosPierre B, Sanchez-Madrid F, Fischer A, Springer TA, Corbi AL. Characterization of two new CD18 alleles causing severe leukocyte adhesion deficiency. *Eur. J. Immunol.* 1993; 23:2792–2798. [PubMed: 7901025]
26. Fiorini M, Vermi W, Facchetti F, Moratto D, Alessandri G, Notarangelo L, Caruso A, Grigolato P, Ugazio AG, Notarangelo LD, Badolato R. Defective migration of monocyte-derived dendritic cells in LAD-1 immunodeficiency. *J. Leukoc. Biol.* 2002; 72:650–656. [PubMed: 12377933]
27. Hinze CH, Lucky AW, Bove KE, Marsh RA, Bleesing JH, Passo MH. Leukocyte adhesion deficiency type I presenting with recurrent pyoderma gangrenosum and flaccid scarring. *Pediatr. Dermatol.* 2010; 27:500–503. [PubMed: 20807363]
28. Tone Y, Wada T, Shibata F, Toma T, Hashida Y, Kasahara Y, Koizumi S, Yachie A. Somatic revertant mosaicism in a patient with leukocyte adhesion deficiency type 1. *Blood.* 2007; 109:1182–1184. [PubMed: 17244687]
29. Allende LM, Hernandez M, Corell A, Garcia-Perez MA, Varela P, Moreno A, Caragol I, Garcia-Martin F, Guillen-Perales J, Olive T, Espanol T, Arnaiz-Villena A. A novel CD18 genomic deletion in a patient with severe leukocyte adhesion deficiency: a possible CD2/lymphocyte function-associated antigen-1 functional association in humans. *Immunology.* 2000; 99:440–450. [PubMed: 10712675]
30. Parvaneh N, Mamishi S, Rezaei A, Rezaei N, Tamizifar B, Parvaneh L, Sherkat R, Ghalehbaghi B, Kashef S, Chavoshzadeh Z, Isaiean A, Ashrafi F, Aghamohammadi A. Characterization of 11 new cases of leukocyte adhesion deficiency type I with seven novel mutations in the ITGB2 gene. *J. Clin Immunol.* 2010; 30:756–760. [PubMed: 20549317]
31. Matsuura S, Kishi F, Tsukahara M, Nunoi H, Matsuda I, Kobayashi K, Kajii T. Leukocyte adhesion deficiency: identification of novel mutations in two Japanese patients with a severe form. *Biochem. Biophys. Res. Commun.* 1992; 184:1460–1467. [PubMed: 1590804]
32. Uzel G, Tng E, Rosenzweig SD, Hsu AP, Shaw JM, Horwitz ME, Linton GF, Anderson SM, Kirby MR, Oliveira JB, Brown MR, Fleisher TA, Law SKA, Holland SM. Reversion mutations in

- patients with leukocyte adhesion deficiency type-1 (LAD-1). *Blood*. 2008; 111:209–218. [PubMed: 17875809]
33. Lorusso F, Kong D, Jalil AK, Sylvestre C, Tan SL, Ao A. Preimplantation genetic diagnosis of leukocyte adhesion deficiency type I. *Fertil. Steril.* 2006; 85:494.e15–494.e18. [PubMed: 16595236]
 34. Hogg N, Stewart MP, Scarth SL, Newton R, Shaw JM, Law SKA, Klein N. A novel leukocyte adhesion deficiency caused by expressed but nonfunctional $\beta 2$ integrins Mac-1 and LFA-1. *J. Clin. Invest.* 1999; 103:97–106. [PubMed: 9884339]
 35. Wardlaw AJ, Hibbs ML, Stacker SA, Springer TA. Distinct mutations in two patients with leukocyte adhesion deficiency and their functional correlates. *J. Exp. Med.* 1990; 172:335–345. [PubMed: 1694220]
 36. Uzel, G.; Kuhns, D.; Hussey, A.; Spalding, C.; Stoddard, J.; Hsu, A.; Holland, S. The new face of leukocyte adhesion deficiency type 1 (LAD-1). XIVth Meeting of the European Society for Immunodeficiencies; 132: 2010. (Abstr. P256)
 37. Corbi AL, Vara A, Ursa A, Garcia Rodriguez MC, Fontan G, Sanchez-Madrid F. Molecular basis for a severe case of leukocyte adhesion deficiency. *Eur. J. Immunol.* 1992; 22:1877–1881. [PubMed: 1352501]
 38. Back AL, Kwok WW, Hickstein DD. Identification of two molecular defects in a child with leukocyte adhesion deficiency. *J. Biol. Chem.* 1992; 267:5482–5487. [PubMed: 1347532]
 39. Ohashi Y, Yambe T, Tsuchiya S, Kikuchi H, Konno T. Familial genetic defect in a case of leukocyte adhesion deficiency. *Hum. Mutat.* 1993; 2:458–467. [PubMed: 7509236]
 40. Malawista SE, de Boisfleury Chevance A, Brown EJ, Boxer LA, Law SKA. Chemotaxis of non-compressed blood polymorphonuclear leukocytes from an adolescent with severe leukocyte adhesion deficiency. *Am. J. Hematol.* 2003; 73:115–120. [PubMed: 12749013]
 41. Simon AJ, Lev A, Wolach B, Gavrieli R, Amariglio N, Rosenthal E, Gazit E, Eyal E, Rechavi G, Somech R. The effect of gentamycin-induced readthrough on a novel premature termination codon of CD18 leukocyte adhesion deficiency patients. *PLoS ONE*. 2010; 5:e13659. [PubMed: 21103413]
 42. Pollard AJ, Heale J-P, Tsang A, Massing B, Speert DP. Nonopsonic phagocytosis of *Pseudomonas aeruginosa*: insights from an infant with leukocyte adhesion deficiency. *Pediatr. Infect. Dis. J.* 2001; 20:452–454. [PubMed: 11332677]
 43. Mathew EC, Shaw JM, Bonilla FA, Law SKA, Wright DA. A novel point mutation in CD18 causing the expression of dysfunctional CD11/CD18 leucocyte integrins in a patient with leucocyte adhesion deficiency. *Clin. Exp. Immunol.* 2000; 121:133–138. [PubMed: 10886250]
 44. Shaw JM, Al-Shamkhani A, Boxer LA, Buckley CD, Dodds AW, Klein N, Roberts I, Roos D, Scarth SL, Simmons DL, Tan S-M, Law SKA. Identification of four CD18 mutations in leukocyte adhesion deficient (LAD) patients with differential abilities to associate with the CD11a, CD11b and CD11c antigens. *Clin. Exp. Immunol.* 2001; 126:311–318. [PubMed: 11703376]
 45. Back AL, Kerkering M, Baker D, Bauer TR, Embree LJ, Hickstein DD. A point mutation associated with Leukocyte Adhesion Deficiency Type I of moderate severity. *Biochem. Biophys. Res. Commun.* 1993; 193:912–918. [PubMed: 7686755]
 46. Kishimoto TK, O'Connor K, Springer TA. Leukocyte adhesion deficiency. Aberrant splicing of a conserved integrin sequence causes a moderate deficiency phenotype. *J. Biol. Chem.* 1989; 264:3588–3595. [PubMed: 2464599]
 47. Tsai Y-C, Lee W-I, Huang J-L, Hung I-J, Jaing T-H, Yao T-C, Chen M-T, Kuo M-L. Neutrophil function and molecular analysis in severe leukocyte adhesion deficiency type I without separation delay of the umbilical cord. *Pediatr. Allergy Immunol.* 2008; 19:25–32. [PubMed: 17651379]
 48. Li L, Jin YY, Cao RM, Chen TX. A novel point mutation in CD18 causing leukocyte adhesion deficiency in a Chinese patient. *Chin. Med. J. (Engl)*. 2010; 123:1278–1282. [PubMed: 20529581]
 49. Cheng M, Foo SY, Shi ML, Tang RH, Kong LS, Law SKA, Tan SM. Mutation of a conserved asparagine in the I-like domain promotes constitutively active integrins α L β 2 and α L β 3. *J. Biol. Chem.* 2007; 282:18225–18232. [PubMed: 17468108]
 50. Fathallah DM, Jamal T, Barbouche MR, Bejaoui M, Hariz MB, Dellagi K. Two novel frame shift, recurrent and de novo mutations in the ITGB2 (CD18) gene causing leukocyte adhesion deficiency

- in a highly inbred North African population. *J. Biomed. Biotechnol.* 2001; 1:114–121. [PubMed: 12488604]
51. Halioui-Louhaichi S, Ben Hariz M, Fathallah Dahmani M, Barbouch MR, Mahmoudi J, Bejaoui M, Fischer A, Dellagi K, Ben Ammar B, Maherzi A. Le défaut d'expression des protéines d'adhésion leucocytaires. A propos d'une nouvelle observation Tunisienne [Deficient expression of leukocyte adhesion proteins. A new Tunisian case]. *Tunis Med.* 2006; 84:464–466. [PubMed: 17039742]
 52. Hixson P, Smith CW, Shurin SB, Tosi MF. Unique CD18 mutations involving a deletion in the extracellular stalk region and a major truncation of the cytoplasmic domain in a patient with leukocyte adhesion deficiency type 1. *Blood.* 2004; 103:1105–1113. [PubMed: 14512306]
 53. Arnaout MA, Dana N, Gupta SK, Tenen DG, Fathallah DM. Point mutations impairing cell surface expression of the common β subunit (CD18) in a patient with leukocyte adhesion molecule (Leu-CAM) deficiency. *J. Clin. Invest.* 1990; 85:977–981. [PubMed: 1968911]
 54. Eyerich K, Cifaldi L, Notarangelo LD, Porta F, Notarangelo L, Mazzolari E, Fiorini M, Paradisi A, Cavani A. Chronic eczema in a patient with Leukocyte Adhesion Deficiency (LAD) type I. *Eur. J. Dermatol.* 2009; 19:78–79. [PubMed: 19171538]
 55. Hogg N, Bates PA. Genetic analysis of integrin function in man: LAD-I and other syndromes. *Matrix Biol.* 2000; 19:211–222. [PubMed: 10936446]
 56. Arnaout, AM.; Michishita, M. Genetic abnormalities in Leukocyte Adhesion Molecule deficiency.. In: Gupta, S.; Griscelli, C., editors. *Immunodeficiency Diseases.* John Wiley; Chichester: 1993. p. 191-202.
 57. Vihinen, M., et al. Primary immunodeficiency mutation database.. In: Hall, JC.; Dunlap, JC.; Friedmann, T.; Gianelli, F., editors. *Advances in Genetics.* Academic Press; San Diego: 2001. p. 103-188.
 58. Moore SW, Sidler D, Zaahl MG. The *ITGB2* immunomodulatory gene (CD18), enterocolitis, and Hirschsprung's disease. *J. Pediatr. Surg.* 2008; 43:1439–1444. [PubMed: 18675632]
 59. Meller S, Jagiello P, Borgmann S, Fricke H, Epplen JT, Gencik M. Novel SNPs in the CD18 gene validate the association with MPO-ANCA⁺ vasculitis. *Genes Immun.* 2001; 2:269–272. [PubMed: 11528520]
 60. Hidalgo A, Ma S, Peired AJ, Weiss LA, Cunningham-Rundles C, Frenette PS. Insights into leukocyte adhesion deficiency type 2 from a novel mutation in the GDP-fucose transporter gene. *Blood.* 2003; 101:1705–1712. [PubMed: 12406889]
 61. Etzioni A, Sturla L, Antonellis A, Green ED, Gershoni-Baruch R, Berninsone PM, Hirschberg CB, Tonetti M. Leukocyte adhesion deficiency (LAD) type II/carbohydrate deficient glycoprotein (CDG) IIc founder effect and genotype/phenotype correlation. *Am. J. Med. Genet.* 2002; 110:131–135. [PubMed: 12116250]
 62. Helmus Y, Denecke J, Yakubenia S, Robinson P, Lühn K, Watson DL, McGrogan PJ, Vestweber D, Marquardt T, Wild MK. Leukocyte adhesion deficiency II patients with a dual defect of the GDP-fucose transporter. *Blood.* 2006; 107:3959–3966. [PubMed: 16455955]
 63. Gazit Y, Mory A, Etzioni A, Frydman M, Scheuerman O, Gershoni-Baruch R, Garty BZ. Leukocyte adhesion deficiency type II: long-term follow-up and review of the literature. *J. Clin. Immunol.* 2010; 30:308–313. [PubMed: 20099014]
 64. Malinin NL, Zhang L, Choi J, Ciocea A, Razorenova O, Ma YQ, Podrez EA, Tosi M, Lennon DP, Caplan AI, Shurin SB, Plow EF, Byzova TV. A point mutation in *KINDLIN3* ablates activation of three integrin subfamilies in humans. *Nat. Med.* 2009; 15:313–318. [PubMed: 19234460]
 65. Robert P, Canault M, Farnarier C, Nurden A, Grosdidier C, Barlogis V, Bongrand P, Pierres A, Chambost H, Alessi MC. A novel Leukocyte Adhesion Deficiency III variant: Kindlin-3 deficiency results in integrin and nonintegrin-related defects in different steps of leukocyte adhesion. *J. Immunol.* 2011; 186:5273–5283. [PubMed: 21441448]
 66. Manevich-Mendelson E, Feigelson SW, Pasvolksy R, Aker M, Grabovsky V, Shulman Z, Kilic SS, Rosenthal-Allieri MA, Ben-Dor S, Mory A, Bernard A, Moser M, Etzioni A, Alon R. Loss of kindlin-3 in LAD-III eliminates LFA-1 but not VLA-4 adhesiveness developed under shear flow conditions. *Blood.* 2009; 114:2344–2353. [PubMed: 19617577]

67. McDowall A, Svensson L, Stanley P, Patzak I, Chakravarty P, Howarth K, Sabnis H, Briones M, Hogg N. Two mutations in the *KINDLIN3* gene of a new leukocyte adhesion deficiency III patient reveal distinct effects on leukocyte function in vitro. *Blood*. 2010; 115:4834–4842. [PubMed: 20357244]
68. Jurk K, Schulz AS, Kehrel BE, Räßle D, Schulze H, Möbest D, Friedrich WW, Omran H, Deak E, Henschler R, Scheele JS, Zieger B. Novel integrin-dependent platelet malfunction in siblings with leukocyte adhesion deficiency-III (LAD-III) caused by a point mutation in *FERMT3*. *Thromb. Haemost.* 2010; 103:1053–1064. [PubMed: 20216991]

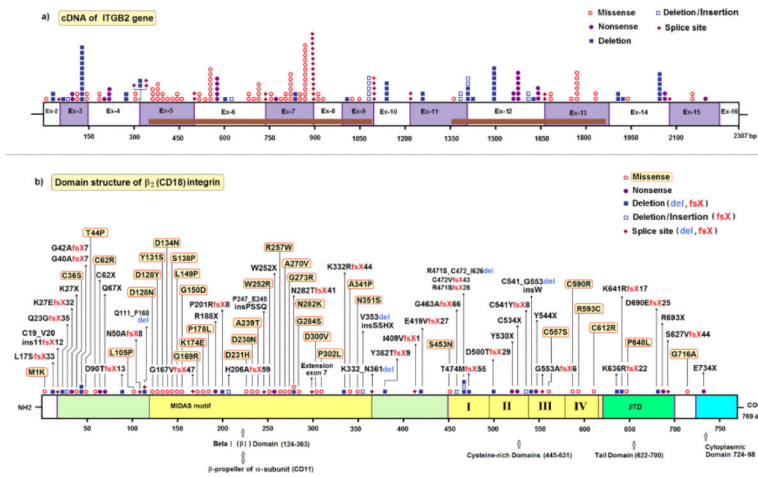
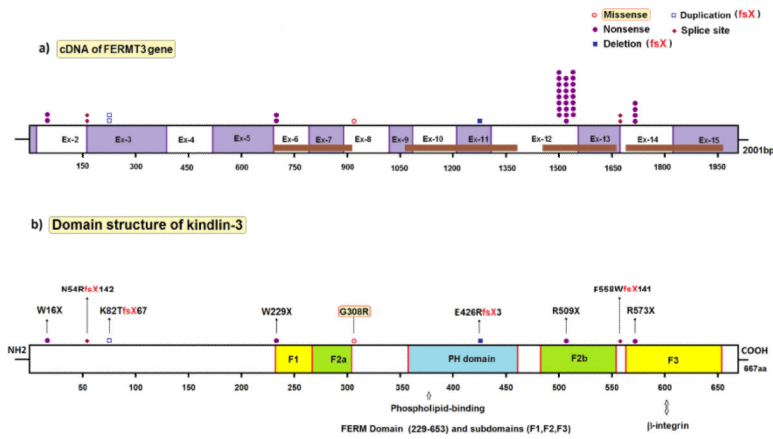


Figure 1. Schematic overview of mutations in *ITGB2*. a. The number of mutated alleles, the type of mutation and their position along the exons of *ITGB2*, with active domains depicted in brown (see underneath fig. 1b). Symbols, explained on the right, represent mutated alleles. Numbers refer to nucleotides in cDNA. b. Domain structure of the β_2 integrin (CD18) and the location of each mutation, with active domains depicted in yellow, green and blue. Protein interaction sites are shown for some domains. Symbols, explained on the right, represent separate mutations. Numbers refer to amino acids in β_2 integrin. Figure drawn by Dr. M. Yavuz Köker.

**Figure 2.**

Schematic overview of mutations in *FERMT3*. a. The number of mutated alleles, the type of mutation and their position along the exons of *FERMT3*, with active domains depicted in brown (see underneath fig. 2b). Symbols, explained on the right, represent mutated alleles. Numbers refer to nucleotides in cDNA.

b. Domain structure of kindlin-3 and the location of each mutation, with active domains depicted in yellow, green and blue. Protein interaction sites are shown for some domains. Symbols, explained on the right, represent separate mutations. Numbers refer to amino acids in kindlin-3.

Figure drawn by Dr. M. Yavuz Köker.

TABLE I

Mutations in the LAD-I gene *ITGB2*

Nucleotide change	Mutation	Amino-acid or mRNA change	LAD-I type	Families (alleles)	Reference	Accession Number
Not determined	Deletion	Entire gene	LAD-I ^o	1(1)	[3, 19]	A0032
Not determined	Deletion	Entire gene	LAD-I ^o	1(1)	[20, 21]	A0051
Not determined	Deletion	Entire gene	LAD-I ^f	1(1)	Unpubl.	A0085
c.-721_58+51del833	Deletion	Deletion exon 2, including start codon	LAD-I ^o	1(2)	[3, 22]	A0063 A0064
c.2T>A	Missense	p.Met1Lys	LAD-I ^f	1(1)	[3, 23]	A0016
c.49delC	Deletion	p.Leu17SerfsX33	LAD-I ^o	1(2)	Unpubl.	A0086
c.59-10C>A ^a	Splice site	Ins 43 nts from intron 2 r.59-43_59-1ins43;59-10C>A p.Cys19_Val20ins1IX12	LAD-I ^o	1(1)	[24]	A0126
c.66_67delTC	Deletion	p.Gln23GlyfsX35	LAD-I ^o	1(1)	[3, 22]	A0028 A0029
c.77dupC	Insertion	p.Lys27GlufsX32	LAD-I ^o	1(1)	Unpubl.	A0079
c.79A>T	Nonsense	p.Lys27X	LAD-I ^o	2(2)	[20, 21] Unpubl.	A0051 A0114
c.106T>A	Missense	p.Cys36Ser	LAD-I ^o	1(2)	Unpubl.	A0088
c.119_128delGGCCCCGGCTG	Deletion	p.Gly40AlafsX7	LAD-I ^o	5(10) ^b	[3, 20, 25, 26] Unpubl.	A0018 A0068 A0083 A0089 A0133
c.120delG	Deletion	p.Gly42AlafsX7	LAD-I ^o	1(1)	Unpubl.	A0127
c.130A>C	Missense	p.Thr44Pro	LAD-I ^o	2(2)	Unpubl.	A0090 A0091
Not identified	Splice site?	Deletion exon 4 p.Asn50AlafsX8	LAD-I ^o	1(2)	[3, 26]	A0081
c.184T>C	Missense	p.Cys62Arg	LAD-I ^f	1(2)	Unpubl.	A0092
c.186C>A	Nonsense	p.Cys62X	LAD-I ^f	1(1)	Unpubl.	A0093
c.199C>T	Nonsense	p.Gln67X	LAD-I ^o	2(3)	[3, 22] Unpubl.	A0005 A0065 A0114
c.268delG	Deletion	p.Asp90ThrfsX13	LAD-I ^f	1(2)	Unpubl.	A0094
c.314T>C	Missense	p.Leu105Pro	LAD-I ^f	1(2) ^b	[27]	A0095

Nucleotide change	Mutation	Amino-acid or mRNA change	LAD-I type	Families (alleles)	Reference	Accession Number
c.328+1G>A ^a	Splice site	Deletion exon 4 p.Asn50AlafsX8	LAD-I ^o	1(1)	[28]	A0050
c.329-37_461del169 ^a	Deletion	Deletion exon 5 p.Gln111_Phe168del	LAD-I ^o	1(2)	[3, 29]	A0038
c.329-6C>A ^a	Splice site	Deletion exon 5? p.Gln111_Phe168del?	LAD-I ^o	1(2)	[30]	A0053
c.382G>A	Missense	p.Asp128Asn	LAD-I ^o	2(3)	[3, 31] Unpubl.	A0002 A0117
c.382G>T	Missense	p.Asp128Tyr	LAD-I ^{o/-}	2(4)	[30]	A0056 A0059
c.392A>C ^c	Missense	p.Tyr131Ser	LAD-I ^o	1(2)	[32]	A0046
c.400G>A	Missense	p.Asp134Asn	LAD-I [?]	1(1)	[33] Unpubl.	A0076
c.412T>C	Missense	p.Ser138Pro	LAD-I ⁺	1(1)	[3,34]	A0003
c.446T>C	Missense	p.Leu149Pro	LAD-I ⁺	2(2)	[3, 5, 22, 35]	A0017 A0078
c.449G>A	Missense	p.Gly150Asp	LAD-I ⁺	1(1)	[36]	A0099
c.500-12T>G ^a	Splice site	Deletion part of exon 6 p.Gly167ValfsX47	LAD-I ^o	1(2)	[3, 22] Unpubl.	A0033
c.505G>A	Missense	p.Gly169Arg	LAD-I ^o	2(2)	[3, 35, 37]	A0010 A0027
c.520A>G	Missense	p.Lys174Glu	LAD-I ^o	1(1)	Unpubl.	A0125
c.533C>T	Missense	p.Pro178Leu	LAD-I ^o	5(7) ^b	[3, 36, 38, 39, 40] Unpubl.	A0020 A0031 A0073 A0074 A0080 A0099
c.562C>T	Nonsense	p.Arg188X	LAD-I [?]	3(5)	[32, 41] Unpubl.	A0076 A0097 A0098 A0122
c.602delC	Deletion	p.Pro201ArgfsX8	LAD-I ^o	1(1)	[28]	A0050
c.614_615insA	Insertion	p.His206AlafsX59	LAD-I ^o	1(1)	[3, 42]	A0021
c.691G>C	Missense	p.Asp231His	LAD-I ⁺	2(2)	[3, 43] Unpubl.	A0013 A0075
c.712G>A	Missense	p.Asp238Asn	LAD-I [?]	1(1)	Unpubl.	A0113
c.715G>A	Missense	p.Ala239Thr	LAD-I ^o	2(4) ^b	[30] Unpubl.	A0055 A0130
c.742-14C>A ^{a,c,d}	Splice site	p.Pro247_Glu248 insProSerSerGln	LAD-I ^o	2(2)	[3, 4, 5, 22, 32]	A0028 A0029 A0030
c.754T>C	Missense	p.Trp252Arg	LAD-I ^o	1(2)	[3, 22]	A0004

Nucleotide change	Mutation	Amino-acid or mRNA change	LAD-I type	Families (alleles)	Reference	Accession Number
c.755G>A	Nonsense	p.Trp252X	LAD-I ^p	1(1)	Unpubl.	A0075
c.769C>T	Missense	p.Arg257Trp	LAD-I ^{o/-}	1(1)	Unpubl.	A0131
c.809C>T	Missense	p.Ala270Val	LAD-I ^{o/-}	2(2)	[3, 44] Unpubl.	A0007 A0111
c.817G>A	Missense	p.Gly273Arg	LAD-I ^p	5(6) ^b	[3, 34] Unpubl.	A0003 A0093 A0113 A0118 A0123
c.843delC	Deletion	p.Asn282ThrfsX41	LAD-I ^p	1(2)	[30]	A0060
c.846C>A	Missense	p.Asn282Lys	LAD-I ^t	1(2) ^b	Unpubl.	A0124
c.850G>A ^c	Missense	p.Gly284Ser	LAD-I ^p	10(12)	[3, 4, 5, 32, 43, 44, 46] Unpubl.	A0011 A0012 A0013 A0014 A0023 A0043 A0044 A0047 A0072 A0091 A0116
c.897+1G>A ^d	Splice site	Extension exon 7 with 64 or 298 nus from intron 7, or with whole intron 7, fsX26 or fsX44	LAD-I ^p	8(14)	[3, 30, 31, 47] Unpubl.	A0039 A0049 A0057 A0058 A0085 A0100 A0105 A0121
c.899A>T	Missense	p.Asp300Val	LAD-I ^p	1(2)	[48]	A0084
c.905C>T	Missense	p.Pro302Leu	LAD-I ^p	1(1)	Unpubl.	A0079
c.995_1004del10	Deletion	p.Lys332ArgfsX44	LAD-I ^t	1(1)	Unpubl.	A0117
c.1021G>C	Missense	p.Ala341Pro	LAD-I ^{o/-}	1(1)	[3, 22, 44]	A0006
c.1052A>G ^c	Missense	p.Asn351Ser	Activated LFA-I	1(1)	[3, 4, 32, 49]	A0030
c.1057_1059delins35	Deletion/insertion	p.Val353del insSerSerHisX	LAD-I ^p	3(5)	Unpubl.	A0104 A0106 A0107
c.1083+3G>C ^d	Splice site	Deletion exon 9 p.Lys332_Asn361del	LAD-I ^p	2(4) ^b	[3, 46] Unpubl.	A0034 A0035 A0036 A0037 A0119
c.1083+5G>C ^d	Splice site	Deletion exon 9? p.Lys332_Asn361del?	LAD-I ^p	1(1)	Unpubl.	A0125
c.1143delC	Deletion	p.Tyr382ThrfsX9	LAD-I ^t	2(4) ^b	[30] Unpubl.	A0052 A0128
c.1225-?_1412+?del	Deletion or splice site	p.Ile409ValfsX1	LAD-I ^p	1(2) ^b	Unpubl.	A0132
c.1256_1257delAG ^d	Deletion	p.Glu419ValfsX27	LAD-I ^t	1(2)	[3, 5]	A0019

Nucleotide change	Mutation	Amino-acid or mRNA change	LAD-I type	Families (alleles)	Reference	Accession Number
c.1358G>A	Missense	p.Ser453Asn	LAD-I ^f	1(1) ^e	Unpubl.	A0112
c.1388_1390delGCTinsCA	Deletion/ insertion	p.Gly463AlafsX66	LAD-I ^f	1(2)	Unpubl.	A0109
c.1413-?_1877+?del2114	Deletion	Deletion exons 12 + 13 p.(Arg471Ser, Cys472_Ile626del)	LAD-I ^o	1(2)	[32, 36]	A0070 A0071
c.1413-149_2080+839 del2518insAAAA	Deletion/insertion	Deletion exon 12_14 p.Cys472ValfsX43	LAD-I ^o	1(2) ^b	Unpubl.	A0134
c.1413-396_?del27703	Deletion	Deletion exon 12 end p.Arg471SerfsX28	LAD-I ^o	1(1)	[24]	A0126
c.1421delC	Deletion	p.Thr474MetfsX55	LAD-I ^f	1(1)	Unpubl.	A0115
c.1498delG	Deletion	p.Asp500ThrfsX29	LAD-I ^o	5(7) ^b	[50, 51]	A0041 A0042 A0043 A0044 A0045 A0108
c.1590C>G	Nonsense	p.Tyr530X	LAD-I ^o	1(2) ^b	[3]	A0025
c.1602C>A	Nonsense	p.Cys534X	LAD-I ^o	4(6) ^b	[3, 22, 25, 44] Unpubl.	A0001 A0006 A0103 A0118
c.1622delGins23	Deletion/ insertion	p.Cys541TyrfsX8	LAD-I ^o	1(2)	Unpubl.	A0101
c.1622_1657del36 (cDNA)	Deletion? Splice site?	p.Cys541_Gly553del insTrp	LAD-I ^f	1(1)	[52]	A0040
c.1632C>G	Nonsense	p.Tyr544X	LAD-I ^o	2(3)	Unpubl.	A0096 A0120
c.1658-2A>G ^d	Splice site	Deletion exon 13? p.Gly553AlafsX6	LAD-I ^f	1(1)	Unpubl.	A0111
Not identified	Splice site	Deletion exon 13 p.Gly553AlafsX6	LAD-I ^o	1(1)	[3, 38]	
c.1670G>C	Missense	p.Cys557Ser	LAD-I ^o	1(2) ^b	Unpubl.	A0129
c.1768T>C	Missense	p.Cys590Arg	Activated LFA-1, no CR3	1(1)	[3, 22, 44]	A0007
c.1777C>T	Missense	p.Arg593Cys	Activated LFA-1, no CR3	5(6) ^b	[3, 22, 44, 50, 53]	A0008 A0009 A0012 A0045 A0066 A0067
c.1834T>C	Missense	p.Cys612Arg	LAD-I ^f	1(2)	[3, 26, 54]	A0082
c.1907delA	Deletion	p.Lys636ArgfsX22	LAD-I ^o	1(2)	[30]	A0061 A0062

Nucleotide change	Mutation	Amino-acid or mRNA change	LAD-I type	Families (alleles)	Reference	Accession Number
c.1920delG	Deletion	p.Lys641ArgfsX17	LAD-I ^o	1(1)	[50]	A0042
c.1943C>T	Missense	p.Pro648Leu	LAD-I ^o	1(1) ^e	Unpubl.	A0110 *
c.2070delT ^c	Deletion	p.Asp690GlufsX25	LAD-I ^o	6(6)	[3, 22, 23, 32, 35] Unpubl.	A0015 A0016 A0017 A0116 A0127
c.2077C>T	Nonsense	p.Arg693X	LAD-I ^o	1(2) ^b	Unpubl.	A0077 *
c.2080+1delG ^a	Splice site	Deletion exon 14 ? p.Ser627ValfsX44?	LAD-I ^o	1(1)	Unpubl.	A0090 *
c.2147G>C ^f	Missense	p.Gly716Ala	LAD-I ^o	1(2)	[30]	A0054 *
c.2200G>T	Nonsense	p.Glu734X	LAD-I ^o	1(1)	[52]	A0040 *

	Number of different alleles	Total number of alleles
Deletions	23 alleles (26.7%)	52 alleles (25.9%)
Insertions	2 alleles (2.3%)	2 alleles (1.0%)
Deletion/insertions	4 alleles (4.7%)	11 alleles (5.5%)
Nonsense mutations	10 alleles (11.6%)	23 alleles (11.4%)
Splice site mutations	12 alleles (14.0%)	31 alleles (15.4%)
Missense mutations	35 alleles (40.7%)	82 alleles (40.8%)
	Total 86 different allelic mutations	Total 109 families with 201 identified alleles in the 123 patients

Additional mutations mentioned in refs. 37 and 55-57 can be disregarded as being nonspecific for LAD-I, unconfirmed or printing errors.

^{o/-} indicates that the mutant CD18 molecule supports differentially the expression of the different CD11/CD18 integrins.

^a Positions of introns in *ITGB2*: intron 1 c.-4_-3; intron 2 c.58_59; intron 3 c.147_148; intron 4 c.328_329; intron 5 c.499_500; intron 6 c.741_742; intron 7 c.897_898; intron 8 c.993_994; intron 9 c.1083_1084; intron 10 c.1224_1225; intron 11 c.1412_1413; intron 12 c.1657_1658; intron 13 c.1877_1878; intron 14 c.2080_2081; intron 15 c.2247_2248.

^b One or more patients presumed to be homozygous for this mutation.

^c And reverse mutations in a small subset of lymphocytes [32].

^d Corrected after consultation of the authors.

^e No other mutation found in this patient; variation of unknown significance.

^f Not certain of clinical significance since transfection of this mutant into a CD18-deficient B cell line induced near normal expression. Moreover, transfection into HEK293 cells together with CD11a induced normal expression and constitutive binding to ICAM-1, and transfection together with CD11b induced expression and constitutive binding to denatured BSA.

Table 2Polymorphisms in the LAD-I gene *ITGB2*

Polymorphic nucleotide	Amino acid change	Reference
c.-403C/T (promoter)	NA	[3, 25]
c.-135A/C (5' UTR)	NA	[Ensembl]
c.-128A/T (5' UTR)	NA	[Ensembl]
c.-111T/C (5' UTR)	NA	[3, 22, 58, 59]
c.-110A/G (5' UTR)		
c.-10_-7AGTT/del	NA	[Ensembl]
c.13C/T	p.Arg5Cys	[Ensembl]
c.24G/T	p.Leu8Leu (silent)	[3, 16, 22, 24, 30, 58]
c.31C/T	p.Leu11Leu (silent)	[Ensembl]
c.58G/A	p.Val20Ile	[Ensembl]
c.117G/A	p.Ser39Ser (silent)	[Ensembl]
c.147+16A/G	NA	Unpubl.
c.162G/A	p.Pro54Pro (silent)	[3, 5, 22]
c.205_206CT/TC	p.Leu69Ser	[Ensembl]
c.222G/A	p.Ala74Ala (silent)	[Ensembl]
c.229G/A	p.Asp77Asn	[58]
c.294G/A ^a	p.Leu98Leu (silent)	[58]
c.328+15G/A	NA	[3, 22]
c.499+7C/T	NA	[3, 22]
c.500-103T/G	NA	Unpubl.
c.500-29T/C	NA	[24]
c.500-11G/T	NA	[3, 22, 24]
c.587A/C ^b	p.Lys196Thr	Unpubl. ^b
c.598C/G	p.Pro200Ala	[Ensembl]
c.741+36T/C	NA	Unpubl.
c.742-13G/A	NA	[3, 22]
c.810G/A	p.Ala270Ala (silent)	[Ensembl]
c.819A/G	p.Gly273Gly (silent)	[3, 5, 22, 25, 31, 38, 45, Ensembl]
c.849C/T	p.Asp283Asp (silent)	[Ensembl]
c.892G/A	p.Glu298Lys	[58]
c.906A/G	p.Pro302Pro (silent)	[Ensembl]
c.993+41T/C	NA	Unpubl.
c.994-47G/A ^a	NA	[3, 22, 24]
c.1002C/T	p.Thr334Thr (silent)	[Ensembl]
c.1062T/A	p.His354Gln	[30, 48]
c.1101C/A	p.Val367Val (silent)	[3, 22, 24, 25, 30, 37, 58]
c.1146C/T	p.Tyr382Tyr (silent)	[Ensembl]
c.1186C/T	p.Pro396Ser	[Ensembl]

Polymorphic nucleotide	Amino acid change	Reference
c.1224+48G/T	NA	[3, 22]
c.1247C/T	p.Thr416Met	[Ensembl]
C.1323C/T	p.Val441Val (silent)	[3, 5, 22, 24, 25, 30, 37, 58]
C.1497G ?A	p.Lys499Lys (silent)	[Ensembl]
C.1542C/T	p.Cys514Cys (silent)	[Ensembl]
C.1635C/T	p.Asn545Asn (silent)	[Ensembl]
C.1700G/C	p.Gly567Ala	[58]
C.1714G/T	p.Ala572Ser	[58]
C.1724G/C	p.Cys575Ser	[58]
C.1756C/T	p.Arg586Trp	[3, 4, 5, 58, 59]
C.1793G/T	p.Cys598Phe	[Ensembl]
C.1888G/A	p.Glu630Lys	[Ensembl]
C.1893C/T	p.Cys631Cys (silent)	[Ensembl]
C.2058C/G	p.Leu686Leu (silent)	[Ensembl]
C.2147G/C ^c	p.Gly716Ala	N. Parvaneh, unp. ^c
C.2349T/C (3' UTR) ^a	NA	[58]
C.2433A/G (3' UTR) ^a	NA	[3, 22, 58]
C.2456C/A (3' UTr) ^a	NA	[3, 22, 58]
C.2482G/C (3' UTR) ^a	NA	[58]
C.2484G/A (3' UTR)	NA	[Ensembl]
C.2579A/C (3' UTR) ^a	NA	[58]
C.2681G/T (3' UTR) ^a	NA	[58]
c.2721_2722delTG (3' UTR)	NA	[Ensembl]

^a Corrected after Consultation of the authors.

^b Originally published as an LAD-I⁻ mutation [53].

TABLE 3Mutations and polymorphisms in the LAD-II gene *SLC35C1*

Nucleotide change	Mutation	Amino-acid or mRNA change	LAD type	Families (alleles)	Reference	Accession Number	
c.439C>T	Missense	p.Arg147Cys	LAD-II	1(2)	[3, 7, 8]	A0001	
c.588delG	Deletion	p.Trp196CysfsX35	LAD-II	1(2)	[60]	A0004	*
c.923C>G	Missense	p.Thr308Arg	LAD-II	2(4)	[3, 8, 61]	A0002 A0003 A0005	
c.969G>A	Nonsense	p.Trp323X	LAD-II	1(2)	[62]	A0008	*
c.1010A>G	Missense	p.Tyr337Cys	LAD-II	1(2)	[63]	A0009	*
c.718A/G	SNP ^a	p.Ile240Val ^b			[61]		
c.772T/C	SNP ^a	p.Phe258Leu ^b			[61]		

^a May also be errors in the GenBank accession number AF323970 [61].

^b Corrected after consultation of the authors.

TABLE 4

Mutations in the LAD-III gene *FERMT3*

Nucleotide change	Mutation	Amino-acid or mRNA change	LAD type	Families (alleles)	Reference	Accession Number	
c.48G>A	Nonsense	p.Trp16X	LAD-III	1(2)	[64]	F0019 F0020	*
c.161-2A>C ^a	Splice site	p.Asn54ArgfsX142	LAD-III	1(2)	[65]	F0025	*
c.238_244dup7	Insertion	p.Lys82ThrfsX67	LAD-III	1(2)	Unpubl.	F0024	*
c.687G>A	Nonsense	p.Trp229X	LAD-III	1(2)	[12, 66]	F0014 F0015	*
c.922G>A	Missense	p.Gly308Arg	LAD-III	1(1)	[67]	F0021	*
c.1275delT	Deletion	p.Glu426ArgfsX3	LAD-III	1(1)	[67]	F0021	*
c.1525C>T	Nonsense	p.Arg509X	LAD-III	12(24)	[11, 12, 14, 64]	F0001 F0002 F0003 F0004 F0005 F0006 F0007 F0008 F0009 F0010 F0011 F0012 F0017 F0018	*
c.1671-2A>G ^a	Splice site	Deletion exon 14 p.Phe558TrpfsX141	LAD-III	1(2)	[14]	F0016	*
c.1717C>T	Nonsense	p.Arg573X	LAD-III	2(4)	[12, 68]	F0013 F0022 F0023	*

^aPosition of introns in *FERMT3*: intron 1 c.-15_-14; intron 2 c.160_161; intron 3 c.394_395; intron 4 c.514_515; intron 5 c.683_684; intron 6 c.786_787; intron 7 c.894_895; intron 8 c.1029_1030; intron 9 c.1079_1080; intron 10 c.1204_1205; intron 11 c.1311_1312; intron 12 c.1545_1546; intron 13 c.1670_1671; intron 14 c.1812_1813.