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Biallelic mutations in LAMA5 disrupts a skeletal noncanonical focal adhesion pathway and produces a distinct bent bone dysplasia

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ABSTRACT

Background: Beyond its structural role in the skeleton, the extracellular matrix (ECM), particularly basement membrane proteins, facilitates communication with intracellular signaling pathways and cell to cell interactions to control differentiation, proliferation, migration and survival. Alterations in extracellular proteins cause a number of skeletal disorders, yet the consequences of an abnormal ECM on cellular communication remains less well understood.

Methods: Clinical and radiographic examinations defined the phenotype in this unappreciated bent bone skeletal disorder. Exome analysis identified the genetic alteration, confirmed by Sanger sequencing. Quantitative PCR, western blot analyses, immunohistochemistry, luciferase assay for WNT signaling were employed to determine RNA, proteins levels and localization, and dissect out the underlying cell signaling abnormalities. Migration and wound healing assays examined cell migration properties.

Findings: This bent bone dysplasia resulted from biallelic mutations in LAMA5, the gene encoding the alpha-5 laminin basement membrane protein. This finding uncovered a mechanism of disease driven by ECM-cell interactions between alpha-5-containing laminins, and integrin-mediated focal adhesion signaling, particularly in cartilage. Loss of LAMA5 altered β1 integrin signaling through the non-canonical kinase PYK2 and the skeletal enriched SRC kinase, FYN. Loss of LAMA5 negatively impacted the actin cytoskeleton, vinculin localization, and WNT signaling.

Interpretation: This newly described mechanism revealed a LAMA5-β1 Integrin-PYK2-FYN focal adhesion complex that regulates skeletogenesis, impacted WNT signaling and, when dysregulated, produced a distinct skeletal disorder.

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1. Introduction

For proper intracellular function, cells need to interact with their extracellular matrix (ECM). The ECM provides a structural scaffold for the cell and aids in the maintenance of contacts between neighboring
Research in context

Evidence before this study

Prior to this work, there remains heritable disorders of the bone and cartilage, the osteochondrodysplasias, for which the molecular basis is unknown. This includes a group of disorders that are referred to as bent bone dysplasias in the recent nosology of the osteochondrodysplasias. Not all clinical and radiographic phenotypes fit into the well classified disorders for which the molecular basis has been established and there remain affected individuals with no basis for their disease.

Added value of this study

This study uncovers a newly described skeletal disorder associated with multiple skeletal defects including distinct bending of long bones, contractures, dysmorphic facies and lethality. Using exome analysis, biallelic mutations in the gene encoding LAMA5 co-segregated with the disorder. These mutations led to diminished LAMA5 expression including in cartilage. Immunohistochemistry showed that LAMA5 is well expressed in musculoskeletal tissues. Loss of LAMA5 led to decreased β1 integrin signaling through a noncanonical pathway that affected focal adhesion complexes, uncovering a network, LAMA5-β1 Integrin-PYK2-FYN. Further, diminished LAMA5 negatively impacted the actin cytoskeleton, vinculin localization, and WNT signaling, tying focal adhesion molecules to key skeletal signaling molecules.

Implications of all available evidence

This work uncovered a role for LAMA5 in regulating skeletogenesis and that LAMA5 mutations can produce a recessively inherited skeletal dysplasia. In chondrocytes, LAMA5 regulated -β1 integrin signaling in a cell context manner. This work further demonstrates the importance of basement membrane proteins and the role of β1 integrin signaling in the skeleton.

Cells. Dependent on the tissue type, cells produce an ECM that is primarily composed of two classes of macromolecules, fibrous proteins and proteoglycans. ECM fibrous proteins include collagens, fibronectins, elastins, and laminins [1]. While collagens are the most abundant fibrous ECM proteins [1], laminins are the major components of the basement membrane, a specialized ECM immediately surrounding the cell [2, 3]. Laminins are composed of α, β and γ polypeptide chains that self-assemble through their coiled coil domains into a T-shape configuration achieved by disulfide bonds along their long arms [4]. To date, five α, four β and three γ chains have been identified, and 16 distinct combinations of these chains have been delineated [4, 5]. Basement membranes provide cells with physical scaffolding and tensile strength and, through binding partners, participate in cellular deformation and homeostasis.

In addition to supplying physical support to cells, the ECM participates in biologic functions due to its ability to bind multiple interacting partners that include growth factors, signal receptors, and adhesion molecules [6]. Adhesion molecules are cell surface glycoproteins involved in cell-to-cell and cell-to-ECM interactions and include the following protein families: integrins, the immunoglobulin superfamily, selectins, and cadherins [7]. Among the adhesion molecules, integrins interact with ECM collagens, fibronectin, and laminins to generate a multi-protein focal adhesion complex required for communication between the ECM and the actin cytoskeleton [8-10]. Upon integrin signaling, talin and paxillin recruit the focal adhesion kinases (FAK or PYK2) and then the cytoskeletal protein vinculin induces clustering and maturation of these integrin focal points [11]. This clustering promotes the growth of focal adhesions through phosphorylation of other focal adhesion kinases with subsequent activation of the SRC family kinases and p130Cas [12]. These focal adhesions compose the integrin adhesome and are crucial in determining downstream signaling events, culminating in changes in cell fate and behavior.

Skeletal dysplasias include over 460 defined heritable disorders of bone and cartilage diagnosed through a combination of clinical, radiographic and molecular findings [13]. Bending of bones is a phenotypic finding in many of the well-defined skeletal disorders including campomelic dysplasia [14], osteogenesis imperfecta [15], hypophosphatasia, and bent bone dysplasia FGR2 [16], among others [17, 18]. Underlying genetic causes in these disorders are distinct and include mutations in genes that encode transcription factors, collagens, enzymes, and receptors. Herein, we delineate a previously undescribed bent bone skeletal dysplasia resulting from biallelic mutations in LAMA5, which encodes the α5 laminin subunit (LAMA5). The mutations resulted in reduced LAMA5 expression and alterations in the integrin focal adhesion pathway involving the non-canonical downstream kinases, PYK2 and FYN. Loss of LAMA5 also altered proper localization of cellular vinculin and diminished WNT signaling. These findings support the importance of proper focal adhesion induced signaling through LAMA5-integrin β1-PYK2-FYN signaling in skeletal development and identify a novel skeletal disorder resulting from disruption of the signaling pathway.

2. Results

2.1. Mutations in LAMA5 result in a new form of bent bone dysplasia

We identified three siblings from a single nonconsanguineous family (International Skeletal Dysplasia Registry reference number R03—206) with skeletal defects that did not phenotypically or radiographically match any of the currently classified skeletal disorders. Clinical findings included atrial septal defects, relative macrocephaly, micrognathia, proptosis, ear abnormalities, widely spaced nipples, elbow dislocations, and severe distal limb contractures (Table S1). Additionally, there was significant polyhydramnios in the prenatal period suggesting placental dysfunction or fetal inability to swallow properly. Radiographic findings included under-mineralization of the calvarium (Fig. 1a-b and S1), distinctly bent, under-mineralized long bones, particularly of the femora, with mid-diaphyseal angulation (Fig. 1c-e and S1), platyspondyly with coronal clefts (Fig. 1f-g and S1), and bilateral upper and lower extremity contractures (Fig. 1h-j and S1). The three affected individuals all delivered at term but died shortly after birth due to presumed pulmonary insufficiency, even though their hearts were not small. Detailed phenotypic, radiographic and histologic findings for each affected individual are described in Supplementary Table 1. The clinical and radiographic findings suggested an unclassified or previously undescribed skeletal disorder.

Exome sequence analyses on the 3 affected individuals, one unaffected sibling, and the parents (Fig. 2a) demonstrated that the affected individuals had biallelic mutations in LAMA5 [OMIM 601,033]. The affected siblings and the father (R03—206-E, -A, -B, -G) (Fig. 2b) were heterozygous for the missense variant, NG_050626.1: g.43161G>A (c.4213G>A), predicted to result in the amino acid substitution p.Ala1405Thr, corresponding to a region close to the central cluster of epithelial growth factor (EGF) domains within the protein (Fig. 2c). The g.43161G>A variant has an allele frequency of 0.0006 in the gnomAD database (https://gnomad.broadinstitute.org/) and has not been seen in the homozygous state. Further this alanine at position 1405 is either an alanine or glycine, among all vertebrates, and threonine is not one seen at position p.1405 in any species. Using the
Mutatntaster prediction program (http://www.mutationtaster.org),
the variant is presumed to be “disease causing” with a score of 0.57
by Bayes classifier, though it was classified as benign by SIFT (http://
sift.jcvi.org). The second variant, NG_050626.1:g.49646C>T
(c.6157C>T), was present in all affected siblings and the mother
(R03-206-D, -A, -B, -G) (Fig. 2b) and it is predicted to result in the
amino acid substitution, p.Arg2053Cys, affecting one of the central
EGF-like domains (Fig. 2c). This variant, g.49655C>T has an allele fre-
quency of 0.0003, also has not been reported in the homozygous
state, and is classified as “disease causing” with a score of 0.99 by
Bayes classifier (Mutationtaster prediction algorithm), though also
classified as benign by the SIFT program. This residue is conserved as
an arginine to c.elegans, though it is a similarly positively charged
histidine in r.norvegicus and g.gallus. One unaffected sibling R03-
206H, was a carrier of the c.4213G>A variant, predicting the p.
Ala1405Thr substitution. Because all three affected siblings were
compound heterozygotes for both mutations and each parent carried
of one of the variants, the inheritance pattern was consistent with an
autosomal recessive disorder. Both mutations affected the protein in
or near EGF-like domains involved in disulide bonding, which are
essential for proper folding and secretion of the mature protein
(Fig. 2c).

At the protein level, studies using primary chondrocytes, fibro-
basts and cartilage derived from one of the affected individuals
(R03-206G), showed statistically significantly decreased levels of
LAMA5, demonstrating that the mutations in LAMA5 destabilized the
protein (Fig. 2d-f), supporting the conclusion that decreased LAMA5
produced this unclassified bent bone dysplasia.

2.2. LAMA5 expression in the musculoskeletal tissues and colocalization
with focal adhesion components around skeletal blood vessels

The role of LAMA5 in the skeleton has not been well investigated.
However, recent evidence suggests that laminins containing the
alpha5 subunit participate in bone homeostasis through cell to
matrix interactions, particularly around type E blood vessels [19]. To
obtain a deeper understanding of the role played by LAMA5 in the
skeleton, we first used publicly available data to examine gene
expression in bone and cartilage tissues; LAMA5 showed broad
expression that included cartilage and bone (Figure S2a) [20, 21]. We
then used immunostaining of several human musculoskeletal tissues
with LAMA5 antibodies and demonstrated expression of the protein
around blood vessels in cartilage and ligament (Fig. 3a, c-d), as well
as in muscle, ligaments, periosteum, trabecular bone and throughout
the cartilage, particularly in the growth plate and in articular chon-
drocytes (Fig. 3a-g). LAMA5 was similarly distributed in mouse mus-
culoskeletal tissues (Figure S2B). CD34, a marker for the specialized
capillary subtype E that is strongly linked to osteogenesis through β1
integrin and LAMA5 interactions [19] showed a similar localization to
LAMA5 in the vascular system of cartilage and in ligaments and bone

Fig. 1. Radiographic findings in R03 -206 G. Full term fetus with enlarged skull (a), micrognathia (b), presence of scapula and elbow dislocations (c), slight bowing of the humerus,
radius and ulna (c, d), severely bent femurs with cortical abnormalities (e), platyspondyly with coronal clefts (f), aplastic acetabular roofs (g) ulnar deviation and contractures in the
hands (h) and bilateral equinovarus (i, j).
Because of the established role of LAMA5 in integrin signaling and focal adhesions, p130-CAS (Figure S2d), a downstream component of the integrin adhesion intracellular cascade was also interrogated and, similar to LAMA5, was expressed in blood vessels, articular cartilage, ligament, and trabecular bone. These results showed that there is a broad distribution of LAMA5 in musculoskeletal tissues, including tissue blood vessels, and that LAMA5 co-localized with p130-CAS, a component of the integrin-focal adhesion pathway.

2.3. LAMA5 mutations alter a non-canonical focal adhesion signaling pathway

Laminins interact with integrins on the cell surfaces of multiple cell lineages. More specifically, LAMA5 laminins have been shown to interact with integrin receptors containing the β1 subunit in specialized bone endothelial lineages supporting osteogenesis [19]. To further study the connection between LAMA5 and integrin signaling in the skeleton, we analyzed different components of the integrin-focal adhesion pathway in primary chondrocytes derived from control and affected individuals and in a gene-edited HeLa cell line, a cell line with a high expression of LAMA5. The HeLa cell mutation is a single nucleotide insertion (317insA) and results in alternative splicing in exon 2, deleting 51 amino acids in the short arm of LAMA5 (Canopy Biosciences, KOCE0011980). The mutation is a hypomorphic allele (LAMA5Hyp) which in the homozygous state reduced LAMA5 protein by 88% (Fig. 4B-D), similar to the levels found in the patient-derived tissues with biallelic LAMA5 mutations. In the R03/C0206 chondrocytes, the level of β1 integrin was decreased (Fig. 4a, e), suggesting the potential for a lack of proper integrin signaling. In the LAMA5Hyp/Hyp cells, not only was there decreased mature β1-integrin (Fig. 4b, f), but also a pattern consistent with accumulation of immature forms of the protein in the secretory pathway [22]. Activation of the integrin receptors by laminins usually initiates the formation of the focal adhesion complex by phosphorylation of Focal Adhesion Kinase (FAK), the main intracellular integrin signal transducer. FAK subsequently triggers phosphorylation of the SRC family kinases p130CAS and Paxillin in order to reorganize the actin cytoskeleton and decrease integrin turnover [23, 24]. Although protein levels of the β1-integrin receptors were altered in defective LAMA5 cells, levels of phosphorylated FAK were not altered in either patient chondrocytes or LAMA5Hyp/Hyp cells (Fig. 4a-f, b-f).

As canonical integrin signaling through FAK was not affected, we interrogated alternative components of the focal adhesion pathway.
Proline-rich tyrosine kinase 2 (PYK2) is a non-receptor tyrosine kinase member of the FAK family with a similar role, but its expression pattern is restricted to only a few tissues, among them bone and cartilage [25-27]. Like FAK, PYK2 is also established as a scaffold for the SRC family of kinases and its translocation to focal adhesions promotes its association with the focal adhesion proteins p130-CAS and Paxillin. Relative to controls, both phosphorylated and total PYK2 levels were decreased in bent bone LAMA5 and HeLa LAMA5Hyp/Hyp mutant cells (Fig. 4a-b and g-j). Further studies of the integrin focal adhesion complex also showed decreased levels in general downstream phosphorylation of p-P130Cas and p-Paxillin (Fig. 4a-b and k-n).

2.4. Noncanonical PYK2 and SRC family member FYN expression in musculoskeletal tissues

Since PYK2 expression was diminished, as well as the total phosphorylated SRC family (Fig. 5a-b and 53e-f), we sought to understand which SRC family kinase member(s) were highly expressed in bone and cartilage, two of the main tissues involved in this skeletal disorder. Publicly available transcriptional data for SRC family members FGR, FYN, YES1, LYN1, BLK, cSRC, LCK, HCK, and FRK, showed high levels of FYN expression in both cartilage and bone [21, 28]. Thus, we interrogated the LAMA5Hyp/Hyp HeLa cells, demonstrating that FYN levels were decreased compared to control (Fig. 5b-c). The data suggested that decreased FYN levels, correlated with decreased phosphorylation levels of general SRC resulted from changes in integrin signaling caused by deficient levels of LAMA5.

As both PYK2 and FYN levels were decreased in LAMA5Hyp/Hyp cells, we investigated their skeletal tissue localization relative to LAMA5 by immunohistochemistry. Very similar to LAMA5, PYK2 was localized in cartilage around blood vessels, in the articular cartilage, the cartilage growth plate, periosteum/perichondrium, ligaments and trabecular bone (Fig. 5d-j). Furthermore, FYN expression was comparable to PYK2 and LAMA5, with expression in articular cartilage, cartilage blood vessels, ligaments, perichondrium, and primary spongiosum, but with no expression in the proliferative and hypertrophic zones of the cartilage growth plate. (Fig. 5k-q). These findings show that key downstream signaling components in the β1 integrin cascade altered by loss of LAMA5 signaling, PYK2 and FYN, share overlapping expression in skeletal tissues, supporting a mechanism involving these molecules in producing this skeletal dysplasia.

Fig. 3. LAMA5 localization in the skeleton. (a) LAMA5 staining in a fetal 18 week musculoskeletal tissues. LAMA5 localized around blood vessels in cartilage, ligaments and bone (c-e) and independently of blood vessels in articular cartilage, periosteum and growth plate (b, g, f). Bars represent 50 μm.
Effects on cell adhesion due to loss of LAMA5

Laminin-integrin interactions activate the focal adhesion pathway, bringing vinculin (VCL) to the complex in order to connect the extracellular matrix to the actin cytoskeleton [29]. While loss of LAMA5 decreased expression of the β1 integrin focal adhesome proteins, no changes were detected in the total protein levels of VCL in LAMA5 mutant chondrocytes or LAMA5Hyp/Hyp HeLa cells (Fig 6a-d). Since the total amount of intracellular VCL was unaltered, we examined whether mutant LAMA5 dysregulated focal adhesion formation through VCL mislocalization by visualizing focal adhesions in LAMA5Hyp/Hyp cell cultures. Actin staining showed a disorganized cytoskeleton and there was a reduced number of VCL focal adhesions in LAMA5Hyp/Hyp cells (Fig. 6e-f). This reduction in focal adhesions was confirmed using functional assays of cell adhesion and proliferation.

Adhesion were measured using a wound healing assay. The assay showed a reduced rate of healing by 10.8% in cells with deficient LAMA5 compared to control cells (92.8 ± 1.6% vs. 82.8 ± 2.2%; p = 0.0013), indicating diminished adhesion properties (Fig. 6g). No changes in cell survival or proliferation were detected (Fig. 6h). These results confirmed a disruption in cellular focal adhesion properties in LAMA5 defective cells by altered cytoskeletal architecture, vinculin localization, and diminished cell healing.

Defective LAMA5 alters WNT signaling

Based on the radiographic phenotype that included decreased mineralization of the axial and appendicular bone, we investigated whether altered WNT signaling could contribute to the phenotype since loss of WNT signaling negatively impacts skeletal mineralization. The results showed a significant reduction in β1 integrin expression in both chondrocytes and HeLa cells from R03-206G compared to control cells, indicating diminished adhesion properties (Fig. 6a-d). Similar changes were observed in HeLa LAMA5Hyp/Hyp, confirming the disruption in cellular focal adhesion properties in LAMA5 defective cells by altered cytoskeletal architecture, vinculin localization, and diminished cell healing.
mineralization [30]. WNT signaling, both canonical and noncanonical, are active in many different cellular contexts to regulate development, growth, and maintenance. In canonical signaling, secreted WNT ligands bind to Frizzled receptors, leading to a larger receptor complex with co-receptors LRP5/6. This leads to the dissolution of a cytoplasmic β-catenin destruction complex consisting of DVL/APC/Axin/GSK3β/CK1α. Dissolution of the destruction complex allows for accumulation of cytoplasmic β-catenin, which then translocates to the nucleus to activate downstream targets (Figure S4). Src family kinases, including Src and FYN, have been implicated in canonical WNT signaling and have been shown to bind to LRP5/6, Frizzled and DVL2 in different contexts [31, 32] and FYN has been shown to directly stabilize β-catenin in osteoarthritic cartilage [33].

To investigate WNT signaling we analyzed the TCF/LEF promoter activity after WNT3A induction. Both patient chondrocytes and LAMA5Hyp/Hyp HeLa cells showed decreased TCF/LEF promoter activity (Fig. 7a-b). Analysis of the intracellular activation of WNT signaling after WNT3A treatment showed a reduction in phosphorylation of the WNT receptor LRP6 in patient chondrocytes at Ser 1490 (Fig. 7c-e). LAMA5Hyp/Hyp cells also showed a reduction in LRP6 phosphorylation after WNT3A treatment, similar to patient chondrocytes (Fig. 7f-h). These results indicate that defective expression of LAMA5 diminished activation of the WNT pathway likely through decreased active Src and FYN kinases.

3. Discussion

We identified biallelic mutations in LAMA5 that resulted in decreased Laminin α5 protein and produced a previously undescribed recessively inherited bent bone skeletal dysplasia. There have been two previous reports of LAMA5 variants in humans. In one report, heterozygosity for a LAMA5 mutation, c.9418G>A (p. V3140M), in one individual was associated with skin anomalies, muscle weakness, osteoarthritis, ligamentous laxity, and with other symptoms; however, although the variant altered a highly conserved residue, it was also found in unaffected individuals in the family [34]. In the other report, there was homozygosity for the LAMA5 variant c.8046C>T (p.Arg2659Trp) in an individual who presented with presynaptic myasthenic syndrome (hypotonia at birth, muscle weakness, respiratory failure requiring intermittent ventilation as a teen, ptosis, myopia, and facial tics). The authors showed that LAMA5 was expressed in nerve endplates and that the patient’s endplates were morphologically abnormal [35]. The substituted arginine is not fully conserved across species, and the affected individual was also homozygous for a variant in LAMA1 (p.Asp210Asn) that altered a highly conserved residue and is predicted to be damaging. The asymptomatic parents were heterozygous for both of these variants [35]. These reports illustrate the potential for defective LAMA5 to produce disease, though the phenotypes differ from each other and our findings.
The phenotype we identified was consistent among the three affected siblings and included defects in both the axial and appendicular skeleton, with radiographic findings showing under-mineralized bone and a distinct angulation of the mid femoral shaft (Fig. 1 and Table S1). The phenotype also included extraskeletal features. There were facial dysmorphisms, abnormally formed ears with tags, wide-spaced nipples, and atrial septal defects, among other findings (Table S1). This disorder also likely includes abnormalities in muscle function based on the presence of elbow fusions, ulnar flexion contractions at the wrist (arthrogryposis multiplex congenita), bilateral talipes equinovarus, and failure to mount a respiratory effort at birth. Immunohistochemistry generated in both human and mouse tissues showed that LAMA5 was expressed in muscle and it has been previously reported that LAMA5 deficient intestine displayed a smooth muscle defect [36]. Laminins, type IV collagen, agrin, nidogen, biglycan, and perlecan form the basal lamina that surrounds skeletal muscle fibers, and it has been suggested that laminin in the ECM stimulates myoblast proliferation and differentiation [37, 38]. Thus, the potential for abnormal muscular function due to loss of LAMA5 is of particular interest in this disorder since muscle defects have also been speculated to contribute to bending of appendicular bones by altered mechanical forces during development [38].

Skeletal growth, in both the prenatal and postnatal periods, requires a series of tightly coordinated processes of cell proliferation, differentiation, migration of cells, mineralization, and expansion of the local vasculature. Both chondrocytes and osteoblasts contribute to angiogenesis by release of vascular endothelial growth factor (VEGF), furthering vessel formation through activation of VEGF receptors in endothelial cells. Blood vessels provide nutrients, oxygen, growth factors and hormones, as well as attracting progenitor cells. Type H vessels are a specialized subtype of capillary that expresses high levels of CD31 and endomucin, and these vessels are located near the growth plate in the metaphysis, the periosteum, and the endosteum of the diaphysis. Type H vessels are densely surrounded by osteoprogenitors [39] and aid in modulating osteoblasts, coupling osteogenesis to angiogenesis. There is also an embryonically expressed vessel termed Type E, with a similar expression pattern to type H vessels, but in mice they are highly expressed at E16.5 and then exhibit diminished expression by P28 [19]. Both Type E and Type H vessel endothelial cells have enriched expression of LAMA4, LAMA5, and β1 Integrin, as well as extracellular matrix proteins, basement membrane and cell adhesion components [19]. Our immunohistochemistry findings expand on the role of LAMA5 expression in musculoskeletal development beyond the...
previously described Type E capillaries, with expression in blood vessels in cartilage and ligament and expression in muscle, ligaments, periosteum, trabecular bone and throughout the cartilage, particularly in the growth plate and articular chondrocytes.

The functionality of laminins in general, including LAMA5, is mediated through interactions with cell membrane receptors; the most established LAMA5-receptor interaction is with β1 Integrin. This study uncovered that the LAMA5-β1 Integrin interaction was

Fig. 7. Defective LAMA5 alters WNT signaling pathway. (a) TCF/LEF promoter activity in R03-206G patient chondrocytes when induced with 100 ng/ml WNT3A for 30 min. (b) Similar changes were observed in LAMA5Hyp/Hyp cells after application of 100 ng/ml WNT3A. (c-h) Defective LAMA5 cells showed reduced LRP6 phosphorylation after a 30-minute incubation with 50 ng/ml WNT3A. *p<0.05, **p<0.01, ****p<0.0001.
altered by mutation and employed a non-canonical focal adhesion signaling pathway and the results are summarized in Table 2. Focal adhesion signaling typically transduces integrin intracellular signals through phosphorylation of FAK [40, 41]. This was not the case for signaling initiated by LAMA5, which induced the phosphorylation of PYK2. PYK2 belongs to the same family as FAK and has similar functions in cell migration, differentiation, and survival [42]. PYK2 has been described as an important factor in osteogenesis with roles in promoting osteoblast differentiation [27, 43-46], regulating osteoclastogenesis and bone remodeling [47-50] and modulating chondrogenesis [51-53]. Our study provides novel data that LAMA5 is required for PYK2 activation in the skeleton. Our results further show that the most relevant SRC family member downstream of PYK2 is FYN showing a similar skeletal localization to LAMA5. The importance of FYN in the skeleton is also supported by several studies that implicate FYN in osteoblast differentiation [54, 55], osteoclastogenesis [56, 57], and in cartilage differentiation/proliferation [58] and osteoarthritides, which is associated with an upregulation of FYN [33].

This skeletal specific signaling of PYK2 and FYN also suggests interactions of these kinases with important differentiation signaling pathways. Although PYK2 and FAK are very similar, they exhibit different effects on cellular behavior, likely dependent on tissue type. FAK activation causes cell spreading and pro-survival, while PYK2 induces reorganization of the cytoskeleton, cell detachment, and apoptosis [59]. PYK2 is known to stimulate multiple oncogenic signaling pathways including WNT [60, 61]. High levels of PYK2 levels are associated with poor cancer prognosis, in part through activation of WNT signaling [62]. Our findings show that diminished WNT3A-stimulated WNT signaling was seen in LAMA5 deficient cells through decreased LRPR receptor levels with decreased responsiveness of the TCF/LEF signaling initiated through LAMA5-Hyp/Hyp cells negatively impacted the organization of the actin cytoskeleton, vinculin localization, an important stabilizer of the focal adhesion complex, and cell migration, illustrating that, in part, the phenotype likely results from altered focal adhesions. The findings also support that, in the skeleton, a novel ECM network is induced through LAMA5-β1 Integrin-PYK2-FYN (Figure S4) signaling, impacting WNT signaling, and underscoring the importance of base-ment membrane proteins in skeletogenesis beyond their structural support role to the cell.

4. Materials and methods

4.1. Ethics

This study was approved by the University of California at Los Angeles Institutional Review Board under protocol number IRB#14-00017. Informed consent was obtained from all participants.

4.2. Study design

Patients and their unaffected family members were ascertained under an IRB-approved human subject’s protocol. Clinical information and imaging were obtained from review of all available medical records. All experiments performed in cells were conducted at least three independent times and all replicates were included in our data analyses.

4.3. Cell culture

Dermal fibroblast cultures were established from explanted skin biopsies from one of the individuals with mutations in LAMA5 (International Skeletal Dysplasia Registry reference number R03–206G) and three controls. Primary chondrocytes were isolated from distal femurs of the same affected individual (R03–206G) and six age-matched normal controls by incubation of fragmented cartilage with 0.03% bacterial collagenase II. LAMA5Hyp/Hyp Hela cells generated by CRISPR/Cas9 and controls were purchased from Canopy Biosciences (catalog no. KOCE01980). All cells were grown in Dulbecco-Vogt Modified Eagle Medium (DMEM) supplemented with 10% fetal bovine serum (FBS).

All cells were grown to 100% confluence for at least 24 h. Stimulation experiments were conducted in DMEM with 10% FBS with either TGFβ-1 (catalog no. 240-B, R&D Biosystems, 5 ng/ml with Ascorbic Acid, 50 ng/ml) or WNT3A (catalog no. 5036-WN, R&D Biosystems, 100 ng/ml). For protein analyses, cells were collected in IP Lysis Buffer (Thermo Scientific, 87,787) supplemented with proteinase inhibitors.

4.4. Exome analysis

DNA was isolated and submitted to the University of Washington Center for Mendelian Genomics for library preparation and exome sequencing. The samples were barcoded, captured using the Nimble-Gen SeqCap EZ Exome Library v2.0 probe library targeting 36.5 Mb of genome, and sequenced on the Illumina GAIIx platform with 50 bp bidirectional reads. Novealign was used to align the sequencing data to the human reference genome (NCBI build 37) and the Genome Analysis Toolkit (GATK) was used for post-processing and variant calling according to GATK Best Practices recommendations. For each sample, at least 90% of targeted bases were covered by at least 10 independent reads. Variants were filtered against NIEHS EGP exome samples (v.0.0.8), exomes from the NHLBI Exome Sequencing Project (ESP6500), 1000 genomes (release 3.20012430), and in-house exome samples. Mutations were further compared with known disease-causing mutations in HGD (2012.2). Variants were annotated using VAX, and mutation pathogenicity was predicted using the programs Polyphen, Sift, Condel, CADD, and MutationTaster (http://www.mutationtaster.org/). Potential disease-associated variants were identified under an autosomal recessive model, identifying either homozygosity for variants in one gene or compound heterozygosity for two variants in the same gene. The mutations reported in this work were confirmed by bidirectional Sanger sequencing of amplified DNA from the proband, siblings and the parents. Sequence trace files were aligned and analyzed using 4peaks.

4.5. Western blot

For Western blot analyses, protein lysates were separated by electrophoresis on 5%, 10%, or gradient (4 to 20%), catalog no. 4,568,093, Bio-Rad) SDS-polyacrylamide gels, transferred to polyvinylidene difluoride membranes, blocked in 5% milk, and probed overnight with primary antibody [anti-LAMA5 antibody (1:500; catalog no. C352437, LifeSpan Biosciences, Inc.), anti-Integrin β1 (1:500; catalog no. AB1952, Chemicon), anti-phospho-FAK (1:1000; catalog no. 8556, Cell Signaling Technology), anti-phospho-Pyk2 (1:1000; catalog no. 3291, Cell Signaling Technology), anti-Pyk2 (1:1000; catalog no. 3292, Cell Signaling Technology), anti-phospho-Src Family (1:1000; catalog no. 6943, Cell Signaling Technology), anti-phospho-p130 Cas (1:1000; catalog no. 4011, Cell Signaling Technology), anti-p130 Cas (1:1000; catalog no. 610,271, BD Biosciences), anti-Vinculin (1:10,000; catalog no. 129,002, Abcam), anti-phospho-Paxillin...
(1:1000; catalog no. 2541, Cell Signaling Technology), anti-phospho-
LRP6 (1:1000; catalog no. 2568, Cell Signaling Technology), anti-LRP6
(1:1000; catalog no. 3395, Cell Signaling Technology), anti-FYN
(1:1000; catalog no. 4023, Cell Signaling Technology), anti-GAPDH
(1:2000; catalog no. 2118, Cell Signaling Technology), anti-β-Actin
(1:1000; catalog no. 4967, Cell Signaling Technology), or anti-
β-Tubulin (1:1000; catalog no. 2128, Cell Signaling Technology).

Peroxidase-conjugated secondary antibodies (1:2000, catalog nos.
7071 and 7072, Cell Signaling Technology) were used, and immuno-
complexes were identified using the ECL (enhanced chemilumines-
cence) Detection Reagent (catalog no. 7003, Cell Signaling Technology).
Fiji was used to quantify bands after gel analysis recommenda-
tions from ImageJ and Gassmann et al. (http://rspb.info.nih.gov/
ij/docs/menus/analyze.html#gels), and the Student’s t-test was per-
formed for statistical analysis using Prism software. Experiments were
repliacted at least three times to perform statistical analysis.

4.6. Histological analyses and immunolocalization

For histology and immunocytochemistry, human (control) and
mouse tissues (E18.5) were decalciﬁed using unimuno decalcifica-
tion solution (catalog no. 1414–1, StatLab), and then parafﬁn-embed-
ded. Parafﬁn blocks were sectioned at 10 μm.

For immunohistochemistry, parafﬁn sections were heated to 48 °C
for 60 min in 0.2 M Sodium Citrate Buffer (pH 3.5). Primary antibodies
used were: anti-LAMA5 (1:100; catalog no. C352437, LifeSpan Bio-
sciences, Inc), anti-CD34 (1:100; catalog no. HPAP036722, Prestige
Antibodies), anti-p130[Cas] (1:100; catalog no. 610,271, BD Biosciences,
sin) anti-PYK2 (1:100; catalog no. 17,592–1-AP, Proteintech) and
anti-FYN (1:2000; catalog no. 7003, Cell Signaling Technology). Sections
were then incubated for 30 min in either R.T.U Biotinylated Goat
Anti-Mouse IgG Antibody or R.T.U Biotinylated Goat Anti-Rabbit IgG
Antibody (Vector Laboratories, catalog no. BP-9100 and BP-9200.
Anti-Mouse IgG Antibody or R.T.U Biotinylated Goat Anti-Rabbit IgG
Antibody (Vector Laboratories, catalog no. BP-9100 and BP-9200.
Antibody (Vector Laboratories, catalog no. BP-9100 and BP-9200.
Secondary antibodies were used: Alexa Fluor goat anti-mouse 488 and
Alexa Fluor goat anti-rabbit 568 for all samples), and DAPI at a 1:1000 dilution
for 1 hour. The primary antibody was incubated overnight at 4 °C.

After a subsequent 30-minute incubation in Streptavidin Horseradish
Peroxidase R.T.U. (Vector Laboratories, catalog no. 5704), the sections
were stained using the ImmPACT DAB Peroxidase Substrate Kit (Vec-
tor Laboratories, catalog no. SK-4105).

Immunofluorescence experiments were performed using a Carl
Zeiss LSM 700 laser scanning confocal microscope. Cultured cells
were ﬁxed in 4% PFA in PBS, then washed and permeabilized with
0.1% Triton X100 for 5 min, followed by blocking in 10% goat serum
for 1 hour. The primary antibody was incubated overnight at 4 °C.
Fluorescent secondary antibody was incubated at a 1:1000 dilution
for 1 hour at room temperature (alexa-fluor goat anti-mouse 488 and
goat anti-rabbit 568 for all samples), and DAPI at a 1:1000 dilution
for 5 min at room temperature was applied before mounting.

For focal adhesion imaging, Hela cells were plated on glass cover-
slips, grown overnight and ﬁxed using 4% paraformaldehyde for
15 min. Polymerized F-actin was visualized using Alexa Fluor
594–conjugated Phalloidin (Life Technologies), and vinculin was
stained with vinculin–ﬂuorescin isothiocyanate antibody (Sigma-
Aldrich) according to the manufacturer’s protocols.

4.7. Cartilage studies

Cartilage studies for protein levels were performed using lysates
from frozen distal femur cartilage from proband R03-206G. Briefly,
frozen cartilage was pulverized in liquid nitrogen and lysates were
preserved in Proteinase and Phosphatase inhibitors (Thermo Scientiﬁc
78440). Western blots were performed as described above.

4.8. Luciferase reporter assay

The TCF/LEF promoter reporter was purchased from Lenti-Cignal,
Qiagen. The Renilla control was used to normalize the luciferase sig-
nal (Lenti CIGNAL, Qiagen). Cells were transduced at 80% confluence.

When cells reached confluence, they were treated with 100 ng/ml
WNT3A, (RnD System) and incubated overnight. Then, cells were
lysed and luciferase activity was determined using a Dual-Luciferase
Reporter Assay (Promega).

4.9. RNA extraction and qPCR

RNA was extracted from primary chondrocytes and Hela cells
using TRIzol reagent (catalog no. 15-596-018, Life Technologies).
Complementary DNA (cDNA) was prepared from 1 μg of RNA using
ReverseAid First strand cDNA synthesis kit (Thermo Fisher Scientiﬁc)
and ampliﬁed using Maxima SYBR Green/ROX qPCR Master Mix (cat-
alog no. 11762100, Thermo Fisher Scientiﬁc). Gene expression
was calculated using the 2ΔΔCT method of analysis against the stable
housekeeping gene β-2-microglobulin (β2M). Three experimental
replicates were performed with three technical replicates. qPCR pri-
mers were: β2M: Fw: TGACTTTGTCAACGCCCCAG and Rev: AGCAAG-
CAAGCAAGATTGG. LAMA5: Fw: CGAGGACCTTACTGCAAGC and
Rev: GGTGACGTGACCTCGTGT.

4.10. Migration and wound healing assays

For the wound healing assay, 250,000 Hela cells were plated in
each well of 24-well plates. The next day, a single wound was intro-
duced to each well using a 20 μl ﬁlter tip scratch; the wells were then
washed with PBS to remove detached cells and given 1 ml of
fresh medium. The wounds were monitored in an automated incuba-
tion microscope BioStation CT (Nikon) in 1-hour intervals for 24 h.
Wound widths were measured in ImageJ (https://imagej.nih.gov/ij/
) using three measurements per image that were averaged to obtain
a single value for each time point. Only wounds with initial widths
of 700–900 μm were included in each analysis.

For the cell tracking migration assay, 4000 Hela cells were plated in
24-well plates. Starting the next day, the cells were monitored in an
automated incubation microscope BioStation CT in 5–10-minute
intervals for 24 h. The trajectories were manually tracked using the
ImageJ Manual Tracking plugin (https://imagej.nih.gov/ij/plugins/
track/track.html), with the beginning and end of the measurement
represented by two cell divisions. The length of the cell trajectories and
the spatial (euclidean) distance between their beginnings and
ends were analyzed using the Chemotaxis and Migration Tool 2.0
(ibidi).

4.11. Statistical analysis

GraphPad Prism was used for statistical analysis. All values are
means ± SEM, as indicated in the Figure Legends. All comparisons in
the study were performed using the Student’s t-test.

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Contributions

M.B., F.C., M.K.B., J.H.M., J.Z., I.D. performed the molecular experi-
ments. W.Z., S.P.T, J.X.C., M.B., D.N. performed and were responsible

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for the genomic analyses. D.H.C., R.S.L., and D.K. performed the clinical and radiographic assessment. D.H.C., P.K.D., and D.K. conceived the study design. M.B., I.D., D.H.C., P.K., and D.K. wrote the paper. The authors read and approved the final version of the manuscript.

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Declaration of Competing Interest

The authors have no financial or personal interests to disclose.

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Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.16116/ebiomed.2020.103075.

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