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ORIGINAL ARTICLE



Lower limb bone geometry in adult individuals with X-linked hypophosphatemia: an observational study

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Abstract

Summary We assessed lower-limb geometry in adults with X-linked hypophosphatemia (XLH) and controls. We found large differences in multiple measures including femoral and tibial torsion, bowing and cross-sectional area and acetabular version and coverage which may contribute to clinical problems such as osteoarthritis, fractures and altered gait common in XLH. **Purpose** Individuals with X-linked hypophosphatemia (XLH) are at risk of lower-limb deformities and early onset of osteoarthritis. These two factors may be linked, as altered biomechanics is a risk factor for osteoarthritis. This exploratory evaluation aims at providing clues and concepts for this association to facilitate future larger-scale and longitudinal studies on that aspect.

Methods For this observational study, 13 patients with XLH, aged 18–65 years (6 female), were compared with sex-, age- and weight-matched healthy individuals at a single German research centre. Femoral and hip joint geometry, including femoral and tibial torsion and femoral and tibial shaft bowing, bone cross-sectional area (CSA) and acetabular version and coverage were measured from magnetic resonance imaging (MRI) scans.

Results Total femoral torsion was 29° lower in individuals with XLH than in controls (p < 0.001), mainly resulting from lower intertrochanteric torsion (ITT) (p < 0.001). Femoral lateral and frontal bowing, tibial frontal bowing, mechanical axis, femoral mechanical–anatomical angle, acetabular version and acetabular coverage were all greater and tibial torsion lower in individuals with XLH as compared to controls (all p < 0.05). Greater femoral total and marrow cavity CSA, greater tibial marrow cavity CSA and lower cortical CSA were observed in XLH (all p < 0.05).

Discussion We observed large differences in clinically relevant measures of tibia and particularly femur bone geometry in individuals with XLH compared to controls. These differences may plausibly contribute to clinical manifestations of XLH such as early-onset osteoarthritis, pseudofractures and altered gait and therefore should be considered when planning corrective surgeries.

Keywords Bone · Femur · Geometry · Shape · XLH

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Introduction

X-linked hypophosphatemia (XLH) is a rare genetic disorder with an assumed incidence rate of 1 in 20,000 [1] caused by pathogenic variants in the phosphate regulating endopeptidase homolog, X-linked (PHEX) gene located at Xp22.1 [2]. The pattern of inheritance is X-chromosomal dominant with supposed complete penetrance. However, family history is negative in about one-third of cases, suggesting a relevant proportion of de novo mutations or an asymptomatic/undiagnosed parental status [3, 4]. By still not completely elucidated mechanisms, pathogenic, loss-of-function variants in PHEX lead to elevated levels of the phosphatonin fibroblast



growth factor 23 (FGF-23). Resulting renal phosphate wasting and compromised production of 1,25 dihydroxyvitamin D along with further disease mechanisms cause a wide spectrum of clinical manifestations [5]. Specifically, skeletal issues including short stature and limb deformities, joint problems and early-onset osteoarthritis are a universal finding in almost all patients [2, 6-9]. Common deformities of the weight-bearing lower limbs in XLH comprise abnormal diaphyseal bowing and torsion including unphysiological femoral torsion (alternatively known as femoral neck anteversion or FNA) [7, 10]. These deformities have clinical relevance, as differences in joint geometry likely contribute to increased risk of (pseudo-)fractures [11-13] and impaired joint congruity both directly through altered biomechanics and indirectly through effects on muscle tractive forces and gait in individuals with XLH [8, 14]. In addition to direct effects of PHEX deficiency and elevated FGF-23 on mineral metabolism and skeletal development, other disease-associated manifestations such as delayed motor development [9, 15] and greater incidence of childhood obesity in children with XLH [16] likely contribute to differences in joint development and wearing [17, 18]. Moreover, hip joint geometry has been identified to have predictive value with regards to developing osteoarthritis independent of clinical risk factors [19] and recent analyses highlight that specifically femoral torsion, on the one hand, is influenced by mechanical loading and, on the other, can contribute to osteoarthritis development [13].

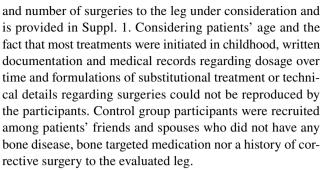
While the incidence of skeletal deformities in XLH has been reported according to clinical assessment thresholds or by self-report [9, 20], an objective characterization of bone geometry in adults with XLH has not been completed. Therefore, we assessed multiple components of lower-limb geometry in individuals with XLH, and compared them to those in an age- and sex-matched control group. We hypothesized that we would observe altered joint geometry in individuals with XLH.

Methods

Cohort

Evaluations are based on data obtained from an exploratory clinical study comparing 13 patients with X-linked hypophosphatemia (XLH) and 13 unaffected, age-, sex- and weight-matched controls. Per selection criterion, participants were 18–65 years old and did not have any acute disorders or conditions that would interfere with planned assessments.

For all XLH subjects, genetic confirmation of the diagnosis was available. Data regarding their previous medical and surgical treatment was based on medical history. Accessible information included age at initiation of medical treatment



While the core study is intended to comparatively evaluate muscle fatigue and physical performance in XLH vs healthy controls, data presented here is dedicated to a subanalysis of an MRI data-based evaluation of leg and hip geometry. Height and body mass were recorded.

Magnetic resonance imaging scan parameters

In this study, 5 sets of 64 transverse images each were recorded from the right leg and body side reaching at least from the foot to the lower part of the lumbar spine using a 3-T Siemens Biograph. Images were recorded using a turbo 2 echo DIXON sequence with the following parameters: flip angle 10°, TR 7.02 ms, TE1 2.46 ms, TE2 3.69 ms, 300 mm × 300 mm field of view at 1.2 mm × 1.2 mm × 4 mm voxel size. In this study, only images covering the region of the femur and tibia were evaluated.

Image analysis

MRI images were analysed using ImageJ (Version1.52 h). The total femoral torsion was defined using the femoral neck line and the posterior condylar line. The femoral neck line was defined as the line between the femoral head centre, and the shaft centre directly distal to the lesser trochanter. The posterior condylar line was defined as the line joining the posterior apices of the condyles in the image where the condyles were most prominent.

The inter-trochanteric torsion (ITT) was defined as the angle between the femoral neck line and the lesser trochanter line. The lesser trochanter line was defined as the line between the apex of the lesser trochanter at peak lesser trochanter size and the shaft centre. The shaft torsion (ST) is the angle between the lesser trochanter line and the posterior distal shaft line, which is aligned with the flat posterior distal shaft surface immediately proximal to the posterior condyles. The condylar torsion (CT) is the angle between the posterior distal shaft line and the posterior condylar line (Fig. 1). Tibial torsion was measured by the angle of the posterior tibial condylar line as reference and the line connecting the medial and lateral malleoli. The angle will be negative when the distal tibia rotates externally respective to the proximal tibia.



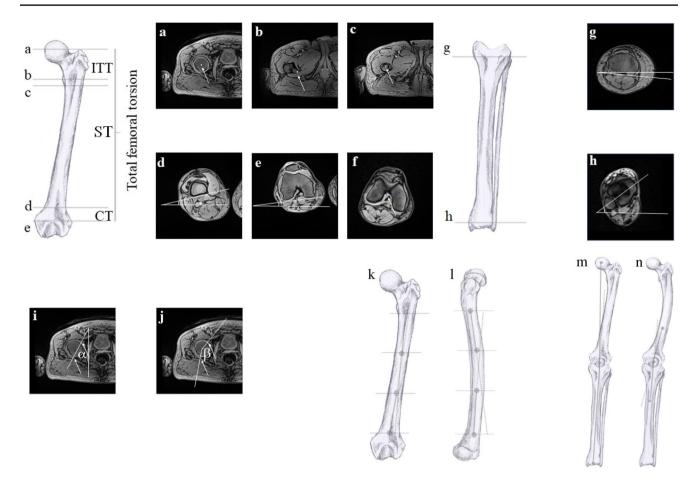


Fig. 1 Angle definitions: **a** femoral head centre, **b** apex of lesser trochanter, **c** shaft centre, **d** posterior distal shaft line, **e** posterior condylar line, **f** intercondylar notch, **g** posterior tibial condylar line and **h** lateral and medial malleoli. **i** Acetabular version α . **j** Acetabular coverage angle β . The slice where the circumference of the femoral

head was largest was chosen for analysis. The arrows show the rim of the acetabulum. **k** Femoral lateral bowing landmarks. **l** Femoral frontal bowing landmarks. **m** Mechanical axis landmarks. **n** femorotibial angle landmarks. Illustrations by X. O'Reilly-Berkeley

To measure femoral lateral and frontal bowing, the shaft centre was identified at four landmarks: immediately distal to the lesser trochanter, at 1/3 proximal—distal femoral shaft length; at 2/3 proximal—distal femoral shaft length (shaft is between slice "c" and "d" in Fig. 1) and at the femoral intercondylar notch. Likewise, tibial lateral bowing and frontal bowing were defined using the shaft centre at the intercondylar notch, the proximal third of the tibia and the distal third of the tibia at the distal tibial plateau.

The two lines used to define the mechanical axis are those connecting the femoral head centre to the femoral intercondylar notch, and from the midpoint of the tibial intercondylar eminence to the midpoint of the distal tibial epiphysis. The two lines defining the femorotibial angle were those connecting the femoral shaft centre at the distal third of the femur and the intercondylar notch, and the intercondylar notch and the proximal third of the tibia. Lastly, the femoral mechanical—anatomical angle is defined by the lines connecting the femoral head centre to the intercondylar notch and the

intercondylar notch to the centre of the femur immediately distal to the lesser trochanter.

The acetabular version was assessed in the cross section where the femoral head was greatest. It was defined as the angle between the anterior and posterior acetabular margins of the acetabulum and the sagittal plane (Fig. 1). On the same slice, the acetabular coverage was assessed as the angle between the posterior margin of the acetabulum and the femoral head centre and the anterior margin of the acetabulum and the femoral head centre.

To calculate the relevant angle λ in the frontal plane, the coordinates of the desired landmarks were calculated using the formula:

$$\lambda = \arctan\left(\frac{\frac{ay - by}{ax - bx} - \frac{cy - dy}{cx - dx}}{1 - \frac{ay - by}{ax - bx} + \frac{cy - dy}{cx - dx}}\right)$$



where a and b represent the respectively the proximal and distal reference points of the proximal axis and c and d the proximal and distal reference points of the distal axis. b and c overlap in the case of the mechanical axis and femorotibial angle (Fig. 1). y and x on the other hand are respectively the antero-posterior coordinates and the medio-lateral coordinates.

Statistical analysis

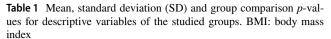
Femoral and tibial geometry, femorotibial alignment and acetabular parameters were compared in individuals with XLH and in control participants. Data from both groups were tested for normal distribution with the Shapiro-Wilk test. Parametric data were expressed as mean \pm SD, non-parametric data as median and interquartile range (IQR).

For parametric data, homogeneity of variance was checked with Levene's test and the Student t-test was implemented to assess group differences. For non-parametric data, the Mann-Whitney U test was applied to test for group differences. Repeatability of the measures performed has been calculated using SPSS statistical package version 23 (SPSS Inc, Chicago, IL) by the intraclass correlation coefficient (ICC). It was calculated based on a mean rating of the 3-image analysis as well as absolute effects, using a 2-way mixed-effect model to assess the repeatability of the image analysis by the same examiner at three different times in random order [21]. ICC was found to be excellent for total femoral torsion (ICC: 0.989), ITT (ICC: 0.968), femoral lateral bowing (ICC: 0.978), femoral frontal bowing (ICC: 0.904), tibial torsion (ICC: 0.93), tibial lateral bowing (ICC: 0.938), mechanical axis (ICC: 0.958) and femoral mechanical-anatomical angle (ICC: 0.974). Values for CT (ICC: 0.843), tibial frontal bowing (ICC: 0.768), acetabular version (ICC: 0.859) and acetabular coverage (ICC: 0.855) showed good reliability.

Results

Out of 13 subjects enrolled per group, whole-leg MRI imaging could be evaluated from 11 individuals with XLH (mean age 48.7 ± 9.2 years, 6 female, 77.3 kg) and 13 age-, sexand weight-matched controls (mean age 48.8 ± 7.6 years, 6 female, 79.9 kg, p > 0.677). The mean height of the XLH individuals was 1.56 m, and the control group height was 1.69 m (Table 1). For two male patients in the XLH group, MRI data could not be analysed due to extensive artefacts from metal implants.

Both basic characteristics and each of the measured angles were normally distributed apart from tibial frontal bowing. Individuals with XLH and control participants had



Variable	XLH (n 5 male/6	,	Control 7 male/6	t-test			
	Mean	SD	Mean	SD	p-value		
Age (years)	48.7	9.2	48.8	7.6	0.995		
Height (m)	1.56	0.10	1.68	0.09	0.006		
Weight (kg)	77.3	16.5	79.9	13.2	0.677		
BMI (kg/m²)	32.0	8.5	28.4	4.6	0.223		

similar age, body mass and BMI although control participants were taller than individuals with XLH (p = 0.006).

The total femoral torsion in the XLH group was $8.4\pm20.2^{\circ}$ and thus significantly lower than in the control group with $37\pm14.2^{\circ}$ (p<0.001; Fig. 2). When evaluating regional torsion on different levels along the femoral axis, group differences in femoral torsion appeared to primarily result from significantly lower inter-trochanteric torsion (ITT) in the XLH group with $36.4\pm12.3^{\circ}$ as compared to the control group with $54.9\pm9.5^{\circ}$ (p<0.001), as well as a lower condylar torsion (CT) in in XLH vs control with $1.5\pm5.3^{\circ}$ vs $7.2\pm2.9^{\circ}$, respectively (p<0.005; Fig. 2). In contrast, the shaft rotation or shaft torsion (ST) was similar in both XLH patients ($-29.5\pm17.9^{\circ}$) and controls ($-26.3\pm12.05^{\circ}$) (p=0.620). The tibia was less externally rotated in individuals with XLH $-8.7\pm13.5^{\circ}$ than in the control group $-37.2\pm11.8^{\circ}$ (p<0.001).

Substantial between-group differences were also evident regarding limb bowing. Femoral bowing was significantly enhanced in individuals with XLH vs controls for both lateral bowing with $13.1\pm7.0^{\circ}$ vs $-1.0\pm2.5^{\circ}$ (p<0.001), as well as frontal bowing with $31.4\pm7.3^{\circ}$ vs $17.8\pm1.4^{\circ}$ (p<0.001), respectively.

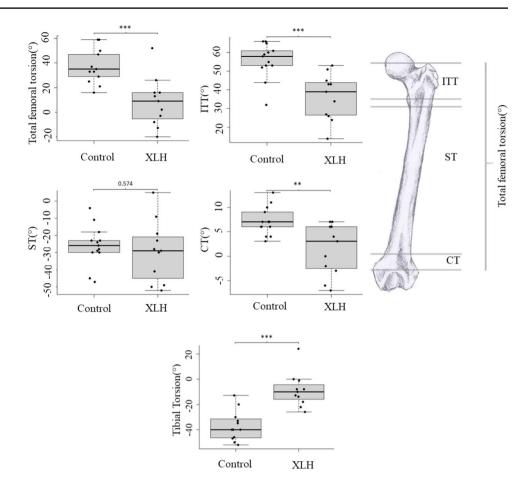
There was no apparent difference in tibial lateral bowing, whereas for tibial frontal bowing (which was not normally distributed) the median value of 11.6° (IQR 7.8° , 19.5°) for individuals with XLH was higher than that in controls 5.0° (IQR: $2.9^{\circ}-7.7^{\circ}$, p < 0.005; Fig. 3).

In terms of lower-limb alignment, there was an enhanced valgus of the mechanical axis in the XLH group $(5.6 \pm 5.3^{\circ})$ as compared to the control group $(1.5 \pm 2.5^{\circ}, p < 0.05)$ In contrast, the average femorotibial angles were not significantly different between the groups. Accordingly, there was a significant difference in the femoral mechanical–anatomical angle, representing the difference between the mechanical and anatomical axis. In that regard, the average femoral mechanical-anatomical angle was $9.1 \pm 2.2^{\circ}$ in the XLH group and only $5.1 \pm 0.8^{\circ}$ in controls (p < 0.001; Fig. 4).

Acetabular version was significantly increased in individuals with XLH $(23.6 \pm 8.4^{\circ})$ as compared to those in



Fig. 2 Total and regional torsion of the femur compared between control participants and individuals with XLH. Boxplots indicate median, interquartile range, minimum and maximum values. Femoral torsion, ITT—intertrochanteric torsion, ST-shaft torsion, CT-condylar torsion. Positive degrees represent external torsion from the frontal plane. Asterisks indicate results of group comparisons—*p < 0.05, **p < 0.01,***p < 0.001. Illustrations by X. O'Reilly-Berkeley



the control group $(17.5 \pm 3.9^{\circ}, p < 0.05)$. Additionally, acetabular coverage was 32.6° higher in individuals with XLH $(192.5 \pm 22.6^{\circ})$ than in the control group $(159.9 \pm 9.1^{\circ}, p < 0.001; Fig. 5)$.

The total CSA of the femoral head and at the lesser trochanter showed no differences between the two groups. The total CSA of femoral shaft sites was ~21–24% greater in individuals with XLH compared to the control group (p < 0.015). The bone marrow cavity of the femoral shaft sites was also greater in XLH individuals (all p < 0.05), such that the cortical bone area was similar between groups. The total CSAs of the three tibial shaft sites and the talus were similar between the groups. The bone marrow cavities of the proximal and mid-tibial shaft sites were greater (p < 0.05) and cortical CSAs at all shaft sites lower (p < 0.05) except proximal sites p = 0.069) in the XLH individuals compared to the control subjects (Table 2).

Discussion

This study was the first to objectively characterise and compare multiple components of lower-limb bone geometry between individuals with XLH and controls. We observed considerable differences in multiple clinically relevant aspects, largely in line with previous clinical and self-reports [9, 14]. Specifically, individuals with XLH exhibited significantly lower femoral torsion as compared to controls. Further detailed analysis revealed this difference to result mainly from a lower proximal, intertrochanteric torsion. While lower distal condylar torsion also appeared to contribute, no group differences in shaft torsion were evident. In that regard, one may speculate that this could be due to enhanced torsional strain towards the joint, i.e. at the intertrochanteric and the condylar area. However, other clinical populations have shown different contributions of subtrochanteric and supertrochanteric torsion to total femoral torsion [22]. This is clinically important, as when pharmacological treatment during growth to prevent femoral deformities has failed and femoral osteotomy is considered [23], it is important to correct torsion in the correct region [13, 22]. To date, surgical protocols focus on frontal plane deformations [7, 24, 25]; therefore, assessment of femoral torsion should also inform development of future guidelines. Furthermore, characterising the overall lower limb geometry of this clinical group could inform about possible sources of joint incongruity, fractures, osteoarthritis and gait disturbances other than frontal plane deformations.



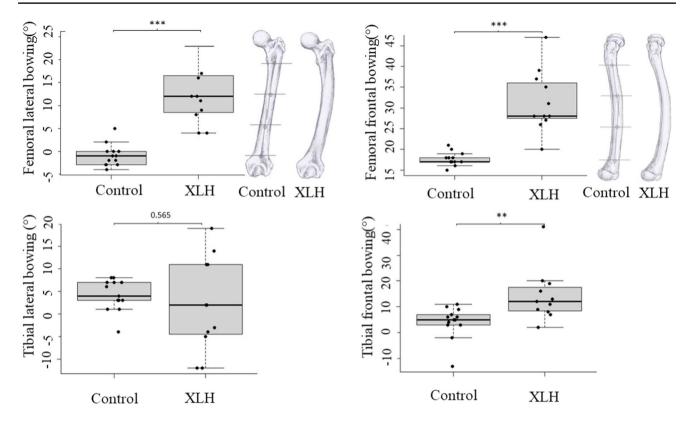
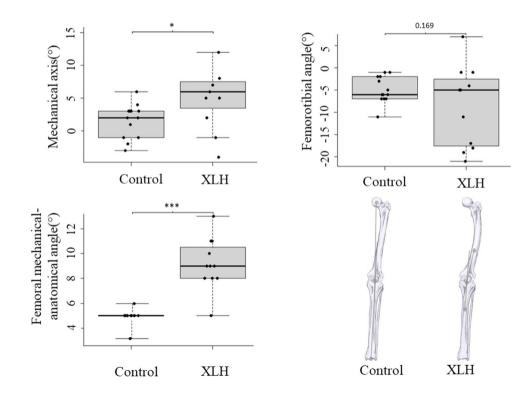


Fig. 3 Femoral and tibial bowing in the two groups; positive values indicate anterior and lateral bowing. Boxplots indicate median, interquartile range, minimum and maximum values. *p < 0.05, **p < 0.01, ***p < 0.001. The illustration in the top left panel shows a visual representation of the average difference between the two groups from a

frontal view. On the control panel, the method of the bowing's measurement is drawn on. The illustration in top right panel shows the average difference from a lateral view. Illustrations by X. O'Reilly-Berkeley

Fig. 4 Mechanical axis, femorotibial angle and femoral mechanical-anatomical angle in the two groups. Boxplots indicate median, interquartile range, minimum and maximum values. Positive degrees are outward lateral. *p < 0.05, **p < 0.01, ***p < 0.001. The bottom right panel shows the visual representation of lower-limb alignment. Also, the mechanical axis is drawn on the control limb and the femorotibial angle on the XLH limb. Illustrations by X. O'Reilly-Berkeley





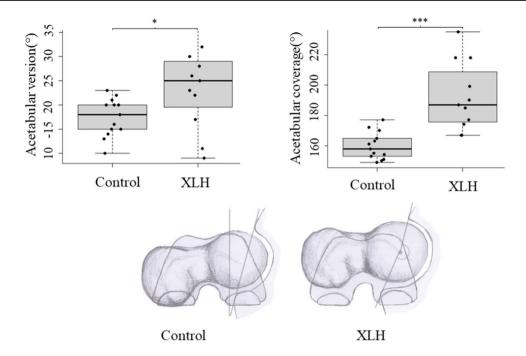


Fig. 5 Acetabular version and acetabular coverage for the two groups. For the acetabular version, positive values indicate anterior rotation from a lateral position, for the acetabular coverage values indicate the portion of the femoral head within the acetabulum. Boxplots indicate median, interquartile range, minimum and maximum values. *p < 0.05, **p < 0.01, ***p < 0.001. The bottom panel shows a visual

representation of the acetabular geometry, the femoro-acetabular relationship and the total femoral torsion with the outline of the knee in the background. On the control panel, the acetabular version measurement is drawn on, and on the XLH panel we can see the acetabular coverage measurement. Illustrations by X. O'Reilly-Berkeley

Moreover, we observed significantly increased acetabular coverage of the hip joints in the XLH cohort. This is particularly important since both reduced femoral torsion and increased acetabular coverage are known to predispose to hip osteoarthritis development [26]. Conversely, a recent large study in a Caucasian population with end-stage knee osteoarthritis [27] revealed a correlation between increasing coronal valgus alignment and increased external condylar torsion as well as increased tibial torsion while in our XLH cohort, coronal valgus deformity correlated with decreased condylar torsion and decreased tibial torsion, suggesting that mechanisms of knee osteoarthritis in XLH might differ from what is observed in the general population. These findings may be important contributions to better understand early osteoarthritis development in XLH [8, 28]. Moreover, it is important to keep in mind that increased acetabular coverage and diminished ITT as the main contributor to low femoral torsion are not directly affected by corrective surgeries in XLH, usually addressing the diaphyseal/distal femur and the proximal tibia. Along the diaphyseal segment, we observed increased frontal and lateral bowing in the XLH cohort at the femur, while differences concerning bowing of the tibia were less pronounced and only significant with regards to frontal bowing. Finally, there was lower external rotation of the tibia (tibial torsion) in line with previous reports [7].

There was larger inter-individual variation for most aspects of skeletal geometry in people with XLH, and the causes and consequences for this need to be identified. The above results agree with current qualitative literature about deformities in patients with XLH [7, 25, 29]; however, detailed quantification of limb geometry in this clinical group was not yet available in the literature.

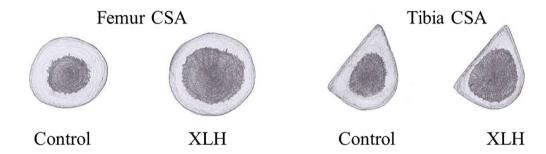
The observation of a significantly greater total crosssectional area of the femur is remarkable, considering the overall reduced body size of XLH patients as compared to controls. This is in line with a previous peripheral quantitative computed tomography (pQCT) study in tibia in children and adults (age 23 ± 13 years) with XLH [30]. Looking at it more closely, the difference results from an enlarged marrow cavity on all height levels along the femoral axis while the cortical cross-sectional area is not significantly different on these levels. In contrast, the generally larger tibia marrow cavity in the absence of a pronounced advantage in total CSA in XLH resulted in lower cortical CSA throughout tibia length. One may speculate if the pronounced femoral hypertrophy results from childhood rickets with compromised bone quality and deficient mineralization leading to enhanced mechanical strain and increased bone turnover with intensified endosteal bone resorption and periosteal apposition.



Table 2 Cross sectional area (CSA) of femur and tibia at multiple locations. Where the marrow cavity was not present, only total CSA has been evaluated. CSA is outlined in mm². In the % column, we show the difference between XLH and control in percentage with the

control group as reference. Bottom panel, visual representation of femur CSA on the left, and tibial CSA on the right. Illustrations by X. O'Reilly-Berkeley

Bone		Site	Total CSA						Bone marrow CSA						Cortical bone CSA					
			XLH		Control					XLH		Control				XLH		Control		
			mean	SD	mean	SD	%	p	mean	SD	mean	SD	%	p	mean	SD	mean	SD	%	p
Femur	Head centre		1840	274	1923	303	-4	0.559			ļ.		ļ	ļ						
	Under lesser trochanter		1093	223	966	159	13	0.182												
	Shaft	Proximal 3rd	823	132	674	104	22	0.015	389	116	172	39	126	<0.001	431	117	502	75	-14	0.147
		Middle	829	137	671	100	24	0.011	355	128	203	64	75	0.004	452	138	468	67	-3	0.752
		Distal 3rd	853	202	703	108	21	0.062	382	54	312	72	22	0.042	487	180	391	59	25	0.136
	Distal flat shaft		1790	277	1551	174	15	0.039						•						
Tibia		Proximal 3rd	701	144	670	93	5	0.592	439	92	344	73	28	0.026	262	87	326	50	-20	0.069
	Shaft	Middle	526	86	516	75	2	0.79	278	81	164	28	70	<0.001	247	61	351	66	-30	0.003
		Distal 3rd	460	78	469	136	-2	0.881	214	60	158	124	35	0.254	246	57	310	44	-21	0.014
	Talus		1755	220	1618	230	8	0.221			•		*	•			•			



Overall, there are likely a number of factors which contribute to the observed findings. Fundamentally, the osteoid content of subjects with osteomalacia results in higher ductility of the bone structure [31]. This abnormality likely leads to changes in deformation patterns caused by mechanical loading during daily activities and exercise, in addition to the resultant adaptive response in bone geometry. In addition, increased bone ductility may also underlie the higher acetabular coverage due to the pressure of the femoral head on the acetabulum. Altered bone composition and mechanical loading may also contribute to the greater variance observed in the group with XLH. In relation to mechanical loading, XLH is associated with muscle weakness [32] and altered gait [8, 14] which likely result in altered bone loading. In a similar vein, childhood motor development, which has been shown to affect development of bone geometry [33, 34], is delayed in children with XLH [35].

These results have both functional and clinical implications for individuals with XLH. Waddling gait and early osteoarthritis at weight-bearing joints are commonly diagnosed in individuals with XLH [7, 8, 14]. In particular, the high prevalence of osteoarthritis in up to 55% even in individuals under the age of 30 is of particular concern [9, 36]. Also, altered femoral geometry likely contributes to the high prevalence of lower limb and specifically femoral

pseudo-fractures in XLH [37]. Femoral torsion and proximal femur varus deformities have been shown to change the lever arms of hip muscles and therefore decrease hip muscle strength [22, 38, 39]. This alteration has been hypothesized to be the cause of waddling gait [25]. Recent gait analysis in subjects with XLH has found that waddling gait was correlated to varus limb alignment [14] and degenerated hip, knee and ankle joints and enthesopathy of the hip abductors [8]. Lower femoral torsion and high acetabular coverage, both confirmed risk factors of hip osteoarthritis [26], impede hip internal rotation in XLH [8, 40] which results in enhanced risk of femoro-acetabular hip impingement and consecutive joint pain [7, 10]. The lower femoral torsion also increases the shear forces on the femoral neck [11]; this could contribute to development of coxa vara which is common in this population. Tibial torsion also contributes to altered gait as identified in paediatric populations with XLH [14].

Corrective surgery is a common practice in individuals with XLH, with > 57% of adults with XLH and frontal plane deformities [7, 24, 25] having already undergone at least one procedure. In the analysis presented here, the average femorotibial angle was similar in both groups. While this finding appears to conflict with the common perception of XLH being associated with varus deformity of the leg axis, it most likely reflects that all but one subject in the XLH cohort



evaluated here have undergone femoral osteotomies which are usually aimed at correction of frontal plane deformities [24]. In addition, the average mechanical valgus angle was actually increased in the XLH group as compared to controls, along with an increased femoral mechanical—anatomical angle, reflective of an enhanced femoral offset and further supporting this perception, while this aspect of previous corrective osteotomies has to be considered a limitation of our study, otherwise.

However, when planning osteotomies taking into account transverse plane parameters such as regional femoral torsion could be beneficial [41, 42], for example to restore advantageous muscular lever arm length. The restoration of limb alignment in multiple planes could decrease the chances of early arthrosis due to malalignment [43].

To our knowledge, this is the first study to quantitatively characterize multiple objectives and clinically relevant measures of lower-limb bone geometry, in individuals with XLH. Previous studies had reported the incidence of identified deformities such as tibial bowing without a description of the magnitude of deviation or the degree of within-group variance [9, 20].

There are several limitations to this study which have to be kept in mind. Importantly, all statistical analyses and interpretations have to be seen against the background of the limited number of cases for this complex assessment in a rare disorder, and lack of significance in some parameters may be related to the low number of patients enrolled. Furthermore, all but two participants with XLH had corrective surgery on the lower limbs during their lifetime; therefore, observed differences are not solely attributable to differences in natural growth. Indeed, the outcome of corrective interventions and the risk for recurrent deformity or overcorrections are subject to multiple modifiers, including but not limited to surgical technique and performance, age at surgery and quality of accompanying medical treatment. However, considering that it is unethical to let XLH patients grow into adulthood without treating their deformities and understanding limitations regarding availability of data on treatments that occurred 10-50 years ago, real-world data provided here still represent best available evidence. Having said that, data presented here should not be perceived to reflect natural bone development in XLH but rather to provide a real-world inventory of bone deformities in adult XLH patients.

Moreover, the fact that specific differences particularly concerning aspects of rotation and bowing persist even following surgery highlights both the scale of the effects and that different interventional approaches, e.g. pharmacological treatments or physiotherapy/exercise, may be required to comprehensively address this issue. Even though the participants were recruited from a broad clinical cohort, the sample size is small. Therefore, while the observed

group differences were large and statistical evidence for them was strong, these findings require replication in a larger cohort. Furthermore, only the left leg was scanned for every subject, as scanning both legs would have quadrupled scanning times. We were therefore unable to assess inter-limb asymmetries, which may contribute to gait and other problems within individuals with XLH. Detailed information on previous pharmacological and other treatments from childhood was not available for all participants, and future studies should examine whether surgical or other treatments impact on bone geometry. Femoral torsion was assessed on two cross-section slices and not on one oblique slice which is now standard practice, while the latter results in slightly smaller torsion values withinpopulation variance is similar [44]; therefore, we believe that this would not substantially impact our main findings. Finally, all frontal plane measurements were assessed using cross-sectional MRIs, and not by the usual clinical method of anteroposterior radiographs.

Conclusions

In conclusion, adults with XLH have substantial differences and greater inter-individual variation in hip and femur bone geometry, principally lower femoral torsion originating in the intertrochanteric region, lower tibial torsion and higher lateral and frontal femoral bowing. Large differences in acetabular coverage and version were also observed, while differences detected between the groups in the mechanical axis and tibiofemoral angle are minimal. These differences are likely to be due to the higher malleability of the bones, variations in gait, delayed motor development and impaired muscle function. These differences were evident despite most individuals within the current study having undergone corrective frontal plane surgery. Accordingly, these observations should be considered when planning corrective surgery in XLH in the future. The presented results have the potential for increased femoral fracture risk and disadvantageous joint mechanics during daily physical activity and could be part of the cause of the osteoarthritis, pain and waddling gait commonly experienced in this clinical group.

Supplementary Information The online version contains supplementary material available at https://doi.org/10.1007/s00198-022-06385-z.

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Declarations

Ethics approval Ethical approval for the study was obtained from the competent ethics committees of the medical faculties at the universities of Cologne (no. 19-1020) and Wuerzburg (128/19). The study is registered with the German Register for Clinical Studies (DRKS00016074).

Consent to participate All participants provided written informed consent prior to any study-related procedures.

Conflict of interest MS, SK, JZ, JJ, OS, ES, JR, and AL have no conflict of interest to disclose. LS has received funding from Kyowa Kirin GmbH to support parts of this research.

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