Clinical characteristics and the identification of novel mutations of *COL1A1* and *COL1A2* in 61 Chinese patients with osteogenesis imperfecta

HAO ZHANG, HUA YUE, CHUN WANG, WEIWEI HU, JIEMEI GU, JINWEI HE,
WENZHEN FU, YUNQIU HU, MIAO LI and ZHENLIN ZHANG

Department of Osteoporosis and Bone Diseases, Metabolic Bone Diseases and Genetic Research Unit,
Shanghai Key Clinical Center for Metabolic Disease, Shanghai Jiao Tong University Affiliated Sixth People's Hospital,
Shanghai 200233, P.R. China

Received December 14, 2015; Accepted September 15, 2016

DOI: 10.3892/mmr.2016.5835

Abstract. Osteogenesis imperfecta (OI) is an inherited connective tissue disorder characterized by brittle bone fractures. The aim of the present study was to investigate the pathogenic gene mutation spectrum and clinical manifestations of mutations in collagen type I, alpha 1 (*COL1A1*) and collagen type I, alpha 2 (*COL1A2*) genes in Chinese patients with OI. A total of 61 unrelated Chinese OI patients with *COL1A1* and *COL1A2* mutations were recruited. All the exons and the exon-intron boundaries of the *COL1A1* and *COL1A2* genes were amplified and directly sequenced and lumbar spine bone mineral density was measured by dual-energy X-ray absorptiometry. The mutations of the 61 probands included 33 missense mutations, 8 nonsense mutations, 7 splicing variants and 13 frameshift mutations in *COL1A1* and *COL1A2* genes. A total of 25 novel mutations were identified, including 18 in *COL1A1* and 7 in *COL1A2*. The mutations p.Gly257Arg, p.Gly767Ser and p.Gly821Ser in *COL1A1* and p.Gly337Ser in *COL1A2* may be located at a mutation hotspot for human OI due to the high repetition rate in OI patients. Family history was positive for OI in 33 probands (54%). All probands had suffered fractures and the most common fracture site was the femur. A total of 49 probands presented with blue sclerae (80.3%), 20 probands suffered from dentinogenesis imperfecta (32.8%) and 1 patient

had hearing loss (1.6%). These findings may improve understanding of the pathogenic gene mutation spectrum and the clinical manifestations of mutations of *COL1A1* and *COL1A2* genes in Chinese patients with OI.

Introduction

Osteogenesis imperfecta (OI) is an inherited connective tissue disorder characterized by brittle bone fractures and short stature. In addition, other connective tissue disorders, including blue or gray sclera, dentinogenesis imperfecta, hyperlaxity of ligaments and skin, and progressive conductive hearing loss, are common in OI patients (1). OI patients have been classified into four types by Sillence and Rimoin (2) according to clinical characteristics and patterns of inheritance. An expanded Sillence classification was published by Rauch and Glorieux in 2004 (3). Forlino *et al* (4) classified OI into 11 types based on the discovery of each novel gene defect. To date, a total of 19 genes have been identified as causing OI: Collagen, type I, alpha 1 (*COL1A1*; MIM 120150), collagen, type I, alpha 2 (*COL1A2*; MIM 120160), cartilage-associated protein (*CRTAP*; MIM 605497), leucine- and proline-enriched proteoglycan 1 (*LEPRE1*; MIM 610339), peptidyl-prolyl isomerase B (*PPIB*; MIM 123841), FK506 binding protein 10 (MIM 607063), serpin family H member 1 (MIM 600943), Wnt family member 1 (MIM 164820), bone morphogenetic protein 1 (MIM 112264), interferon induced transmembrane protein 5 (*IFITM5*; MIM 614757), plastin 3 (MIM 300131), procollagen-lysine, 2-oxoglutarate 5-dioxygenase 2 (MIM 601865), serpin peptidase inhibitor, clade F (alpha-2 antiplasmin, pigment epithelium derived factor), member 1 (MIM 172860), Sp7 transcription factor (MIM 606633), transmembrane protein 38B (MIM 611236), cAMP responsive element binding protein 3-like 1 (HGNC ID 18856; https://oi.gene.le.ac.uk/home.php?select_db=CREB3L1), prolyl 4-hydroxylase, beta polypeptide (MIM 176790), SEC24 family member D (MIM 607186) and Secreted protein, acidic, cysteine-rich (*SPARC*; MIM 182120) (5-8). Although numerous genetic causes of OI have been reported, mutations in *COL1A1* and *COL1A2* are responsible for ~90% of OI cases (9). To date,

Correspondence to: Professor Zhenlin Zhang, Department of Osteoporosis and Bone Diseases, Metabolic Bone Diseases and Genetic Research Unit, Shanghai Key Clinical Center for Metabolic Disease, Shanghai Jiao Tong University Affiliated Sixth People's Hospital, 600 Yishan Road, Shanghai 200233, P.R. China
E-mail: zzl2002@medmail.com.cn

Abbreviations: OI, osteogenesis imperfecta; *COL1A1*, collagen type I, alpha 1; *COL1A2*, collagen type I, alpha 2; BMD, bone mineral density; DXA, dual-energy X-ray absorptiometry

Key words: osteogenesis imperfect, collagen type I, alpha 1, collagen type I, alpha 2, mutation, Chinese

the *COL1A1* and *COL1A2* mutations in Chinese patients with OI have been reported only in certain sporadic cases (10-14). To investigate the pathogenic gene mutation spectrum and the clinical manifestations of mutations in *COL1A1* and *COL1A2* genes in Chinese patients with OI requires a large sample size.

Our previous study (14) identified 56 heterozygous mutations in *COL1A1* and *COL1A2*, including 43 mutations in *COL1A1* and 13 mutations in *COL1A2*. Of those Chinese patients, 56.9% had OI type I, 19.0% had type III and 20.7% had type IV. In addition, the study identified 2 novel compound heterozygous mutations in the *LEPRE1* gene in two probands with OI. Our other previous study identified mutations in the *IFITM5* gene in four Chinese families with OI type V (15). To more accurately reflect the Chinese OI clinical characteristics and pathogenic gene mutations, the present study recruited a further 61 OI patients from 2012 to 2015. Mutations in the *COL1A1* and *COL1A2* genes were identified in the 61 Chinese OI patients. Although the exact incidence of OI in China remains unknown, this is a further large report of Chinese OI cases. The aim of the present study was to investigate the pathogenic gene mutation spectrum and clinical manifestations of mutations of *COL1A1* and *COL1A2* genes in Chinese patients with OI. These data may be useful for future clinical diagnosis and genetic counseling.

Materials and methods

Subjects. A total of 61 unrelated probands from 61 separate families were recruited from the Department of Osteoporosis and Bone Diseases, Shanghai Jiao Tong University Affiliated Sixth People's Hospital (Shanghai, China) over a 2-year period (2012-2015). The probands came from 16 provinces of China, with the majority from Eastern cities. All patients were of Han ethnicity. A total of 410 DNA samples were collected from the 61 probands, 99 family members and 250 healthy control donors. Samples from healthy controls were sequenced to determine whether mutations occurred as polymorphisms. Clinical characteristics, including scleral hue, the presence of dentinogenesis imperfecta and hearing loss, a history of fractures, height, and OI type, were recorded. The OI type was classified in all probands according to the Sillence classification (2). None of the patients belonged to a consanguineous family or had received any treatment prior to the present study. All family members were examined by a single experienced clinician familiar with OI.

Ethics statement. The present study was approved by the Ethics Committee of Shanghai Jiao Tong University Affiliated Sixth People's Hospital. Written informed consent was obtained from all adult participants prior to recruitment. In addition, written informed consent was obtained from parents on behalf of pediatric participants. This consent procedure was approved by the Ethics Committee of the Shanghai Jiao Tong University Affiliated Sixth People's Hospital.

Collagen type I mutation analysis. Genomic DNA was extracted from 2 ml peripheral blood samples using the QuickGene DNA whole blood kit (Kurabo Industries Ltd., Osaka, Japan) and a Nucleic Acid Isolation system (QuickGene-610L; Autogen, Inc., Holliston, MA, USA).

All exons of the *COL1A1* and *COL1A2* genes, including the exon-intron boundaries, were amplified by polymerase chain reaction using the primers described in our previous study (14). The genomic (AF017178.2 and AF004877.1) and mRNA (Z74615.1 and Z74616.1) sequences of the *COL1A1* and *COL1A2* genes were used as reference sequences. DNA mutations were numbered according to the cDNA sequence and the A of the ATG codon was designated as nucleotide +1. Novel mutations were identified according to the Type 1 Collagen Mutation Database (https://oi.gene.le.ac.uk/variants.php?select_db=COL1A1&action=view_all and https://oi.gene.le.ac.uk/variants.php?select_db=COL1A2&action=view_all). In addition, control alleles from 250 healthy individuals were sequenced to determine whether novel mutations occurred as polymorphisms.

To perform a cross-species analysis of the gene sequences of the mutations that were not substitution mutations of glycine, the Uniprot database (<http://www.uniprot.org/uniprot/?query=COL1A1&sort=score> and <http://www.uniprot.org/uniprot/?query=COL1A2&sort=score>) was used to align 10 protein sequences, and the Clustal Omega program (<http://www.ebi.ac.uk/Tools/msa/clustalo/>) was used to view their characteristics alongside each other. The 10 *COL1A1* protein sequences compared were from human, zebrafish, Japanese common newt, chicken, African clawed frog, mouse, rat, cow, dog and cat. The 10 *COL1A2* protein sequences compared were human, zebrafish, green anole or horse, chicken, African clawed frog, mouse, rat, cow, dog and cat, since the protein sequence of Japanese common newt of *COL1A2* was not included in the database.

Bone densitometry. The bone mineral density (BMD; g/cm²) of the lumbar spine (L1-L4) was measured using dual-energy X-ray absorptiometry (DXA). Vertebrae were excluded from analysis if they were affected by fractures. All subjects were assessed using Lunar Prodigy equipment (GE Healthcare Life Sciences, Chalfont, UK). The Lunar Prodigy device was calibrated daily. The coefficient of variability of the lumbar spine DXA measurements was 1.39% (16). All DXA scans were conducted by the same trained specialist. Lumbar spine BMD results were converted to age- and gender-specific Z-scores as previously described (17).

Statistical analysis. Statistical analyses were performed in SPSS software version 11.0 (SPSS, Inc., Chicago, IL, USA). Group differences in enumeration data were analyzed for significance using the chi-square test. Data with normal distribution was tested for significance using independent-samples *t*-test and data with non-normal distribution using the Mann-Whitney U test. All tests were two-sided. *P* < 0.05 was considered to indicate a statistically significant difference.

Results

Clinical characteristics of the 61 patients. A total of 44 probands had a mutation in the *COL1A1* gene, and 17 had a mutation in *COL1A2* (Table I). Of the patients, 35 were male and 26 female, with a median age of 11.0 years (range, 0.4-60 years). Family history was positive for OI in 33 probands (54%), negative for OI in 27 probands (44%) and unknown

Table I. Clinical findings of OI in the 61 patients.

Mutation	Sex (M/F)	Age (years)	Fr >10	Blue sclerae	DI	Hearing loss	Height Z score	Weight Z score	LS BMD Z score
<i>COL1A1</i> (n=44)	28/16	11.0 (4.5-22.5)	6 (13.6%)	37 (84.1%)	15 (34.1%)	1 (2.3%)	-0.8 (-2.5, 0.6)	-0.5 (-1.2, 2.1)	-1.0±1.4
<i>COL1A2</i> (n=17)	7/10	9.0 (7.0-20.0)	0	12 (70.6%)	5 (29.4%)	0	-1.2 (-2.5, -0.7)	0.2 (-1.1, 0.9)	-2.7±0.6

Normally distributed data are expressed as the mean ± standard deviation. Data that are not normally distributed are expressed as the median (inter-quartile range). M, male; F, female; Fr, fracture; DI, dentinogenesis imperfecta; LS BMD, lumbar spine bone mineral density; *COL1A1*, collagen type I, alpha 1; *COL1A2*, collagen type I, alpha 2.

in 1 proband (2%). All probands had suffered fractures, and 13.2% of probands with *COL1A1* mutations had suffered >10 fractures (Table I). A total of 49 probands presented with blue sclerae (80.3%), 20 probands suffered from dentinogenesis imperfecta (32.8%) and 1 patient had hearing loss (1.6%).

***COL1A1* and *COL1A2* mutations.** A total of 44 patients had a mutation within the *COL1A1* gene, and 17 had a mutation in *COL1A2* (Table I). These mutations included 33 missense mutations, 8 nonsense mutations, 7 splicing variants and 13 frameshift mutations (Table II). Almost half the probands (42.6%; 26/61) had a substitution mutation of the glycine within the Gly-X-Y triplet domain of the triple helix, of which 13 were in *COL1A1* and 13 were in *COL1A2*. Serine substitutions were the most common in the present study (42.3%).

In total, 25 of the mutations of 26 probands (2 probands had the same novel mutation in *COL1A2*) identified were novel: 18 in *COL1A1* (Table III) and 7 in *COL1A2* (Table IV). Novel mutations were identified according to the Type 1 Collagen Mutation Database. In addition, these novel mutations were not present in the 250 healthy controls. Of the 25 novel mutations, 7 missense mutations, 7 frameshift mutations and 4 nonsense mutations were identified in *COL1A1*, and 6 missense mutations and 1 nonsense mutation were identified in *COL1A2*. Of these mutations only 6 were not substitution mutations of glycine. Therefore, the gene sequences of these 6 mutations were compared across species. Although the p.Ala508Thr missense mutation of *COL1A1* is not at a highly conserved position, other vertebrate species including cow, dog and cat all contained alanine (Fig. 1A). The proband was classified as OI type III and *COL1A2*, *CRTAP* and *LEPRE1* gene mutations were excluded. The mutation is potentially functionally damaging as alanine is a nonpolar and threonine a polar amino acid. The p.Thr766Ser (Fig. 1B) and p.Thr1298Ile (Fig. 1C) missense mutations of *COL1A1*, and the p.Asp37Gly (Fig. 2A), p.Arg1258His (Fig. 2B) and p.Phe4Leu (Fig. 2C) missense mutations of *COL1A2* are at highly conserved positions. In addition, these missense variants were searched for in the large Exome Aggregation Consortium (ExAC) database (exac.broadinstitute.org), and the frequencies of p.Ala508Thr of *COL1A1*, p.Arg1258His and p.Phe4Leu missense mutations of *COL1A2* were revealed to be 2.471×10^{-5} , 2.471×10^{-5} and 8.236×10^{-6} , respectively. The remaining three missense variants were absent from

Table II. Types of *COL1A1* and *COL1A2* mutations in the present study.

Mutation	Missense	Nonsense	Splicing	Frameshift
<i>COL1A1</i>	17	7	7	13
<i>COL1A2</i>	16	1	0	0

COL1A1, collagen type I, alpha 1; *COL1A2*, collagen type I, alpha 2.

the ExAC database. Furthermore, the combined annotation dependent depletion (CADD) pathogenicity scores of these missense variants were high. The CADD pathogenicity score of the p.Ala508Thr, p.Thr766Ser and p.Thr1298Ile mutations of *COL1A1* and the p.Aso37Gly, p.Arg1258His and p.Phe4Leu mutations of *COL1A2* were 13.3, 26.6, 15.8, 17.8, 23.4 and 19.8, respectively.

The number of fractures in patients with the missense mutation close to the carboxyl-terminal end was greater compared with patients with missense mutations close to the amino terminal. In addition, 2 probands had a p.Gly257Arg (c.769G>A) mutation at exon 11 of *COL1A1*, which is the mutation identified in 2 unrelated patients in our previous study (14); 1 proband had a p.Gly767Ser (c.2299G>A) mutation at exon 33, which is the mutation identified in 4 unrelated patients in our previous study; 1 proband had a p.Gly821Ser (c.2461G>A) mutation at exon 37, which is the same mutation identified in 2 unrelated patients in our previous study; and 2 probands had a p.Gly337Ser (c.1009G>A) mutation at exon 19 of *COL1A2*, which is the same mutation identified in 1 unrelated patient in our previous study.

According to the Sillence classification, no patients in the present study had OI type II. A total of 40 patients (65.6%) had OI type I, 11 patients (18.0%) had type III and 10 patients (16.4%) had type IV. As in our previous study, *COL1A1* mutations were more frequent than *COL1A2* mutations in patients with OI types I and III ($P < 0.05$; Fig. 3). In the *COL1A1* mutation group, the most common fracture site was the femur (n=21; 21% of all fractures), followed by tibia/fibula (n=19; 19%) and radius/ulna (n=19; 19%). In the *COL1A2* mutation group, the most common fracture site was the femur (n=12; 40% of all fractures), followed by tibia/fibula (n=5; 17%), radius/ulna (n=3; 10%) and humerus (n=3; 10%).

Table III. Clinical and genetic characteristics of 44 probands with mutations in collagen type I, alpha 1.

No.	Sex	Age (years)	F/S ^a	Blue sclerae	DI	Hearing loss	Height, cm (Z value)	Weight, kg (Z value)	Fracture rates ^b	Nucleotide change ^c	Predicted amino acid change ^d	Novel
C1	F	12	F	+	-	-	134.0 (-2.5)	40 (0.6)	+	c.144delT	p.His48GlnfsX26	Yes
C2	M	2	F	+	-	-	87 (-0.6)	13 (0.3)	+	c.157-158delTG	p.Trp53GlufsX19	Yes
C3	F	14	F	+	-	-	162.6 (0.7)	68 (2.8)	+	c.268G>T	p.Glu90X	Yes
C4	M	13	S	+	-	-	151 (-0.3)	51 (2.7)	+	c.433_434insC	p.Gly145AlafsX24	Yes
C5	M	12	S	+	-	-	150.8 (0.6)	47 (1.4)	+	c.441dupC	p.Gly148ArgfsX21	-
C6	M	18	F	+	+	-	163 (-1.5)	70 (2.1)	+++	c.484C>T	p.Gln162X	Yes
C7	M	2.3	S	+	-	-	88 (-1.4)	12 (-0.8)	+	c.569delC	p.Pro190LeufsX75	Yes
C8	M	8	F	+	+	-	133.6 (0.7)	28 (0.5)	+	c.573_574delCCinsG	p.Pro193LeufsX72	Yes
C9	M	2	F	+	-	-	84 (-1.5)	12 (-0.4)	+	c.579delT	p.Gly194ValfsX71	-
C10	F	33	F	+	+	-	156.1 (-0.5)	47 (-1.0)	++	c.643-2A>G	Splicing variant	-
C11	F	30	F	+	+	+	152.8 (-1.0)	49 (-0.8)	+	c.757C>T	p.Arg253X	-
C12	F	28	F	+	-	-	147.5 (-2.1)	44 (-1.2)	+	c.769G>A	p.Gly257Arg	-
C13	M	42	S	+	+	-	150 (-2.8)	45 (-2.3)	+	c.769G>A	p.Gly257Arg	-
C14	M	11	F	+	-	-	150 (1.0)	50 (2.2)	+	c.898C>T	p.Gln300X	Yes
C15	M	4	S	+	-	-	116 (3)	22 (3.3)	+	c.903+1G>A	Splicing variant	-
C16	F	6	F	-	+	-	115.5 (-0.4)	17 (-0.9)	+	c.1121G>C	p.Gly374Ala	-
C17	M	2.8	S	+	-	-	101 (1.7)	20 (3.7)	+	c.1128delT	p.Gly377AlafsX164	-
C18	F	65	F	+	-	-	150.2 (-0.5)	53 (-0.5)	+	c.1155+1G>A	Splicing variant	-
C19	F	27	S	+	-	-	161.1 (0.2)	51 (-0.4)	+	c.1200+1G>A	Splicing variant	-
C20	M	7	F	+	-	-	134.5 (1.9)	35 (3.0)	+	c.1243C>T	p.Arg415X	-
C21	F	4	F	+	-	-	105.0 (0.6)	17 (0.7)	+	c.1299+1G>A	Splicing variant	-
C22	M	11	F	+	-	-	137.2 (-0.8)	30 (-0.5)	+	c.1299+1G>A	Splicing variant	-
C23	F	2.5	S	+	+	-	80.0 (-3.3)	10 (-2.2)	+	c.1522G>A	p.Ala508Thr	Yes
C24	M	2.5	-	+	+	-	76.0 (-4.9)	9 (-3.5)	+	c.1588G>A	p.Gly530Ser	-
C25	M	29	F	-	+	-	157.0 (-2.3)	58 (-0.9)	+	c.1678G>T	p.Gly560Cys	-
C26	M	8	F	-	-	-	112.5 (-4.2)	20 (-1.2)	+	c.1787G>C	p.Gly596Ala	Yes
C27	M	12	F	+	-	-	152.0 (0.8)	52 (2.0)	+	c.1866delT	p.Gly623AlafsX143	-
C28	F	11	S	+	+	-	160.0 (1.9)	52 (2.6)	+	c.1930-2A>C	Splicing variant	-
C29	M	12	S	+	-	-	150.0 (0.5)	55 (2.4)	+	c.2183G>A	p.Gly728Glu	Yes
C30	F	26	S	+	+	-	134.5 (-4.2)	40 (-1.7)	++	c.2297C>G	p.Thr766Ser	Yes
C31	F	24	S	+	+	-	118.0 (-6.9)	44 (-1.0)	++	c.2299G>A	p.Gly767Ser	-
C32	M	8	S	+	-	-	120.0 (-2.5)	20 (-1.2)	+	c.2410G>T	p.Glu804X	Yes
C33	M	28	F	+	+	-	173.5 (0.3)	64 (-0.6)	+	c.2450delC	p.Pro817LeufsX291	-
C34	F	16	S	-	+	-	114.5 (-7.3)	32 (-2.6)	+++	c.2461G>A	p.Gly821Ser	-

Table III. Continued.

No.	Sex	Age (years)	F/S ^a	Blue sclerae	DI	Hearing loss	Height, cm (Z value)	Weight, kg (Z value)	Fracture rates ^b	Nucleotide change ^c	Predicted amino acid change ^d	Novel
C35	F	10	F	-	-	-	110.0 (-4.1)	23 (-1.4)	++	c.2560G>A	p.Gly854Ser	-
C36	M	7	F	+	-	-	117.0 (-0.8)	21 (-0.7)	+	c.2775delT	p.Gly926ValfsX182	-
C37	F	25	S	+	+	-	126.0 (-5.7)	40 (-1.7)	++	c.2867G>C	p.Gly956Ala	Yes
C38	M	14	S	+	-	-	159.5 (-0.8)	45 (-0.6)	+	c.3076C>T	p.Arg1026X	-
C39	M	8	F	+	-	-	127.4 (-0.8)	24 (-0.4)	+	c.3328delC	p.His110ThrfsX129	Yes
C40	M	13	F	+	-	-	154.8 (0.1)	75 (8.0)	+	c.3559G>T	p.Gly1187Cys	Yes
C41	M	3	S	+	-	-	81.0 (-4.3)	11 (-2.3)	+	c.3638delG	p.Gly1213AlafsX26	Yes
C42	M	4	F	-	-	-	106.0 (0.6)	20 (2.1)	+	c.3655G>A	p.Asp1219Asn	-
C43	M	0.4	S	+	-	-	60.0 (-1.2)	6 (-1.1)	++	c.3893C>T	p.Thr1298Ile	Yes
C44	M	9	S	-	-	-	139.0 (1.4)	40.0 (2.0)	+	c.4363G>A	p.Gly1455Ser	-

^aF, familial; S, sporadic. ^b+, <10; ++, 10-20; +++, >20. ^cMutation numbering is based on cDNA sequences. +1 corresponds to the A of the ATG translation initiation codon. ^dAmino acids are numbered from the translation initiator methionine. DI, dentinogenesis imperfecta.

There were no significant differences in the phenotypes of patients with glycine to serine mutations of *COL1A1* and *COL1A2* genes (Table V).

Discussion

The present study identified 61 mutations in *COL1A1* and *COL1A2*, 25 of them novel. Of the 25 novel mutations, 13 missense mutations, 5 nonsense mutations and 7 frameshift mutations were identified. To date, 912 unique mutations in *COL1A1* are listed on the OI variant database (oi.gene.le.ac.uk/home.php?select_db=COL1A1), 571 unique mutations in *COL1A2* (oi.gene.le.ac.uk/home.php?select_db=COL1A2), plus a further 5 novel mutations that have recently been discovered (18-20). Despite the number of mutations identified, the 25 novel mutations observed in the present study may contribute to revealing the pathogenesis of OI and improve the disease-causing gene spectrum of OI in humans; the novel glycine substitution and frameshift mutations are of particular interest in this regard. In addition, the present study identified various missense mutations with unclear pathogenicities, including the p.Ala508Thr, p.Thr766Ser and p.Thr1298Ile mutations of *COL1A1* and the p.Asp37Gly, p.Arg1258His and p.Phe4Leu mutations of *COL1A2*. They occur at highly or relatively highly conserved positions and their ExAC control frequencies were very low or absent. Furthermore, the CADD pathogenicity scores of these missense variants were high. Therefore, these 6 novel missense mutations may be pathogenic.

As in our previous study, almost three times the number of mutations was observed in *COL1A1* compared with *COL1A2*. The *COL1A1* group contains patients with stop or frameshift mutations that lead to haploinsufficiency and OI type I. However, these haploinsufficiency mutations are not observed in *COL1A2*, as they do not result in a phenotype.

In the present study, of the *COL1A1* mutations, 4 probands had a p.Gly257Arg mutation at exon 11, 5 probands had a p.Gly767Ser mutation at exon 33 and 3 probands had a p.Gly821Ser mutation at exon 37. The Type 1 Collagen Mutation Database lists these mutations 30, 26 and 21 times, respectively. In addition, 3 probands had a p.Gly337Ser mutation at exon 19 of *COL1A2*. The Type 1 Collagen Mutation Database lists this mutation 23 times. Therefore, these 3 mutations in *COL1A1* and 1 mutation in *COL1A2* may occur at a mutation hotspot for human OI.

Type I procollagen chains form a heterotrimer containing two copies of the $\alpha 1$ (I) chain and one copy of the $\alpha 2$ (I) chain; it has therefore been suggested that phenotypes resulting from mutations in *COL1A1* were more severe compared with those from *COL1A2* (21). However, the present study observed no significant differences between *COL1A1* and *COL1A2* glycine to serine mutation groups, in contrast to a previous study by Rauch *et al* (22). The number of patients examined in the present study was small, which may limit the statistical power. As mutations toward the carboxyl-terminus are more disruptive to helix formation, it has been suggested that mutations closest to the carboxyl-terminal end would be more severe compared with those closer to the amino-terminal end of the α (I) chains (23,24). However, Rauch *et al* (22) demonstrated this 'gradient model' of disease severity for $\alpha 2$ (I) mutations but not for mutations affecting $\alpha 1$ (I). In the present study,

Table IV. Clinical and genetic characteristics of 17 probands with mutations in collagen type I, alpha 2.

No.	Sex	Age (years)	F/S ^a	Blue sclerae	DI	Hearing loss	Height, cm (Z value)	Weight, kg (Z value)	Fracture rates ^b	Nucleotide change ^c	Predicted amino acid change ^d	Novel
H1	M	13	F	+	+	-	142.3 (-1.2)	48 (2)	+	c.12T>G	p.Phe4Leu	Yes
H2	F	9	F	+	-	-	122 (-1.6)	26.5 (0.2)	+	c.110A>G	p.Asp37Gly	Yes
H3	M	7	S	+	+	-	113.3 (-0.8)	23 (0.7)	+	c.812G>A	p.Gly271Asp	Yes
H4	F	8	S	-	-	-	120 (-1.3)	21 (-0.8)	+	c.946G>A	p.Gly316Ser	-
H5	M	6	S	+	-	-	113 (-0.8)	19 (-0.5)	+	c.1009G>A	p.Gly337Ser	-
H6	F	7	F	+	+	-	112.8 (-1.1)	17.2 (-1.5)	+	c.1009G>A	p.Gly337Ser	-
H7	F	47	F	+	-	-	158 (-0.1)	47.5 (-1.4)	+	c.1081C>T	P.Arg361X	Yes
H8	F	9	F	-	-	-	126.5 (-2.3)	48.3 (1.5)	+	c.1648G>A	p.Gly550Ser	-
H9	M	19	F	+	-	-	160 (-2)	76 (1.8)	+	c.1666G>T	p.Gly556Cys	Yes
H10	M	21	S	-	-	-	145 (-2.6)	50 (-1.1)	+	c.2081G>A	p.Gly694Asp	Yes
H11	M	25	S	+	+	-	136.5 (-5.6)	56 (-1.0)	+	c.2081G>A	p.Gly694Asp	Yes
H12	M	7	F	+	-	-	116.5 (-0.6)	23 (0.5)	++	c.2441G>A	p.Gly814Glu	-
H13	M	13	S	-	-	-	150 (-0.4)	42 (0.6)	+	c.2764G>A	p.Gly922Ser	Yes
H14	F	43	S	-	-	-	136.9 (-4.2)	64 (0.8)	+	c.2918G>A	p.Gly973Asp	-
H15	M	11	S	+	+	-	135.0 (-1.2)	30 (-0.5)	+	c.3197G>T	p.Gly1066Val	-
H16	M	3	S	+	-	-	87 (-2.6)	11 (-2.3)	+	c.3304G>A	p.Gly1102Ser	-
H17	F	8	F	+	-	-	131.5 (-0.5)	30.7 (0.9)	+	c.3773G>A	p.Arg1258His	Yes

^aF, familial; S, sporadic. ^b+, <10; ++, 10-20; +++, >20. ^cMutation numbering is based on cDNA sequences. +1 corresponds to the A of the ATG translation initiation codon. ^dAmino acids are numbered from the translation initiator methionine. DI, dentinogenesis imperfecta.

Figure 1. Missense mutations occur at conserved positions in COL1A1, as demonstrated by a comparison of the sequences of ten species. (A) The missense mutation p.Ala508Thr occurs at a relatively conserved position in COL1A1. (B) p.Thr766Ser occurs at a highly conserved position in COL1A1. (C) p.Thr1298Ile occurs at a highly conserved position in COL1A1. COL1A1, collagen type I, alpha 1.

Figure 2. Missense mutations occur at conserved positions in COL1A2, as demonstrated by a comparison of the sequences of ten species. (A) The missense mutation p.Asp37Gly occurs at a highly conserved position in COL1A2. (B) p.Arg1258His occurs at a highly conserved position in COL1A2. (C) p.Phe4Leu occurs at a highly conserved position in COL1A2. COL1A2, collagen type I, alpha 2.

Figure 3. Clinical types of OI caused by COL1A1 and COL1A2 mutations. (A) Clinical types of OI caused by COL1A1 mutations. (B) Clinical types of OI caused by COL1A2 mutations. Checks, OI type I; dots, OI type III; stripes, OI type IV. OI, osteogenesis imperfecta; COL1A1, collagen type I, alpha 1; COL1A2, collagen type I, alpha 2.

proband with the missense mutation at the amino-terminal end had a milder phenotype compared with those closer to the carboxyl-terminal end. It has previously been demonstrated that the most common mutations associated with OI are substitutions for glycine by another amino acid in the triple helical domain of COL1A1 or COL1A2 (22). In the present study, 33 missense mutations were identified from 61 probands. Substitutions of glycine by serine were the most common in COL1A1 and COL1A2, in accordance with our previous study and Rauch *et al* (14,22).

In contrast to our previous study, disease in the 61 patients of the present study was 54% familial and 44% sporadic disease. This was in accordance with a previous study on Korean OI patients (51% familial and 49% sporadic OI cases) (25). In the present study, OI type I (65.6%) was the most common type. A total of 80.3% probands presented with blue sclerae and all patients with mutations in the amino-terminal end of the $\alpha 1(I)$ chain had blue sclera, as described by Rauch *et al* (22), who additionally observed that dentinogenesis imperfecta was absent in patients with mutations in the amino-terminal end of the $\alpha 1(I)$ or $\alpha 2(I)$ triple helical domain. In the present study, patients were identified with COL1A1 mutations at the amino-terminal and carboxyl-terminal ends without dentinogenesis imperfecta. Hearing loss is a common secondary feature of OI in adults, often with mixed conductive and sensorineural deficiency (4). Previous studies have reported that in a Scottish population ~50% of patients have subjective hearing loss (26), and >60% of Finnish OI adult patients had hearing loss (27); however, only ~5% of OI pediatric patients have been reported to suffer hearing loss (28). In the present study, only 1.6% of probands had hearing loss; this may due to the fact that the majority of the patients were children. The most common fracture site was the femur, followed by tibia/fibula, radius/ulna in the patients of the COL1A1 and COL1A2 mutation groups; this is in accordance with the study by Ben Amor *et al* (29).

The present study is the first, to the best of our knowledge, to observe Chinese OI patients in such large numbers.

Table V. Clinical characteristics of patients with glycine to serine substitutions.

Parameter	<i>COL1A1</i> (n=5)	<i>COL1A2</i> (n=6)	P-value
Age (years)	12.3±18.1	7.7±3.3	0.23
Height (Z score)	-4.4±3.5	-1.4±0.9	0.08
Weight (Z score)	-1.3±2.1	-0.5±1.4	0.47

Data are expressed as the mean ± standard deviation. *COL1A1*, collagen type I, alpha 1; *COL1A2*, collagen type I, alpha 2.

However, the present study had certain limitations. The study was limited to a single center and was cross-sectional. No patients had lethal type II OI as the patients were recruited from the outpatient center and newborns with the lethal type were not yet diagnosed.

In conclusion, the present study identified 61 mutations in *COL1A1* and *COL1A2*, 25 of which were novel. Serine substitutions were the most frequently encountered mutation type. The mutations p.Gly257Arg, p.Gly767Ser and p.Gly821Ser in *COL1A1*, and p.Gly337Ser in *COL1A2* may be located at a mutation hotspot for human OI. Family history was positive for OI in 33 probands (54%). All probands had suffered fractures and the most common fracture site was the femur. A total of 49 probands presented with blue sclerae (80.3%), 20 probands suffered from dentinogenesis imperfecta (32.8%) and 1 patient had hearing loss (1.6%). These findings may improve understanding of the pathogenic gene mutation spectrum and clinical manifestations of mutations in *COL1A1* and *COL1A2* genes of Chinese patients with OI, and may be useful for future clinical diagnosis and genetic counseling.

Acknowledgements

The authors thank the patients and their family members for their invaluable cooperation, and the Center for Genetic & Genomic Analysis, Genesky Biotechnologies Inc. (Shanghai, China) for their assistance with gene identification. The present study was supported by the National Basic Research Program of China (grant no. 2014CB942903), the Science and Technology Commission of Chongqing Municipality (grant no. CSTC2013jcyjC00009), the Science and Technology Commission of Shanghai Municipality (grant nos. 14JC1405000 and 14ZR1431900), the National Natural Science Foundation of China (grant nos. 81170803, 81370978 and 30800387), the Frontier Technology Joint Research Program of the Shanghai Municipal Hospitals (grant no. SHDC12013115) and the Shanghai Leading Talent Plan (grant no. 051).

References

- Basel D and Steiner RD: Osteogenesis imperfecta: Recent findings shed new light on this once well-understood condition. *Genet Med* 11: 375-385, 2009.
- Sillence DO and Rimoin DL: Classification of osteogenesis imperfect. *Lancet* 1: 1041-1042, 1978.
- Rauch F and Glorieux FH: Osteogenesis imperfecta. *Lancet* 363: 1377-1385, 2004.
- Forlino A, Cabral WA, Barnes AM and Marini JC: New perspectives on osteogenesis imperfecta. *Nat Rev Endocrinol* 7: 540-557, 2011.
- Van Dijk FS and Sillence DO: Osteogenesis imperfecta: Clinical diagnosis, nomenclature and severity assessment. *Am J Med Genet A* 164A: 1470-1481, 2014.
- Rauch F, Fahiminiya S, Majewski J, Carrot-Zhang J, Boudko S, Glorieux F, Mort JS, Bächinger HP and Moffatt P: Cole-Carpenter syndrome is caused by a heterozygous missense mutation in *P4HB*. *Am J Hum Genet* 96: 425-431, 2015.
- Garbes L, Kim K, Rieß A, Hoyer-Kuhn H, Beleggia F, Bevo A, Kim MJ, Huh YH, Kweon HS, Savarirayan R, *et al*: Mutations in *SEC24D*, encoding a component of the COPII machinery, cause a syndromic form of osteogenesis imperfecta. *Am J Hum Genet* 96: 432-439, 2015.
- Mendoza-Londono R, Fahiminiya S, Majewski J; Care4Rare Canada Consortium; Tétreault M, Nadaf J, Kannu P, Sochett E, Howard A, Stimec J, *et al*: Recessive osteogenesis imperfecta caused by missense mutations in *SPARC*. *Am J Hum Genet* 96: 979-985, 2015.
- Cheung MS and Glorieux FH: Osteogenesis imperfecta: Update on presentation and management. *Rev Endocr Metab Disord* 9: 153-160, 2008.
- Liu W, Gu F, Ji J, Lu D, Li X and Ma X: A novel *COL1A1* nonsense mutation causing osteogenesis imperfecta in a Chinese family. *Mol Vis* 13: 360-365, 2007.
- Qin W, He JX, Shi J, Xing QH, Gao JJ, He L, Qian XQ, Liu ZJ, Shu AL and He L: Mutation detection of *COL1A1* gene in a pedigree with osteogenesis imperfecta. *Yi Chuan Xue Bao* 32: 248-252, 2005 (In Chinese).
- Xia XY, Cui YX, Huang YF, Pan LJ, Yang B, Wang HY, Li XJ, Shi XC, Lu HY and Zhou YC: A novel RNA-splicing mutation in *COL1A1* gene causing osteogenesis imperfecta type I in a Chinese family. *Clin Chim Acta* 398: 148-151, 2008.
- Lu Y, Ren X, Wang Y, Li T, Li F, Wang S, Xu C, Wu G, Li H, Li G, *et al*: Mutational and structural characteristics of four novel heterozygous C-propeptide mutations in the *proα1(I)* collagen gene in Chinese osteogenesis imperfecta patients. *Clin Endocrinol (Oxf)* 80: 524-531, 2014.
- Zhang ZL, Zhang H, Ke YH, Yue H, Xiao WJ, Yu JB, Gu JM, Hu WW, Wang C, He JW and Fu WZ: The identification of novel mutations in *COL1A1*, *COL1A2*, and *LEPRE1* genes in Chinese patients with osteogenesis imperfecta. *J Bone Miner Metab* 30: 69-77, 2012.
- Zhang Z, Li M, He JW, Fu WZ, Zhang CQ and Zhang ZL: Phenotype and genotype analysis of Chinese patients with osteogenesis imperfecta type V. *PLoS One* 8: e72337, 2013.
- Zhang H, He JW, Gao G, Yue H, Yu JB, Hu WW, Gu JM, Hu YQ, Li M, Fu WZ, *et al*: Polymorphisms in the *HOXD4* gene are not associated with peak bone mineral density in Chinese nuclear families. *Acta Pharmacol Sin* 31: 977-983, 2010.
- Maynard LM, Guo SS, Chumlea WC, Roche AF, Wisemandle WA, Zeller CM, Towne B and Siervogel RM: Total-body and regional bone mineral content and areal bone mineral density in children aged 8-18 y: The Fels Longitudinal Study. *Am J Clin Nutr* 68: 1111-1117, 1998.
- Wang Y, Cui Y, Zhou X and Han J: Development of a high-throughput resequencing array for the detection of pathogenic mutations in osteogenesis imperfecta. *PLoS One* 10: e0119553, 2015.
- Hoefele J, Mayer K, Marschall C, Alberer M, Klein HG and Kirschstein M: Rare co-occurrence of osteogenesis imperfecta type I and autosomal dominant polycystic kidney disease. *World J Pediatr*: April 8, 2016. (Epub ahead of print). doi: 10.1007/s12519-016-0014-1.
- Lin HY, Chuang CK, Su YN, Chen MR, Chiu HC, Niu DM and Lin SP: Genotype and phenotype analysis of Taiwanese patients with osteogenesis imperfecta. *Orphanet J Rare Dis* 10: 152, 2015.
- Prockop DJ, Constantinou CD, Dombrowski KE, Hojima Y, Kadler KE, Kuivaniemi H, Tromp G and Vogel BE: Type I procollagen: The gene-protein system that harbors most of the mutations causing osteogenesis imperfecta and probably more common heritable disorders of connective tissue. *Am J Med Genet* 34: 60-67, 1989.
- Rauch F, Lalic L, Roughley P and Glorieux FH: Genotype-phenotype correlations in nonlethal osteogenesis imperfecta caused by mutations in the helical domain of collagen type I. *Eur J Hum Genet* 18: 642-647, 2010.

23. Byers PH: Brittle bones-fragile molecules: Disorders of collagen gene structure and expression. *Trends Genet* 6: 293-300, 1990.
24. Marini JC, Forlino A, Cabral WA, Barnes AM, San Antonio JD, Milgrom S, Hyland JC, Körkkö J, Prockop DJ, De Paepe A, *et al*: Consortium for osteogenesis imperfecta mutations in the helical domain of type I collagen: Regions rich in lethal mutations align with collagen binding sites for integrins and proteoglycans. *Hum Mutat* 28: 209-221, 2007.
25. Lee KS, Song HR, Cho TJ, Kim HJ, Lee TM, Jin HS, Park HY, Kang S, Jung SC and Koo SK: Mutational spectrum of type I collagen genes in Korean patients with osteogenesis imperfecta. *Hum Mutat* 27: 599, 2006.
26. Paterson CR, Monk EA and McAllion SJ: How common is hearing impairment in osteogenesis imperfecta? *J Laryngol Otol* 115: 280-282, 2001.
27. Kuurila K, Kaitila I, Johansson R and Grénman R: Hearing loss in Finnish adults with osteogenesis imperfecta: A nationwide survey. *Ann Otol Rhinol Laryngol* 111: 939-946, 2002.
28. Kuurila K, Grenman R, Johansson R and Kaitila I: Hearing loss in children with osteogenesis imperfecta. *Eur J Pediatr* 159: 515-519, 2000.
29. Ben Amor IM, Roughley P, Glorieux FH and Rauch F: Skeletal clinical characteristics of osteogenesis imperfecta caused by haploinsufficiency mutations in COL1A1. *J Bone Miner Res* 28: 2001-2007, 2013.