Pathophysiology and therapeutic options in osteogenesis imperfecta: an update

Abstract: Osteogenesis imperfecta (OI) is a rare, heritable systemic disorder of bone and connective tissue, which in almost 90% of cases is due to mutations affecting the normal synthesis of type I collagen. In 1979, four OI phenotypes were categorized which were inherited as autosomal dominant characteristics. Individuals with OI present both genetic and phenotypic variabilities. Major characteristics of OI are bone fragility, blue sclerae, dentinogenesis imperfecta, short stature, scoliosis, and joint hyperextensibility. Both autosomal dominant and recessive inheritance are now recognized. Advances in molecular diagnosis have led to a major expansion in our understanding of the genetic basis for different OI phenotypes. To date, sequence variants in 17 genes are described as causative of OI. These genes regulate the synthesis of type I collagen pro-alpha polypeptide chains, proteins involved in type I collagen processing in the endoplasmic reticulum and proteins involved in osteoblast function. These new genetic associations have also led to uncertainty with regard to the current classification of OI phenotypes. Bisphosphonates have been widely used to improve bone mass and decrease fractures in both children and adults with OI. While effective in many but not all children when administered for 2–4 years, bisphosphonates have not proven effective in adults with OI. Studies are limited for treatment of adults with teriparatide and denosumab. Advances have been reported in the surgical management of OI. Although the role of physical therapy in the management of children and adults was previously described, this important treatment modality is significantly underutilized.

Keywords: osteogenesis imperfecta, sequence variants, collagen, bisphosphonates, bone, treatment

Introduction

Osteogenesis imperfecta (OI) is a rare clinically and genetically heterogeneous systemic disorder of bone and connective tissue characterized by bone fragility and physical findings related to the underlying connective tissue disorder. The incidence of OI is approximately 6–7/100,000 births, and approximately 500,000 persons worldwide have OI.1 In USA, there are between 25,000 and 50,000 affected individuals.2 Individuals with OI have a lifelong risk of fracture, which may occur at times spontaneously or following minimal trauma. Other systemic manifestations are blue sclerae, dentinogenesis imperfecta (DI), early-onset hearing loss, short stature, kyphoscoliosis, and joint hyperextensibility. Less frequent are basilar invagination, cardiac valvular and vascular disease, and ocular complications, including glaucoma and decreased corneal thickness.1–5

There has been remarkable progress in understanding the molecular basis for this disease. Over the last decade, advances in molecular biology have greatly expanded
the association of previously unrecognized gene sequence variants with phenotypes now categorized as OI. As a consequence, the spectrum of OI phenotypes is now considerably enhanced compared to the four phenotypes first described by Sillence et al. That classification employed clinical severity, radiological features, and inheritance, dividing OI into four types: mild (OI type I), lethal (OI type II), severe progressive (OI type III), and moderately deforming (OI type IV). These four types were inherited in an autosomal dominant manner and each was subsequently shown to involve mutations affecting type I collagen pro-alpha chains.

The Online Mendelian Inheritance in Man database currently lists OI types I–XVII based on recently defined sequence variants including several proteins not in the classic collagen pathway. Approximately 90% of OI cases involve autosomal dominant inheritance with mutations in the COL1A1 or COL1A2 genes. To date, the Human Collagen Mutation Database (http://www.le.ac.uk/genetics/collagen/) lists the following genes associated with a phenotype within the broad clinical spectrum considered OI: BMP1, COL1A1, COL1A2, CREB3L1, CRTAP, FKB10, IFITM5, P3H1, P4HB, PLOD2, PLS3, PPIB, SEC24D, SERPINF1, SERPINH1, SP7, TMEM38B, WNT1, and SPARC.

**OI classification**

A classification of OI can be based on a definitive genotype or on a phenotype that is highly variable and more difficult to categorize in many individuals even within the same family. Furthermore, expression of one gene locus can translate into different skeletal phenotypes. Access to genetic testing is not available for many individuals. Therefore, many individuals with OI do not know what clinical type they represent or the genetic basis for their phenotype. Recognizing the emerging confusion due to the expansion of genetic data, the International Skeletal Dysplasia Society kept the Sillence classification with the inclusion of type V OI and without any direct molecular reference. The new causative OI genes have been arranged according to clinical severity into the five OI types (Table 1). In 2010, Van Dijk et al proposed a revised classification of OI with the exclusion of type VII and VIII OI, since these types were added based on genotype but with clinical and radiological characteristics indistinguishable from Sillence types II and IV OI.

It is important to emphasize that the diagnosis of OI still requires historical, clinical, and radiological evidence of the disorder. Fracture history per se in a child does not establish the diagnosis of OI, and several other skeletal disorders including secondary causes of osteoporosis can mimic OI clinically. These include hypophosphatasia, idiopathic juvenile osteoporosis, celiac disease, Ehlers–Danlos syndrome, vitamin D deficiency, and hypophosphatemic disorders. In addition, child physical abuse is relevant considering that this is more frequent than OI and usually presents a history of multiple fractures, but in a different OI.

The recommendation of this review, from a clinical standpoint, is that a clinical classification such as the one suggested by Bonafe et al seems to be the best option. Probably, clinical features will continue being the basis for OI diagnosis and newer genetic findings will be adapted into recognizable phenotypes.

**Phenotypes and genotypes of OI**

The discovery of new genotypes related to OI phenotypes has been significantly expanded with the advent of single-nucleotide polymorphism array technology and the

**Table 1 OI nomenclature according clinical severity**

<table>
<thead>
<tr>
<th>OI classification</th>
<th>Clinical feature</th>
<th>Gene</th>
<th>Inheritance</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Nondeforming OI</td>
<td>COL1A1</td>
<td>AD</td>
<td></td>
</tr>
<tr>
<td>form</td>
<td>COL1A2</td>
<td>AD</td>
<td></td>
</tr>
<tr>
<td>2 Perinatally lethal OI</td>
<td>COL1A1</td>
<td>AD</td>
<td></td>
</tr>
<tr>
<td>form</td>
<td>COL1A2</td>
<td>AD</td>
<td></td>
</tr>
<tr>
<td>CRTAP</td>
<td>LEPRE1</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>PPIB</td>
<td></td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>3 Progressively deforming OI</td>
<td>COL1A1</td>
<td>AD</td>
<td></td>
</tr>
<tr>
<td>form</td>
<td>COL1A2</td>
<td>AD</td>
<td></td>
</tr>
<tr>
<td>BMP1</td>
<td>CRTAP</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>FKB10</td>
<td>LEPRE1</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>PLOD2</td>
<td>PPIB</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>SERPINF1</td>
<td></td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>SERPINH1</td>
<td>TMEM38B</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>WNT1</td>
<td>CREB3L1</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>4 Moderate OI form</td>
<td>COL1A1</td>
<td>AD</td>
<td></td>
</tr>
<tr>
<td></td>
<td>COL1A2</td>
<td>AD</td>
<td></td>
</tr>
<tr>
<td></td>
<td>WNT1</td>
<td>AD</td>
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<tr>
<td>CRTAP</td>
<td>PPIB</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>FKB10</td>
<td>SP7</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>SERPINF1</td>
<td>PLS3</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td></td>
<td>IFITM5</td>
<td>AR</td>
<td></td>
</tr>
<tr>
<td>5 OI with calcification in interosseous membranes</td>
<td>COL1A1</td>
<td>AD</td>
<td></td>
</tr>
</tbody>
</table>


**Abbreviations:** AD, autosomal dominant; AR, autosomal recessive; XL, X-linked; OI, osteogenesis imperfecta.
increased application of whole-exome sequencing. The genes described as OI-causative regulate the synthesis of type I collagen alpha polypeptide chains, several proteins involved in type I collagen processing, or proteins involved in osteoblast development and function. Nonetheless, the correlation between genotype and phenotype in OI is extremely complex due to high variability in the expressivity of the disorder (Table 2).

The majority of OI cases are related to autosomal dominant inheritance caused by heterozygosity for mutations in the COL1A1 or COL1A2 genes corresponding to OI types I, II, III, and IV. Individuals with OI type I may experience few fractures associated or not to mild deformities (Figure 1A). OI type II is characterized by severe bone fragility and neonatal mortality. OI type III is a severe form leading to multiple fractures, even in utero, and bone deformities (Figures 1B and 2). OI type IV presents moderate bone involvement with high clinical variability. De novo mutations are responsible for approximately 60% of mild (OI-I) or moderately deforming OI (OI-IV), almost 100% of severely deforming OI (OI-III), and practically all perinatal lethal cases (OI-II). Occasionally, type III and IV OI are the result of autosomal recessive inheritance.

OI type V is an autosomal dominant disorder with a distinctive phenotype representing 4%–5% of OI cases. OI type V is the result of heterozygous mutations (approximately −14C>T) in the 5′UT region of the IFITM5 gene, which encodes interferon-induced transmembrane protein. It is characterized by variable degree of severity and specific features such as hyperplastic callus, dislocation of the radial head, and progressive calcification of the interosseous membranes of forearms and legs. However, a child with clinical features of type III/IV OI was identified with a de novo IFITM5 mutation (approximately −14C>T) but without the classic features described for OI-V.

OI types VI–XVI are recessively inherited. Collectively, these account for approximately 3%–5% of individuals with OI and express moderately severe to lethal phenotypes, which in some variants clinically resemble OI types II and III.

OI-VI is associated with homozygous or compound heterozygous mutations in SERPINH1. This gene encodes pigment epithelium derived factor that is a multifunctional secreted glycoprotein of the serpin family with potent antiangiogenic activity. It functions in bone homeostasis and osteoid mineralization. Affected individuals have a moderate-to-severe OI, presenting decreased bone mineral density (BMD) with no fractures at birth and without blue sclerae or DI. Bone biopsy shows a characteristic “fish-scale” pattern caused by disorganization of the bone matrix, with large amount of unmineralized osteoid tissue simulating osteomalacia.

Individuals with OI types VII, VIII, and IX have severe to lethal phenotypes caused, respectively, by mutations in the CRTAP, LEPRE1, and PPIB genes. Approximately 1.5% of West Africans and 0.4% of African Americans carry a founder mutation in LEPRE1 gene and three lethal cases of OI-VIII were reported in individuals of these ethnicities. In addition to features common to severe OI phenotypes, mutations in CRTAP and LEPRE1 genes (OI-VII, OI-VIII) are associated with formation of bulbous epiphysis with popcorn calcifications at the distal femurs.

Type X OI is the result of mutations involving the SERPINH1 gene, which regulates folding and trafficking of the collagen triple helix in the endoplasmic reticulum. This mutation was reported in a child from Saudi Arabia with triangular facies, blue sclerae, micrognatia, severe bone deformities, and bilateral renal calculi leading to impaired renal function.

The main features of Bruck syndrome are bone fragility with congenital contractures of large joints. It is classified as types 1 and 2 according to the gene mutated, FKBP10 (OI-XI) and PLOD2, respectively; however, the phenotypes are indistinguishable (Figure 3). PLOD2 mutation expression in a boy from nonconsanguineous Turkish parents led to congenital joint contractures with pterygium and severe OI-like osteopenia including multiple fractures. Interestingly, two brothers from a Spanish family also with PLOD2 mutations had phenotypes of different severity, one had a mild form with few fractures and white sclerae and the second had a more severe form with multiple fractures and blue sclerae; however, neither had congenital joint contractures.

SP7 gene is associated with OI type XII. This moderate form of OI was described in one child from an Egyptian family displaying decreased BMD, recurrent fractures, mild bone deformities, white sclerae, delayed teeth eruption, and absence of DI.

Mutations in BMP1 gene are categorized as OI type XIII. It was identified in a few individuals with severe but variable phenotypes. Individuals of Egyptian origin had generalized decreased BMD, light blue sclerae, delay of motor development, severe growth deficiency, bone deformities, platispondyly, kyphoscoliosis, umbilical hernia, and several spontaneous fractures. Nonetheless, Turkish individuals had increased BMD, white sclerae, and normal motor development, also, multiple fractures after minimal
<table>
<thead>
<tr>
<th>OI type</th>
<th>OMIM</th>
<th>Gene(s)</th>
<th>Locus</th>
<th>Protein(s)</th>
<th>Defective mechanism</th>
<th>Phenotype</th>
<th>Inheritance</th>
</tr>
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<tbody>
<tr>
<td>I</td>
<td>166200</td>
<td>COL1A1/COL1A2</td>
<td>17q21.3</td>
<td>Collagen alpha-1(i) chain/Collagen alpha-2(i) chain</td>
<td>Collagen quantity</td>
<td>Mild</td>
<td>AD</td>
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<tr>
<td>II</td>
<td>166220</td>
<td>COL1A1/COL1A2</td>
<td>7q21.3</td>
<td>Collagen alpha-1(i) chain/Collagen alpha-2(i) chain</td>
<td>Collagen structure</td>
<td>Lethal</td>
<td>AD</td>
</tr>
<tr>
<td>III</td>
<td>259420</td>
<td>COL1A1/COL1A2</td>
<td>17q21.3</td>
<td>Collagen alpha-1(i) chain/Collagen alpha-2(i) chain</td>
<td>Collagen structure</td>
<td>Progressive deformity</td>
<td>AD; AR</td>
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<tr>
<td>IV</td>
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<td>COL1A1/COL1A2</td>
<td>7q21.3</td>
<td>Collagen alpha-1(i) chain/Collagen alpha-2(i) chain</td>
<td>Collagen structure</td>
<td>Moderate</td>
<td>AD; AR</td>
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<tr>
<td>V</td>
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<td>IFRM5</td>
<td>11p15.5</td>
<td>Interferon-induced transmembrane protein 5</td>
<td>Matrix mineralization/ossification</td>
<td>Moderate, hypertrophic callus and ossification of the interosseous membrane</td>
<td>AD</td>
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<td>VI</td>
<td>613982</td>
<td>SERPINF1</td>
<td>17p13.3</td>
<td>Pigment epithelium derived factor (PEDF)</td>
<td>Matrix mineralization/ossification</td>
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<td>610682</td>
<td>CRTAP</td>
<td>3p22.3</td>
<td>Cartilage-associated protein (CRTAP)</td>
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<td>LEPRE1/P3H1</td>
<td>1p34.2</td>
<td>Prolyl 3-hydroxylase 1 (P3H1)</td>
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<td>Severe to lethal</td>
<td>AR</td>
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<td>IX</td>
<td>259440</td>
<td>PPIB</td>
<td>15q22.3</td>
<td>Peptidyl-prolyl cis-trans isomerase B/Cyclophilin B</td>
<td>Collagen modification</td>
<td>Severe to lethal</td>
<td>AR</td>
</tr>
<tr>
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<td>613848</td>
<td>SERPINC1</td>
<td>11q13.5</td>
<td>Serpin H1/Heat shock protein 47 (HSP47)</td>
<td>Collagen chaperoning</td>
<td>Severe</td>
<td>AR</td>
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<tr>
<td>XI</td>
<td>610968</td>
<td>FBZP10</td>
<td>17q21.2</td>
<td>Peptidyl-prolyl cis-trans isomerase FBZP10</td>
<td>Telopeptide hydroxylation</td>
<td>Progressive deformity with contractures (Bruck syndrome)</td>
<td>AR</td>
</tr>
<tr>
<td>XII</td>
<td>613849</td>
<td>SP7</td>
<td>12q13.13</td>
<td>Transcription factor 7/Osterix</td>
<td>Osseoblast development</td>
<td>Moderate</td>
<td>AR</td>
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<tr>
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<td>614856</td>
<td>BMP1</td>
<td>8p21.3</td>
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<td>XIV</td>
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<td>TMEM38B</td>
<td>9q31.2</td>
<td>Trimeric intracellular cation channel B (TRIC-B)</td>
<td>Osseoblast development</td>
<td>Variable severity</td>
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<td>615220</td>
<td>WNT1</td>
<td>12q13.12</td>
<td>Wingless-type MMTV integration site family, member 1 (WNT1)/Proto-oncogene Wnt-1</td>
<td>Osseoblast development</td>
<td>Variable severity</td>
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<tr>
<td>XVI</td>
<td>616229</td>
<td>CREB3L1</td>
<td>11p11.2</td>
<td>cAMP responsive element binding protein 3-like 1/Old astrocyte specifically induced substance (OASIS)</td>
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<td>3q24</td>
<td>Procollagen-lysine, 2-oxoglutarate, 5-dioxogenase 2</td>
<td>Collagen cross-linking</td>
<td>Severe with congenital joint contractures (Bruck syndrome)</td>
<td>AR</td>
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<tr>
<td></td>
<td>176790</td>
<td>P4HB</td>
<td>17q25.3</td>
<td>Protein disulfide-isomerase (PDI)</td>
<td>Collagen chaperoning</td>
<td>Severe (Cole-Carpenter syndrome-1)</td>
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<td>SEC24D</td>
<td>4q26</td>
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<td>300131</td>
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<td>Xq23</td>
<td>Plastin-3</td>
<td>&quot;Bone development&quot;/&quot;assembly and disassembly of the actin cytoskeleton&quot;</td>
<td>Mild</td>
<td>X-linked</td>
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Abbreviations: AD, autosomal dominant; AR, autosomal recessive; OI, osteogenesis imperfecta; OMIM, Online Mendelian Inheritance in Man database.

trauma, and not all of them had deformities in lower limbs. In addition, a boy with consanguineous parents of Pakistani descent was identified with a homozygous mutation in BMP1 associated with white sclerae, normal bone density, multiple fractures up to age 2, and delay of motor development. A homozygous mutation in TMEM38B gene is associated with OI type XIV, which appears to be an ancient mutation.
Pathophysiology and therapeutic options in OI

Research in Israeli Bedouin and Saudi families from the Arabian Peninsula.\textsuperscript{30,31} Affected individuals had bone involvement of variable severity with decreased BMD, multiple fractures ranging from prenatal onset up to 6 years of age, white or blue sclerae, mild-to-moderate short stature, and normal teeth and hearing.\textsuperscript{30,31}

Mutations in \textit{WNT1} gene are related to OI-XV. Components of the canonical Wnt/\(\beta\)-catenin signaling pathway are integral for normal osteoblast differentiation as well as the differentiation of several types of stem cells.\textsuperscript{32} Fahiminiya et al\textsuperscript{33} have reported \textit{WNT1} mutations in four children from three families. These children had short stature, low bone density, and multiple vertebral compression fractures in addition to multiple long-bone fractures during the first years of life.\textsuperscript{33} A homozygous nonsense mutation was identified in two siblings, this led to brain malformation and severe intellectual disability not associated with major OI phenotypes.\textsuperscript{34}

Keupp et al\textsuperscript{35} have also reported that hypofunctional alleles of \textit{WNT1} (one frameshift mutation, two missense mutations, one splice site mutation, and one nonsense mutation) cause autosomal-recessive OI in consanguineous families. Of note, these authors identified a heterozygous \textit{WNT1} missense mutation in a family affected by dominantly inherited early-onset osteoporosis.\textsuperscript{35} Studies indicated that altered WNT1 proteins fail to activate canonical LRP5-mediated WNT-regulated \(\beta\)-catenin signaling.\textsuperscript{31,34,35} Pyott et al\textsuperscript{36} have reported biallelic loss-of-function mutations in \textit{WNT1} causing a phenotype consistent with recessively inherited type III OI.

OI-XVI is associated with mutations in \textit{CREB3L1} gene. It was reported in a Turkish family with consanguinity. One affected was a fetus from a pregnancy medically interrupted at 19 weeks where postmortem exam showed thin ribs and long-bone fractures. The second child showed intrauterine
fractures, pulmonary infections, and cardiac insufficiency leading to death in the first year of life.\textsuperscript{37}

Mutations in \textit{P4HB} and \textit{SEC24D} genes are causative of Cole–Carpenter syndrome 1 and 2 (CCS-1 and CCS-2), respectively.\textsuperscript{38,39} CCS-1 is an OI-like disorder that is characterized by severe bone fragility, fractures, craniosynostosis, ocular proptosis, hydrocephalus, and dysmorphic facial features.\textsuperscript{38} Individuals initially described with CCS-2 were from a German family and displayed severe pre- and postnatal bone fragility with multiple fractures, diminished ossification of the skull, craniofacial dysmorphism, and short stature.\textsuperscript{39} A study with the zebrafish model showed that Sec24D plays an important role in the extracellular matrix secretion in cartilage. With reference to facial dysmorphism, Sec24D deficiency is shown to affect craniofacial morphogenesis in late stages of chondroblast development.\textsuperscript{40}

In 2013, X-linked osteoporosis with early-onset fractures occurring in childhood was described in five families associated with a mutation in \textit{PLS3}, which encodes the actin-binding protein, plastin 3. The rare heterozygous sequence variant (rs140121121) was associated with decreased BMD and an increased risk of fracture among women in the general population.\textsuperscript{41} \textit{SPARC} (OI-XVII) is the gene most recently associated with an OI phenotype. Two distinctive homozygous missense mutations in \textit{SPARC} were reported from two unrelated girls with severe bone fragility. Besides severe bone fragility, one had severe early-onset scoliosis requiring spinal fusion at age 6 years. The second girl was born prematurely of a consanguineous couple, with hypotonia and gross motor developmental delay. Histomorphometric study showed matrix hypermineralization.\textsuperscript{42}

**Current pharmacological management options**

There is no “optimal treatment” for OI if the primary object of treatment is to limit or prevent fractures in children or adults. Relief from musculoskeletal pain and improvement of compressed vertebral bodies and mobility can also be sought as secondary outcomes. Critical assessment of treatment outcomes is limited by the small numbers of participants in clinical trials and the short duration of many trials, which is frequently limited to 1 or 2 years of observation.

**Bisphosphonates**

For over 2 decades, bisphosphonates (BPs) have been considered the first-line drug for treating children and adults with moderate or severe forms of OI. Individuals with type I OI may be treated with BPs depending on the frequency of fractures. As will be discussed, the use of BPs in OI has recently come under critical review. Third-generation BPs are nitrogen-containing analogs of pyrophosphate, which inhibit osteoclast bone resorption by interfering with the mevalonic acid pathway and protein prenlation.\textsuperscript{43} Bone formation is subsequently increased as osteoblastic bone formation completes mineralization at sites of active bone formation (transients). BPs have strong affinity for bone mineral, particularly at sites of increased bone turnover.\textsuperscript{44} In OI, the response to BP therapy differs between children and adults. BP treatment is generally efficient during childhood due to high bone turnover; however, after puberty there is a decrease in bone turnover, which limits the effectiveness of BP in decreasing fracture rate.\textsuperscript{2,44}

Currently, individual centers in the USA and Europe may use different BPs. Nonetheless, alendronate and risedronate by the oral route and pamidronate and zoledronate by intravenous (IV) route are the most frequently used. It is important to note that there are significant differences with regard to schedules, doses, and duration of treatment among various centers and published studies (Table 3).\textsuperscript{45} Oral BPs are advantageous as they do not require IV access; however, the bioavailability is low and compliance may be poor. Several studies have reported the treatment effects of oral BPs in children and adults, but to date there is no conclusive evidence of its effect on sustained reduction of fracture rate. Two years of alendronate treatment in children with OI was associated with improved lumbar spine areal BMD (LS-aBMD) and decreased bone turnover; however, there was no significant difference in regard to the height of the vertebral bodies, cortical thickness, mobility, bone pain, and fracture incidence.\textsuperscript{46}

Equally beneficial effects following 2 years of BP therapy including increase in BMD, decrease in bone turnover, and a trend to decrease fracture incidence were observed comparing alendronate to pamidronate in a sample of 18 children with OI.\textsuperscript{47} In a randomized, double-blind, placebo-controlled trial, a daily dose of risedronate was administered for 1 year to children with OI.\textsuperscript{48} Approximately 81% had a mild form of OI. After 12 months, risedronate treatment increased LS-aBMD and reduced the risk of recurrent fractures by 42%. However, during an extension open-label phase, 2–3 years after the placebo-controlled phase, there was no difference in lumbar spine and total body aBMD between the groups. In addition, 53% of patients in the treated group versus 65% of patients receiving placebo reported vertebral or nonvertebral fractures.\textsuperscript{48}
Palomo et al reported the results of pamidronate treatment in 37 children with OI who started BP therapy before 5 years of age, and who had a subsequent follow-up period of at least 10 years. At baseline, 35% of vertebrae were affected by compression fractures, whereas only 6% appeared compressed at the final evaluation, indicating vertebral reshaping during growth. Although long-term pamidronate therapy was associated with higher Z-scores for LS-aBMD and vertebral reshaping, the long-bone fracture rate remained high and the majority of patients developed scoliosis.

Bradbury et al treated 37 adults with type I OI with risendronate (35 mg/weekly) for 24 months following which there was an increase on lumbar spine BMD and a decrease in bone biomarkers; however, posttreatment, there was no difference in bone pain or fracture incidence. In addition, Shapiro et al reported a marginal decreased fracture incidence associated with IV BP therapy in adult patients with moderate to severe OI but not in individuals with mild OI, and no significant change was observed with oral BP in any type of OI.

Pamidronate is the most used BP for children with OI; however, more recently zoledronic acid also has been adopted as a treatment option for this population. Zoledronic acid given intravenously is 200-fold more potent than pamidronate and longer-lasting, possibly up to 2 years in children, in suppressing bone turnover. To date, information is limited as regards efficacy and safety in OI patients. However, recent publications indicate the relative short-term safety of zoledronic acid administered at 6-month intervals in children with OI. The advantage of this drug compared to pamidronate is the schedule of infusion, that is, a shorter infusion requires short time and is less frequent. Zoledronate is given for 30 minutes every 6 months and pamidronate is infused for 4 hours every 3–4 months.

There is no data in neonates with OI regarding BP use. In addition, the recent studies by Plotkin et al and Letocha et al. demonstrate that zoledronate and pamidronate can be used in neonates with OI and are effective in increasing bone mass and strength.

### Table 3: Bisphosphonate treatment protocols for pediatric OI

<table>
<thead>
<tr>
<th>Study</th>
<th>Drug</th>
<th>Dose</th>
<th>Route</th>
<th>Age</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sakkers et al</td>
<td>Olpadronate</td>
<td>10 mg/m² daily</td>
<td>Oral</td>
<td>3–18 yrs</td>
</tr>
<tr>
<td>Antoniassi et al</td>
<td>Neridronate</td>
<td>2 mg/kg for 2 d, every 3 mo</td>
<td>IV</td>
<td>Neonatal</td>
</tr>
<tr>
<td>Rauch et al</td>
<td>Risedronate</td>
<td>15 mg weekly (&lt;40 kg); 30 mg weekly (&gt;40 kg)</td>
<td>Oral</td>
<td>6–17 yrs</td>
</tr>
<tr>
<td>Bishop et al</td>
<td>Risedronate</td>
<td>2.5 mg daily (10–30 kg); 5 mg daily (&gt;30 kg)</td>
<td>Oral</td>
<td>4–15 yrs</td>
</tr>
<tr>
<td>Cho et al</td>
<td>Alendronate</td>
<td>10 mg/d (&gt;35 kg), 10 mg every other day (20–35 g), 10 mg every 3 d (&lt;20 kg)</td>
<td>Oral</td>
<td>6.3–15 yrs</td>
</tr>
<tr>
<td>Ward et al</td>
<td>Alendronate</td>
<td>5 mg daily (&lt;40 kg); 10 mg daily (&gt;40 kg)</td>
<td>Oral</td>
<td>4–19 yrs</td>
</tr>
<tr>
<td>DiMeglio and Peacock</td>
<td>Pamidronate</td>
<td>1 mg/kg/d for every 3 d, every 4 mo</td>
<td>Oral/IV</td>
<td>&gt;3 yrs</td>
</tr>
<tr>
<td>Glorieux et al</td>
<td>Pamidronate</td>
<td>1 mg/kg/d for every 3 d, every 4 mo</td>
<td>IV</td>
<td>&gt;3 yrs</td>
</tr>
<tr>
<td>Plotkin et al</td>
<td>Pamidronate</td>
<td>0.5–1.0 mg/kg/d for every 3 d, every 2–4 mo</td>
<td>IV</td>
<td>≤2 yrs</td>
</tr>
<tr>
<td>Letocha et al</td>
<td>Pamidronate</td>
<td>10 mg/m²/d for every 3 d, every 3 mo</td>
<td>IV</td>
<td>4–16 yrs</td>
</tr>
<tr>
<td>Kusumi et al</td>
<td>Pamidronate</td>
<td>0.5 mg/kg/d every 3 d, every 2 mo (&lt;2 yr), 0.75 mg/kg/d every 3 d, every 3 mo (2–3 yr), 1 mg/kg/d every 3 d, every 3 mo (&gt;3 yr)</td>
<td>IV</td>
<td>≤24 mo</td>
</tr>
<tr>
<td>Panigrahi et al</td>
<td>Zoledronate</td>
<td>2 mg (age &lt;6 mo) to 4 mg (age &gt;6 mo) over 30 min to 1 h, every 3–4 mo</td>
<td>IV</td>
<td>2.5 weeks to 7 yr</td>
</tr>
<tr>
<td>Vuorimies et al</td>
<td>Zoledronate</td>
<td>0.05 mg/kg up to 4.0 mg/d over 45 min every 6 mo</td>
<td>IV</td>
<td>1.5–16.8 yr</td>
</tr>
<tr>
<td>Barros et al</td>
<td>Zoledronate</td>
<td>0.025 mg/kg/d every 2 d, every 4 mo (&lt;3 yr), 0.05 mg/kg/day every 2 d, every 4 mo (&gt;3 yr)</td>
<td>IV</td>
<td>2–10 yr</td>
</tr>
<tr>
<td>Otaify et al</td>
<td>Zoledronate</td>
<td>0.1 mg/kg, every 6 mo</td>
<td>IV</td>
<td>0.2–16 yr</td>
</tr>
</tbody>
</table>

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**Abbreviations:** d, day; mo, months; yr, years of age; IV, intravenous; min, minutes; h, hour; OI, osteogenesis imperfecta.

## References

stated an increase of BMD for children and adults with OI following either oral or IV BP treatment, but the effect on decreasing fracture rate was inconclusive. Moreover, the Dwan et al\textsuperscript{60} review found no significant evidence of positive effect of BP on improvement of clinical status such as reduction of pain and functional mobility.

Rijks et al\textsuperscript{61} reviewed ten randomized controlled trials (519 participants) with regard to effectiveness and safety of BP in children with OI. LS-aBMD increased and bone resorption biomarkers decreased following either oral or IV BP. Significant decrease in fracture incidence was reported in 70% of the studies following BP therapy. However, the authors highlighted a common problem in interpreting these treatment reports: 1) fracture incidence was reported in several ways, 2) differentiation between vertebral fractures and fractures of extremities required definition, and 3) confounding factors need to be considered, for example, age, pubertal status, and physical activity.

In 2015, Hald et al\textsuperscript{62} conducted a meta-analysis specifically to assess the effects of BP therapy on the prevention of fractures. Six randomized, placebo-controlled trials (424 subjects) were included. The results of BP therapy on fracture prevention were considered “inconclusive,” and the proportion of participants who experienced a fracture was not significantly reduced.\textsuperscript{62} The meta-analysis by Shi et al\textsuperscript{63} reported the efficacy of BP treatment on BMD and fracture rate in patients with OI. In children, BPs were efficacious in reducing fractures (relative risk [RR] = 0.80; 95% confidence interval [CI] = 0.66--0.97), whereas in adults, BP seemed equivalent to placebo (RR = 0.82; 95% CI = 0.42--1.59). There was also no significant difference in fracture reduction when comparing oral and IV BPs.

BPs are stored in the skeleton for years, although more rapid turnover in children may shorten residence in bone compared to adults.\textsuperscript{64} Although data regarding the safety of BP is reported, issues such as bone turnover after several years of BP treatment go unreported. Delayed eruption of primary teeth following BP therapy is reported in children with OI, and Kamoun-Goldrat and Le Merrer\textsuperscript{65} described a mean delay of 1.67 years in tooth eruption associated to BP treatment. Atypical femoral fracture following prolonged BP therapy previously described in adults with osteoporosis is now reported in a small number of children and adults with OI.\textsuperscript{66--68} It raises questions about oversuppression of bone remodeling and its contribution to microdamage accumulation and increased bone fragility.\textsuperscript{66,69} Hegazy et al\textsuperscript{60} reported the occurrence of unusual femoral stress fractures in six children with OI on long-term pamidronate therapy. These fractures occurred around intramedullary rods in the subtrochanteric or diaphyseal regions of the femur and were caused by no or minimal trauma, similar to the atypical femoral fractures previously described in adults with osteoporosis on long-term BP therapy and with suppression of bone turnover.

Delayed healing of osteotomy sites secondary to BP therapy was previously reported in children and adolescents with OI.\textsuperscript{71} However, the same group recently published new data based on a protocol focused on patients treated with zoledronate, interval of 4 months without BP infusion after osteotomy, and a different surgical technique using a manual osteotomy rather than a high-powered saw.\textsuperscript{72} It was observed that, with the newer approach, delayed healing of the osteotomies over 12 months occurred in 42% of the osteotomies performed versus 72% with the previous approach ($P=0.001$).

In a review published in 2014 of four retrospective studies and one case report involving mean duration of BP from 4.5 to 6.8 years, the authors concluded that there was no evidence to support a causal relationship between BP and osteonecrosis of the jaw in children and adolescents with OI.\textsuperscript{73}

Other effects also associated with BP therapy in OI patients are gastrointestinal pain associated with oral BPs, flu-like symptoms after the first BP infusion, transient asymptomatic hypocalcemia, and fracture nonunion.\textsuperscript{42,46,74,75}

**Teriparatide**

Teriparatide (human recombinant parathyroid hormone) is a parathyroid hormone analog, which increases bone mass by increasing osteoblast bone formation. Highly effective in the treatment of age-related osteoporosis, teriparatide decreases the risk of fracture by increasing trabecular number and trabecular thickness.\textsuperscript{76,77} A randomized, double-blind placebo-controlled trial assessed the effect of teriparatide treatment on 79 adults with OI.\textsuperscript{77} Positive effects on bone markers were observed with a significant increase in hip and spine aBMD, vertebral volumetric BMD, and estimated vertebral strength. This response was more pronounced in type I OI than in individuals with OI types IV and III. There was no significant difference in self-reported fractures when treated and control groups were compared.\textsuperscript{77}

Thirteen postmenopausal women with type I OI previously treated with BPs received daily injections of teriparatide for 18 months.\textsuperscript{78} Lumbar spine BMD increased significantly up to 3.5% but no difference was observed for total hip BMD. As in the Orwoll et al\textsuperscript{77} study, teriparatide significantly increased markers of bone formation and bone resorption suggesting a normal osteoblastic response to therapy. Fracture response...
was not reported. Teriparatide is reported to promote healing of an atypical femoral fracture in an adult patient with OI. This may be related to the ability of this drug to stimulate chondrocyte proliferation and differentiation and cartilage production at the site of fracture.

**Denosumab (anti-RANK-ligand antibody)**

The RANK, RANKL complex regulates bone-remodeling cycles by regulating osteoblast/osteoclast coupling and osteoclast differentiation. RANK is present on the osteoclast precursor, and RANKL produced by the osteoblast is part of the TNF superfamily and, along with the soluble decoy receptor osteoprotegerin, are essential regulators of osteoclast development and function. Denosumab is a human monoclonal antibody to RANKL; studies involving age-related osteoporosis have showed the efficacy of denosumab in reducing signaling via RANK, leading clinically to prevention of bone loss. Due to poor response to BP treatment in OI-VI, Semler et al treated four children with denosumab who had shown continuously elevated urinary bone resorption markers during a previous treatment with BP. Treatment was well tolerated and bone resorption markers decreased to the normal range. However, this report did not address change in BMD or the impact on fracture rate in these children. Subsequently, these authors described long-term denosumab effects over 8–12 cycles of treatment on these same patients. There was a BMD increase, normalization of vertebral shape, and decrease in fracture rate. Denosumab treatment also improved BMD and longitudinal bone growth in two children with COL1A1/A2 mutations previously treated with BPs.

**New therapeutic approaches**

**Antisclerostin antibody**

Sclerostin is a negative regulator of bone formation released from osteocytes that modulates osteoblast activity acting through Wnt/β-catenin pathway. Preclinical studies have demonstrated that treatment with antisclerostin monoclonal antibody improves bone mass and bone strength, and enhances repair of fractures in animal models. Interestingly, at the tissue level, different mechanisms are involved when comparing osteoanabolic therapy (teriparatide) and sclerostin antibody (Scl-Ab). While osteoanabolic therapy increases bone remodeling through increase in both bone formation and resorption, Scl-Ab increases bone formation while decreasing bone resorption.

In humans, anti-sclerostin antibody (romozumab) was administrated to postmenopausal women and healthy men (aged 45–59 years), and the results showed an increase of bone formation markers and a decrease of bone resorption markers. In addition, there was a significant increase in lumbar spine and hip aBMD. In patients with OI, Palomo et al did not find a relationship between circulating serum levels of sclerostin and LS-aBMD.

Sinder et al treated rapidly growing 3-week-old Brtl/+ mice model of OI for 5 weeks with Scl-Ab. Scl-Ab had anabolic effects in Brtl/+ and led to new cortical bone formation and increased cortical bone mass. This anabolic action resulted in improved mechanical strength to wild-type levels without altering the underlying brittle nature of the material.

However, Roschger et al reported somewhat different results using male Coll1a1 (Jrt)/+ mice. Once-weekly IV Scl-Ab injections for growing and adult mice had no effect on weight or femur length. No significant treatment-associated differences occurred in alkaline phosphatase activity, pro-collagen type I N-propeptide, or C-telopeptide. Micro-CT analyses at the femur showed that Scl-Ab treatment was associated with higher trabecular bone volume and higher cortical thickness in wild-type mice at both ages and in growing OI mice, but not in adult OI mice.

**Cathepsin K antibody**

Cathepsin K is highly expressed in osteoclasts and is an essential enzyme involved in the degradation of type I collagen in the organic bone matrix. In an animal model, the cathepsin K monoclonal antibody (Odanacatib) effectively suppressed bone resorption. A Phase III randomized, placebo-controlled trial assessed the effect of Odanacatib on fracture risk over 5 years of treatment in 16,713 women with osteoporosis aged 65 years or older. There was an increase in lumbar spine and total hip BMD over the 5 years and a significant reduction in the risk of fractures of hip (47% reduction, P<0.001), spine (23% reduction, P<0.001), and nonvertebral (23% risk reduction, P<0.001). Applicability to the collagen defect in OI remains to be determined.

**Transforming growth factor-β**

TGF-β is produced by osteoblasts and acts to coordinate bone remodeling by coupling osteoblasts and osteoclasts in the process of bone remodeling. TGF-β is secreted predominantly in an inactive latent form and is deposited into the bone matrix. Several studies have reported on the effect of TGF-β overexpression in bone cells and bone. In 1996, Erlebacher et al used transgenic mice to evaluate the role of TGF-β2 in bone development and observed that
increased expression of TGF-β2 in osteoblasts resulted in an osteoporosis-like phenotype.

In 2000, Gebken et al. studied the cell-surface expression and functional properties of TGF-β receptors I, II, and III on osteoblasts from a group of OI patients compared to healthy controls. The number of TGF-β receptors was significantly higher on OI osteoblasts than on age-matched control osteoblasts in spite of similar steady-state levels for TGF-β receptor II mRNA in OI and control cells. Furthermore, receptor affinity was not significantly diminished. Thus, osteoblasts from OI patients were shown to have an elevated number of cell-surface receptors for TGF-β, without any evidence for a transcriptional regulation of TGF-β receptor II. Excessive TGF-β signaling has been identified in Marfan syndrome involving abnormalities in both lung and aorta.

Grafe et al. have reported that excessive TGF-β signaling is a mechanism of OI in both recessive (Crtap−/−) and dominant (Col1a2tm1.1Mcb) OI mouse models. Higher expression of TGF-β target genes, a higher ratio of phosphorylated Smad2 to total Smad2 protein, and higher in vivo Smad2 reporter activity were observed in these models. The binding of type I collagen of Crtap (−/−) mice showed reduced binding to decorin, a known regulator of matrix TGF-β activity. Treatment of Crtap (−/−) with the anti-TGF-β neutralizing antibody 1D11 corrected the bone phenotype and improved lung abnormalities in both recessive and dominant forms of OI.

Prenatal and postnatal transplantation of mesenchymal stem cells
Severe to lethal forms of OI may be diagnosed in utero by ultrasonography starting at the 16th week. Signs include diffuse osteopenia, poor ossification of the skull, increased nuchal translucency, fractures, delay of growth, micromelia, ventriculomegaly, and bowing of long bones. In the oim/oim mouse, allogenic transplanted wild-type donor mesenchymal stem cells (MSCs) homed to bone, improving collagen content and mineralization. In humans, improvement of linear growth and reduction of fracture rate followed prenatal and postnatal cell transplantation in OI. Additionally, prenatal transplantation of allogeneic MSCs in three OI pregnancies indicated that it has appeared to be safe. A clinical trial in human pregnancy is currently in progress.

Multidisciplinary management
Orthopedic treatment
Orthopedic management and/or surgical treatment may be necessary in cases of severe bone deformity impairing function, with recurrent fractures, nonunion of fracture, severe scoliosis, and basilar invagination where neurosurgical intervention may be required. Long periods of immobilization secondary to recurrent fractures or surgery may lead to reduction of bone mass and loss of muscle strength, thereby increasing the risk of further fractures. Straightening, realignment, and fixation of long bones by intramedullary rodding is a classic technique for limb stabilization in OI patients. The purpose of rodding is to provide fracture protection and to improve alignment of long bone, thus improving function in growing children. The main complication following intramedullary rod fixation is proximal migration of the rod and breakage or disassembly of the rod creating an unprotected segment in the bone.

The need for rodding is limited to severe types of OI. Lower limb deformities were treated with bone splint technique (osteotomies and internal plating combined with cortical strut allografts) in nine children with type I OI (aged 5–12 years). Bone healing occurred within 12–16 weeks after surgery in all patients, and significant improvement on mobility was observed in eight patients. No participant experienced further fractures, deformity, or nonunion. Nonunion of fractures occurs in approximately 20% of OI patients (Figure 4). Surgical repair can be difficult due to poor

**Figure 4** Male, 18-years-old, Oi type iii.

**Notes:** (A, B) Nonunion at the site of osteotomies in the left humerus around an intramedullary rod. Patient previously treated with pamidronate.

**Abbreviation:** OI, osteogenesis imperfecta.
bone quality. Recently, stabilization of long-bone nonunion fractures was performed through compressed sandwich allograft cortical struts method in 12 OI patients (aged 11–78 years) at 13 sites of nonunion fractures. All nonunions healed with allograft incorporation to the bone diaphysis, with all patients recovering to the prefracture level of function, thus reinforcing this approach as durable and safe.

Rehabilitation

An increased incidence of long-bone fractures is the defining feature in individuals with OI. The majority of these fractures occur in early childhood, directly affecting the child’s motor development. The clinical type and severity of OI have a major bearing on functional status, including motor development, range of joint motion, and muscle strength. Children with moderate to severe forms of OI have greater functional limitation, which negatively affects the development of motor function and the level of ambulation when compared to children with mild forms of OI. In addition, gait acquisition is directly affected by joint range of motion and muscle strength. Young adults with type I OI demonstrate full independence, while limitations in self-care, mobility, and domestic life activities were observed in some individuals with OI types IV and V.

To date, there are no physiotherapeutic treatment protocols available for children and adults with OI. A recent study investigated a rehabilitation approach combining resistance training, body-weight-supported treadmill training, and neurodevelopmental treatment associated with side-alternating whole-body vibration in 53 individuals with OI (ages 2.5–24.8 years) for 6 months within a period of 12 months of treatment. There was improvement of mobility between 0 and 6 months of treatment, and also an increase in lean mass and aBMD was observed. However, 46 patients had received BP treatment for years, and the program involved different approaches to treatment.

Further investigation is required, including targeted therapy, the acquisition of specific clinical data, and the evaluation of physiotherapy over short- and long-term courses of intervention. The overall approach to effective physiotherapy for the OI patient will improve when experience is enlarged through the publication of more case reports and cohort studies aimed at outcome assessment.

Conclusion

Research in clinical and genetic fields has expanded speedily. Multiple new genes are now recognized as OI causative, reinforcing the need for extensive clinical investigation focused on improving pharmacological and nonpharmacological therapy linked to these new genetic discoveries. After 20 years of BP therapy in children and adults with OI, questions have arisen about the long-term efficacy of this mode of treatment. In addition, new therapeutic approaches and surgical techniques have been investigated in OI. Efforts are required to improve techniques for gene replacement therapy, which to date has shown only limited success. Nevertheless, there is no cure for OI and the treatment remains based on drug therapy, orthopedic care, rehabilitation, and nutrition.

Disclosure

The authors report no conflicts of interest in this work.

References


90. Sinder BP, Salemi JD, Ominsky MS, Caird MS, Marini JC, Kozloff KM. Rapidly growing Brl/+ mouse model of osteogenesis imperfecta improves bone mass and strength with sclerostin antibody treatment. Bone. 2015;71:115–123.


