

# Concert Genetic Testing: Orthopedics

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Reference Number: V2.2025  
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[Coding implications](#)  
[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

## OVERVIEW

This policy addresses the use of tests for rare skeletal dysplasias and other bone disorders. Pre-test and post-test genetic counseling that facilitates informed decision-making, addresses the possibility of secondary or incidental findings, and a plan for returning results before testing occurs is strongly advised.

For additional information see the [Rationale](#) section.

## POLICY REFERENCE TABLE

### Coding Implications

This clinical policy references Current Procedural Terminology (CPT®). CPT is a registered trademark of the American Medical Association. All CPT codes and descriptions are copyrighted 2024, American Medical Association. All rights reserved. CPT codes and CPT descriptions are from the current manuals and those included herein are not intended to be all-inclusive and are included for informational purposes only. Codes referenced in this clinical policy are for informational purposes only and may not support medical necessity. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

The tests, CPT codes, and ICD codes referenced in this policy are not comprehensive, and their inclusion does not represent a guarantee of coverage or non-coverage. Please see the [Concert Platform](#) for additional registered tests.

**NOTE: Coverage is subject to each requested code's inclusion on the corresponding LDH fee schedule. Non-covered codes are denoted (\*) and are reviewed for Medical Necessity for members under 21 years of age on a per case basis. The non-covered codes will only be denoted in the table below and not throughout the policy. Please only reference the policy reference table for covered and non-covered codes.**

<u>CRITERIA SECTIONS</u>	<b>EXAMPLE TESTS (LABS)</b>	<b>COMMON BILLING CODES</b>	<u>REF</u>
<b><u>Osteogenesis Imperfecta (OI)</u></b>			
<u>Osteogenesis Imperfecta</u>	Osteogenesis imperfecta COL1A1 & COL1A2 NGS Panel (HNL Genomics)	81406*, 81408*, 81479, Q78.0, Z82.79	3
	Osteogenesis Imperfecta Panel (PreventionGenetics, part of Exact Sciences)		
	Osteogenesis Imperfecta NGS Panel - Dominant & Recessive (HNL Genomics)		
<b><u>Skeletal Dysplasias and Rare Bone Disorders</u></b>			
<u>Multigene Panel Analysis for Skeletal Dysplasia or Rare Bone Disorder</u>	Skeletal Disorders Panel (Invitae)	81400*, 81401*, 81402*, 81403*, 81404*, 81405*, 81406*, 81407*, 81408*, 81479, M85, Q77, Q78	1, 7, 8
	Skeletal Dysplasia Core & Extended NGS Panel (HNL Genomics)		
	Comprehensive Skeletal Dysplasias and Disorders Panel (Blueprint Genetics)		
<b><u>Other Covered Skeletal Dysplasias and Rare Bone Disorders</u></b>			
<u>Other Covered Skeletal Dysplasias and Rare Bone Disorders</u>	See list below	81400*, 81401*, 81402*, 81403*, 81404*, 81405*, 81406*, 81407*, 81408*, 81479, M85, Q77, Q78	2, 4, 5, 6

## RELATED POLICIES

This policy document provides criteria for testing related to skeletal dysplasia and rare bone disorders. Please refer to:

- **Specialty Testing: Cardiovascular** for criteria related to diagnostic tests for inherited and sporadic cardiovascular conditions.
- **Specialty Testing: Multisystem Genetic Conditions** for criteria related to diagnostic tests for genetic disorders that affect multiple organ systems (e.g. whole exome and genome sequencing, chromosomal microarray, and multigene panels for broad phenotypes).
- **General Approach to Laboratory Testing** for criteria related to skeletal dysplasias and rare bone disorders, including known familial variant testing, that is not specifically discussed in this or another non-general policy.

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## CRITERIA

It is the policy of Louisiana Healthcare Connections that that the specific genetic testing noted below is **medically necessary** when meeting the related criteria:

### OSTEOGENESIS IMPERFECTA (OI)

#### Osteogenesis Imperfecta

- I. *COL1A1* and *COL1A2* variant analysis or multigene panel analysis that includes *COL1A1* and *COL1A2* to establish or confirm a diagnosis of osteogenesis imperfecta (OI) is considered **medically necessary** when:
  - A. The member/enrollee has any of the following:
    1. Fractures with minimal or no trauma in the absence of other factors, such as [non-accidental trauma \(NAT\)](#) or other known disorders of bone, **OR**
    2. Short stature, often with bone deformity, **OR**
    3. Blue/gray scleral hue, **OR**
    4. Dentinogenesis imperfecta (DI), **OR**
    5. Progressive, postpubertal hearing loss, **OR**
    6. Ligamentous laxity or other signs of connective tissue abnormality, **OR**
    7. Family history of OI, **OR**
    8. Fractures of varying ages and stages of healing (often of the long bones), **OR**
    9. “Codfish” vertebrae, **OR**
    10. Wormian bones, **OR**
    11. Protrusio acetabuli, **OR**

12. Low bone mass or osteoporosis.
- II. Current evidence does not support *COL1A1* and *COL1A2* variant analysis or multigene panel analysis that includes *COL1A1* and *COL1A2* to establish or confirm a diagnosis of osteogenesis imperfecta for all other indications.

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## SKELETAL DYSPLASIAS AND RARE BONE DISORDERS

### Multigene Panel Analysis For Skeletal Dysplasia Or Rare Bone Disorder

- I. Multigene panel analysis to confirm or establish a post-natal diagnosis of a skeletal dysplasia or a rare bone disorder may be considered **medically necessary** when:
  - A. The differential diagnosis includes more than one type of skeletal dysplasia or bone disorder, **AND**
  - B. The member/enrollee displays one or more of the following clinical features of a skeletal dysplasia:
    1. Prenatal ultrasound that showed shortening of the bones of the arms and legs more than 3 standard deviations below the mean, **OR**
    2. Prenatal ultrasound that showed head circumference greater than 75th percentile, **OR**
    3. Prenatal ultrasound that showed bone irregularities (e.g., bowed, fractured, thickened, thin, undermineralized, etc.), **OR**
    4. Prenatal ultrasound that showed abnormal ribs or a small chest circumference, **OR**
    5. Postnatal short stature with height or length less than 3<sup>rd</sup> percentile.
- II. Current evidence does not support multigene panel analysis to confirm or establish a diagnosis of a skeletal dysplasia or a rare bone disorder for all other indications.

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## OTHER COVERED SKELETAL DYSPLASIAS AND RARE BONE DISORDERS

### Other Covered Skeletal Dysplasias and Rare Bone Disorders

The following is a list of conditions that have a known genetic association. Due to their relative rareness, it may be appropriate to cover these genetic tests to establish or confirm a diagnosis.

- I. Genetic testing to establish or confirm one of the following skeletal dysplasias or rare bone disorders to guide management is considered **medically necessary** when the

member/enrollee demonstrates clinical features<sup>1</sup> consistent with the disorder (the list is not meant to be comprehensive, see II below):

- A. Achondroplasia Group
  - 1. [Achondroplasia](#)
  - 2. [Hypochondroplasia](#)
  - 3. [Thanatophoric Dysplasia](#)
- B. [Type II Collagenopathies](#)
  - 1. [Hypochondrogenesis](#)
  - 2. [Spondyloepiphyseal Dysplasia](#)
- C. Type XI Collagen Disorders
  - 1. [Fibrochondrogenesis](#)
  - 2. [Otospondylomegaepiphyseal Dysplasia \(OSMED\)](#)
- D. Sulfation Disorders
  - 1. [Achondrogenesis IB](#)
  - 2. [Atelosteogenesis II](#)
  - 3. [Diastrophic Dysplasia](#)
  - 4. [Chondrodysplasia with Congenital Joint Dislocations](#)
- E. Filamin Disorders and Similar Disorders
  - 1. [Atelosteogenesis Type I](#)
  - 2. [Atelosteogenesis Type III](#)
  - 3. [Larsen Syndrome](#)
  - 4. [Spondylo-Carpal-Tarsal Dysplasia](#)
- F. Short-Rib Dysplasias (with and without Polydactyly)
  - 1. [Chondroectodermal Dysplasia \(Ellis-van Creveld \(EVC\)\)](#)
  - 2. [Short-Rib Polydactyly Syndrome I, II, III, IV including Asphyxiating Thoracic Dystrophy](#)
- G. Metaphyseal Dysplasias
  - 1. [Cartilage-Hair Hypoplasia](#)
- H. Spondylo-Epi-(Meta)-Physeal Dysplasia
  - 1. SEMD, Short Limb Abnormal Calcification Type
- I. Acromesomelic Disorders
  - 1. Acromesomelic Dysplasia, Type Maroteaux
- J. Mesomelic and Rhizo-Mesomelic Dysplasias
  - 1. [Langer Type \(Homozygous Dyschondrosteosis\)](#)
- K. Bent Bone Dysplasias
  - 1. [Campomelic Dysplasia](#)
  - 2. Stuve-Wiedemann Dysplasia
  - 3. Bent Bone Dysplasia FGFR2 Type
- L. Slender Bone Dysplasia
  - 1. [Microcephalic Osteodysplastic Primordial Dwarfism](#)
  - 2. Osteocraniostenosis
- M. Neonatal Osteosclerotic Dysplasias

1. Bloomstrand Dysplasia
  2. [Caffey Disease \(Infantile\)](#)
  3. Raine Dysplasia
- N. Increased Bone Density Group
1. [Osteopetrosis](#)
- O. Abnormal Mineralization Group
1. [Hypophosphatasia](#)
- P. Multiple Epiphyseal Dysplasia and Pseudoachondroplasia Group
1. [Multiple Epiphyseal Dysplasia \(MED\) - Autosomal Dominant](#)
  2. [Multiple Epiphyseal Dysplasia \(MED\) - Autosomal Recessive](#)
  3. [Stickler Syndrome](#)
- Q. [Hereditary Multiple Osteochondromas](#)
- II. Genetic testing to establish or confirm the diagnosis of all other skeletal dysplasias or rare bone disorders not specifically discussed within this or another medical policy will be evaluated by the criteria outlined in *General Approach to Laboratory Testing* (see policy for criteria).

<sup>1</sup>Clinical features for a specific disorder may be outlined in resources such as [GeneReviews](#), [OMIM](#), [National Library of Medicine](#), [Genetics Home Reference](#), or other scholarly sources.

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## RATIONALE

### Osteogenesis Imperfecta

*GeneReviews: COL1A1/2 Osteogenesis Imperfecta*

*GeneReviews is an expert-authored review of current literature on a genetic disease, and goes through a rigorous editing and peer review process before being published online.*

The recommended diagnostic testing for osteogenesis imperfecta is as follows:

COL1A1/2 osteogenesis imperfecta (OI) should be suspected in individuals with the following clinical, radiographic, and laboratory features.

- Fractures with minimal or no trauma in the absence of other factors, such as non-accidental trauma (NAT) or other known disorders of bone
- Short stature or stature shorter than predicted based on stature of unaffected family members, often with bone deformity
- Blue/gray scleral hue

- Dentinogenesis imperfecta (DI)
- Progressive, postpubertal hearing loss
- Ligamentous laxity and other signs of connective tissue abnormality
- Family history of OI, usually consistent with autosomal dominant inheritance

Radiographic features of OI change with age. The major findings include the following:

- Fractures of varying ages and stages of healing, often of the long bones but may also rarely involve ribs and skull. Metaphyseal fractures can be seen in a very small number of children with OI. Rib fractures are much more common in NAT than in OI.
- "Codfish" vertebrae, which are the consequence of spinal compression fractures, seen more commonly in adults.
- Wormian bones, defined as "sutural bones which are 6 mm by 4 mm (in diameter) or larger, in excess of ten in number, with a tendency to arrange in a mosaic pattern." Wormian bones are suggestive of but not pathognomonic for OI.
- Protrusio acetabuli, in which the socket of the hip joint is too deep and the acetabulum bulges into the cavity of the pelvis causing intrapelvic protrusion of the acetabulum.
- Low bone mass or osteoporosis detected by dual energy x-ray absorptiometry (DEXA). Bone density can be normal, especially in individuals with OI type I, as DEXA measures mineral content rather than collagen.

“A multigene panel that includes *COL1A1*, *COL1A2*, and other genes of interest is most likely to identify the genetic cause of the condition at the most reasonable cost while limiting identification of variants of uncertain significance and variants in genes that do not explain the underlying phenotype.”

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### **Multigene Panel Analysis For Skeletal Dysplasia Or Rare Bone Disorder**

*Krakow et al 2009*

A guideline for prenatal diagnosis of fetal skeletal dysplasias (Krakow, Lachman, Rimoin, 2009) recommends the following criteria:

- Fetuses with long bone measurements at or less than the 5th centile or greater than 3 SD below the mean should be evaluated in a center with expertise in the recognition of skeletal dysplasias. If the patient cannot travel, arrangements may be able to be made for evaluation of ultrasound videotapes or hard copy images.
- The following fetal ultrasound measurements should be visualized and plotted against normative values: fetal cranium (biparietal diameter and head circumference), facial profile, mandible, clavicle, scapula, chest circumference, vertebral bodies, all fetal long bones, and the hands and feet. Fetuses with long bone parameters more than 3 SD below the mean should be strongly suspected of having a skeletal dysplasia, especially if the head circumference is greater than the 75th centile

- Lethality should be determined by chest circumference to abdominal circumference ratio and/or femur length to abdominal circumference measurement ratio. A chest-to abdominal circumference ratio of less than 0.6 or femur length to abdominal circumference ratio of 0.16 strongly suggests a perinatal lethal disorder, although there are exceptions. The findings should be conveyed to the physicians caring for the patient and to the patient (p. 5).

In addition, close attention should be paid to the shape and mineralization pattern of the fetal calvarium and fetal skeleton (poor or ectopic mineralization). Determining the elements of the skeleton that are abnormal, coupled with the findings of mineralization and shape of the bones can aid in diagnosis (p. 3).

*American College of Medical Genetics and Genomics (ACMG)*

For diagnosis of genetic causes of short stature, the American College of Medical Genetics clinical practice resource for evaluation of short stature (Seaver et al, 2009) is as follows:

“The definition most commonly used for short stature is height-for-age less than two standard deviations below average for gender, which is demonstrated on the standard growth curves as a length or height less than the 3rd centile” (p. 466).

*Scocchia, et al.*

A 2021 study of the clinical utility of multigene panel testing for an unselected population of individuals with suspected skeletal dysplasia demonstrated a high diagnostic yield in individuals with a suspected skeletal dysplasia or growth disorder (p. 1).

A molecular diagnosis was established in 42% of patients (228/543). Diagnostic variants were identified in 71 genes, with variation in nearly half of these genes contributing to a molecular diagnosis for a single patient in this cohort. Overall, the most common genes in which molecular diagnoses were identified included: *COL2A1* associated with type II collagenopathies; *FGFR3* associated with achondroplasia, thanatophoric dysplasia, hypochondroplasia, and other conditions such as FGFR-related craniosynostoses; and *COL1A1* or *COL1A2*, associated with osteogenesis imperfecta. Together, these four genes accounted for over one third of all molecular diagnoses across the cohort (p. 2-3).

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## Other Covered Skeletal Dysplasias and Rare Bone Disorders

*International Skeletal Dysplasia Society*

The International Skeletal Dysplasia Society published an updated categorization of skeletal dysplasias (Unger, 2023):

“The ‘Nosology of genetic skeletal disorders’ has undergone its 11th revision and now contains 771 entries associated with 552 genes reflecting advances in molecular delineation of new disorders thanks to advances in DNA sequencing technology....As with the previous versions, the list of disorders and genes in the Nosology may be useful in considering the differential diagnosis in the clinic, directing bioinformatic analysis of next-generation sequencing results, and provide a basis for novel advances in biology and medicine” (p. 1165).

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## DEFINITIONS

1. **Non-accidental Trauma (NAT)** - This refers to injury that is purposely inflicted upon a child (e.g., child abuse). NAT often occurs as injury to the skin and soft tissue, but approximately a third of NATs are fractures.

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Reviews, Revisions, and Approvals	Revision Date	Approval Date	Effective Date
Converted corporate to local policy	09/23	11/27/23	
Semi-annual review. Updated title to reflect V1.2024 version. Overview, coding, reference-table, background and references updated. Throughout policy: replaced “coverage criteria” with “criteria. For Policy Reference Table: added “81401, 81402, 81403, 81404, 81405, 81406, 81407”. For Other Related Policies: added “and Molecular”. For Criteria; under Multigene Panel Analysis for Skeletal Dysplasia or Rare Bone Disorder Panel: added “post-natal”; for Other Covered Skeletal Dysplasia and Rare Bone Disorders Panel: under I.O.I. removed “(also in Metabolic Policy)”; under II. added “and Molecular”.	12/23	2/27/24	
Semi-annual review. Updated title to reflect V2.2024 version. In Overview and Clinical Considerations, policy overview updated to include information from the Clinical Considerations section, which has been consolidated into the Overview section. Minor rewording for clarity throughout. Coding, reference-table, background and references updated.	06/24	9/4/24	10/4/24
Semi-annual review. Updated title to reflect V1.2025 version. Osteogenesis Imperfecta: Updated example tests in Policy Reference Table; Updated GeneReviews copyright dates in Reference list; Streamlined portions of Background and Rationale section for brevity. Multigene Panel Analysis for Skeletal Dysplasia or Rare Bone Disorder: Updated References; Streamlined portions of Background and Rationale section for brevity; Updated tests in Policy Reference Table. Other Covered Skeletal	1/25	3/31/25	5/1/25

Reviews, Revisions, and Approvals	Revision Date	Approval Date	Effective Date
Dysplasias and Rare Bone Disorders: Updated Background and Rationale with information from newer version of reference; updated reference.			
Annual review. Policy name changed from Concert Genetic Testing: Skeletal Dysplasia and Rare Bone Disorders to Concert Genetic Testing: Orthopedics. Policy reference table, rationale, background, and references updated. “Investigational” policy statements changed to state “current evidence does not support.”	03/26	5/27/26	6/26/26

## REFERENCES

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### **Important Reminder**

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. LHCC makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved.

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by providing services to member/enrollees and/or submitting claims for payment for such services.

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