



Discover Dysplasias™ provides no-charge genetic testing for patients age 16 and younger in the US and Canada with short stature & signs and symptoms suggestive of, or consistent with, a diagnosis of skeletal dysplasia. Discover Dysplasias™ can help facilitate diagnosis and in some cases, put patients on the path to disease-specific management sooner.<sup>1,2</sup>



### Who it's for

Eligible patients aged 16 years and younger in the US and Canada with short stature and  $\geq 1$  sign/symptom associated with genetic skeletal dysplasia\*



### Genes tested

Over 350 genes including those associated with the most common skeletal dysplasias<sup>3</sup>



### Sample type

3 different sample collection methods (whole blood, saliva, buccal swab)



### How to order

Visit [DiscoverDysplasias.com](https://DiscoverDysplasias.com) or contact Invitae Client Services at [clientservices@invitae.com](mailto:clientservices@invitae.com) or 1-800-436-3037



### Receiving results

Once Invitae receives the sample, results are typically available within 10-21 days (14 days on average)



### Genetic counseling

Expert genetic counselors are available to help clinicians and patients review results and specific cases<sup>4</sup>

Order a sample collection kit today at [DiscoverDysplasias.com](https://DiscoverDysplasias.com)



\*specific signs and symptoms listed at [DiscoverDysplasias.com](https://DiscoverDysplasias.com)

References: 1. Solari P, Clarke L, Deignon B, Raggio C, Wood T, Pollard L. Increasing awareness and earlier testing for mucopolysaccharidoses to improve patient outcomes: updated results from Simply Test for MPS™ enzyme-panel program (ENCORE). Poster presented at WORLDSymposium™ 2020; February 10-13, 2020; Orlando, FL. 2. Souza CFM, Siqueira AC, Antunes NS, et al. Perthes-like disease masquerading as non-classical MPS. J Inborn Errors Metab Screen. 2020;8(e20200003):1-11. doi:10.1590/2326-4594-JIEMS-2020-0003. 3. Krakow D. Skeletal dysplasias. Clin Perinatol. 2015;42(2):301-319. doi:10.1016/j.clp.2015.03.003. 4. Seratti G, Pansare V, Pang TY, et al. Clinical utility of a sponsored, no-cost skeletal dysplasia gene panel testing program: results from 850 tests. Poster presented at the 17th Annual WorldSymposium™; February 8-12, 2021.