Advancing care and research for chordoma together

CHORDOMA FOUNDATION

Resources for physicians and researchers

Chordoma is diagnosed in just 1 in 1,000,000 people per year. The Chordoma Foundation is here to support your efforts to achieve the best possible outcomes for every patient. We offer trusted, evidence-based clinical guidance, patient support services in multiple languages, and a range of research tools to accelerate discovery and improve care. Since 2007, we've assembled the world's largest collection of chordoma models, samples, and other research tools, and partnered with clinicians and researchers around the globe to fast-track the development of new therapies.

Clinical resources

Evidence-based diagnosis and treatment guidance

- Chordoma Global Consensus Group papers on diagnosis, primary and recurrent disease
- NCCN and ESMO bone-sarcoma guidelines
- Clinician-friendly summaries available

Clinical trial finder

A curated list of drug therapies with scientific rationale for chordoma, including trials open worldwide, is designed to save time when discussing treatment options with patients.

Supporting your research

Chordoma Foundation Biobank

With over 250 chordoma tumor samples collected, our biobank offers immediate access to well-annotated specimens, and models for target validation, drug screening, and biomarker discovery — accelerating your translational research and derisking the path from the laboratory to clinical trials.

Chordoma Foundation Labs

Located in North Carolina in the US, our dedicated laboratory can provide you with rapid, go/no-go preclinical studies using chordoma cell lines; expert support to accelerate drug development; and cost-effective, collaborative partnerships tailored to your goals.

Clinical trial development

Our team partners with investigators to design and launch clinical trials, offering help with protocols, feasibility, recruitment, and funding.

Global collaboration

Join our global network of chordoma researchers and clinicians at events like our International Chordoma Research Workshop to exchange insights, advance ideas, and forge collaborations.

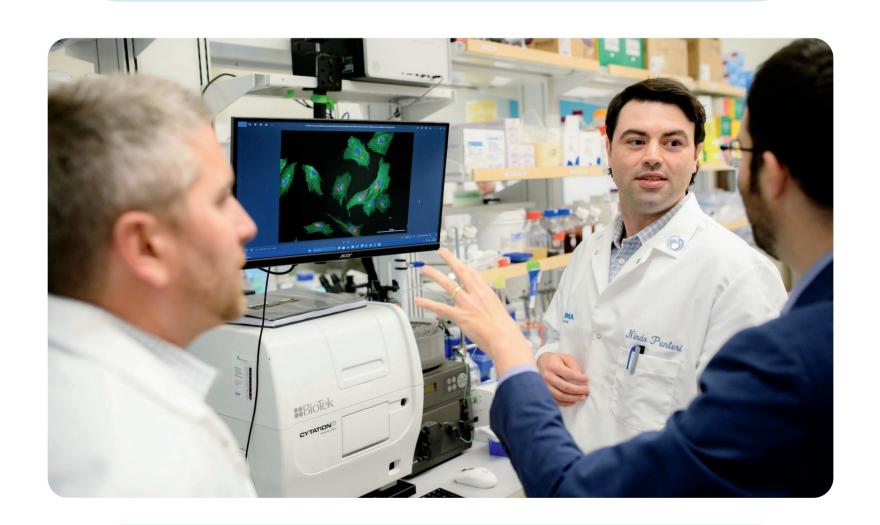
Helping your patients navigate chordoma

Multilingual patient navigation

Our navigators provide personalized support to help patients find experienced treatment and clinical trials, and connect them with peer support.

Survivorship and quality of life

Our survivorship program offers chordoma patients in-depth guidance on managing treatment side effect, and functional challenges-developed with input from patients and clinicians.



Contact us

Resources for physicians: support@chordoma.org

Resources for researchers: research@chordoma.org



Email signup

Scan the QR code to join our mailing list and hear about new resources for clinicians and researchers.