Resource Summary Report

Generated by Kravitz on Sep 10, 2026

Cincinnati Children's Hospital Discover Together Biobank Core Facility

RRID:SCR_022632
Type: Tool

Proper Citation

Cincinnati Children's Hospital Discover Together Biobank Core Facility (RRID:SCR_022632)

Resource Information

URL: https://www.cincinnatichildrens.org/service/c/clinical-trials/biobank

Proper Citation: Cincinnati Children's Hospital Discover Together Biobank Core Facility (RRID:SCR_022632)

Description: Provides access to services for standardized and centralized acquisition, processing, storage and distribution of biospecimens for research.

Synonyms: Cincinnati Children's Hospital Discover Together Biobank Core, Discover Together Biobank

Resource Type: access service resource, core facility, service resource

Keywords: USEDit, ABRF, biospecimens standardized and centralized acquisition, biospecimens processing, biospecimens storage and distribution

Resource Name: Cincinnati Children's Hospital Discover Together Biobank Core Facility

Resource ID: SCR_022632

Alternate IDs: ABRF_1481

Alternate URLs: https://coremarketplace.org/?FacilityID=1481&citation=1

Record Creation Time: 20220803T050137+0000

Record Last Update: 20260905T133428+0000

Ratings and Alerts

No rating or validation information has been found for Cincinnati Children's Hospital Discover Together Biobank Core Facility.

No alerts have been found for Cincinnati Children's Hospital Discover Together Biobank Core Facility.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We found 5 mentions in open access literature.

Listed below are recent publications. The full list is available at Kravitz .

Theobald K, et al. (2026) Expanding Access to Genome Sequencing: Higher Diagnostic Yield in Self-Referred Participants From the CincyKidsSeq Study and Implications for Hybrid Models of Genetic Service Delivery. Clinical genetics, 109(4), 717.

Buasri K, et al. (2026) Genotypic and phenotypic spectrum of hereditary spastic paraplegia 56: insights from novel CYP2U1 variants and a literature review. BMC neurology, 26(1).

Kottyan LC, et al. (2026) Sequencing and health data resource of children of African ancestry. Genetics in medicine : official journal of the American College of Medical Genetics, 28(3), 101660.

Kottyan LC, et al. (2025) Sequencing and health data resource of children of African ancestry. medRxiv : the preprint server for health sciences.

Marsili L, et al. (2024) Studying Rare Movement Disorders: From Whole-Exome Sequencing to New Diagnostic and Therapeutic Approaches in a Modern Genetic Clinic. Biomedicines, 12(12).