Resource Summary Report

Generated by T1D on Aug 9, 2026

University of Cologne Center for Genomics (CCG) Core Facility

RRID:SCR_027822
Type: Tool

Proper Citation

University of Cologne Center for Genomics (CCG) Core Facility (RRID:SCR_027822)

Resource Information

URL: https://ccg.uni-koeln.de/

Proper Citation: University of Cologne Center for Genomics (CCG) Core Facility (RRID:SCR_027822)

Description: Represents central facility and does not operate for profit, engaging in academic research collaborations and service provision on collaborative basis. Provides infrastructure for all kinds of sequencing, gene mapping and whole genome association studies in complex diseases as well as the validation of copy number variations.

Synonyms: , University of Cologne Center for Genomics (CCG), Cologne Center for Genomics (CCG)

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

Keywords: ABRF, sequencing, gene mapping, whole genome association studies,

Availability: Restricted

Resource Name: University of Cologne Center for Genomics (CCG) Core Facility

Resource ID: SCR_027822

Alternate IDs: ABRF_5714

Alternate URLs: https://coremarketplace.org/RRID:SCR_027822/?citation=1

Record Creation Time: 20251219T055114+0000

Record Last Update: 20260808T190815+0000

Ratings and Alerts

No rating or validation information has been found for University of Cologne Center for Genomics (CCG) Core Facility.

No alerts have been found for University of Cologne Center for Genomics (CCG) Core Facility.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We found 1 mentions in open access literature.

Listed below are recent publications. The full list is available at T1D .

J??schke C, et al. (2026) CACNB3 defects are associated with infantile idiopathic nystagmus. Brain communications, 8(2), fcag034.