

Resource Summary Report

Generated by [NIF](#) on Aug 11, 2026

University of Chicago Proteomics Platform Core Facility

RRID:SCR_022928

Type: Tool

Proper Citation

University of Chicago Proteomics Platform Core Facility (RRID:SCR_022928)

Resource Information

URL: <https://voices.uchicago.edu/proteomics/>

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Description: Core to develop and utilize proteomics techniques to answer both basic and translational scientific questions.

Synonyms: University of Chicago Proteomics Platform, Proteomics Platform

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

Keywords: USEDit, ABRF

Availability: Open

Resource Name: University of Chicago Proteomics Platform Core Facility

Resource ID: SCR_022928

Alternate IDs: ABRF_1607

Alternate URLs: <https://coremarketplace.org/?FacilityID=1607&citation=1>

Record Creation Time: 20221028T050209+0000

Record Last Update: 20260811T043714+0000

Ratings and Alerts

No rating or validation information has been found for University of Chicago Proteomics Platform Core Facility.

No alerts have been found for University of Chicago Proteomics Platform Core Facility.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 4 mentions in open access literature.

Listed below are recent publications. The full list is available at [NIF](#) .

Hajirahimkhan A, et al. (2026) Reprogramming SREBP1-dependent lipogenesis and inflammation in high-risk breast with licochalcone A: A novel path to cancer prevention. International journal of cancer, 158(7), 1927.

Song X, et al. (2024) RNA splicing analysis deciphers developmental hierarchies and reveals therapeutic targets in adult glioma. The Journal of clinical investigation, 134(11).

Mousaei Ghasroldasht M, et al. (2024) A Comparative Analysis of Na⁺⁺ve Exosomes and Enhanced Exosomes with a Focus on the Treatment Potential in Ovarian Disorders. Journal of personalized medicine, 14(5).

Hou Y, et al. (2024) Engineered mischarged transfer RNAs for correcting pathogenic missense mutations. Molecular therapy : the journal of the American Society of Gene Therapy, 32(2), 352.