

Resource Summary Report

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University of Michigan Medical School BRCF Flow Cytometry Core Facility

RRID:SCR_026697

Type: Tool

Proper Citation

University of Michigan Medical School BRCF Flow Cytometry Core Facility
(RRID:SCR_026697)

Resource Information

URL: <https://medresearch.umich.edu/office-research/about-office-research/biomedical-research-core-facilities/flow-cytometry-core>

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Description: Flow Cytometry Core provides instrumentation, expertise and services. Samples are prepared by individual investigators, who then deliver to the Core for flow cytometric analysis or cell sorting. The Core also provides assistance in grant and publication preparation, publication-quality graphics, and the development of experimental designs.

Synonyms: University of Michigan BRCF Flow Cytometry

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

Keywords: ABRF, flow cytometry, instrumentation, expertise, services, assistance in grant and publication preparation,

Availability: Open

Resource Name: University of Michigan Medical School BRCF Flow Cytometry Core Facility

Resource ID: SCR_026697

Alternate IDs: ABRF_3191

Alternate URLs: https://coremarketplace.org/RRID:SCR_026697/?citation=1

Record Creation Time: 20250403T060323+0000

Ratings and Alerts

No rating or validation information has been found for University of Michigan Medical School BRCF Flow Cytometry Core Facility.

No alerts have been found for University of Michigan Medical School BRCF Flow Cytometry Core Facility.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 2 mentions in open access literature.

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

Shafer AM, et al. (2025) Immunocompetent Murine Models Recapitulate the Heterogeneous Tumor-Immune Microenvironment of Human Liposarcoma. Clinical cancer research : an official journal of the American Association for Cancer Research, 31(23), 5078.

Dennison DD, et al. (2025) Ubiquitin chain variability directs substrates of the Tul1 ubiquitin ligase complex to different degradation pathways. The Journal of cell biology, 224(9).