

Resource Summary Report

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University of Utah Flow Cytometry Core Facility

RRID:SCR_012210

Type: Tool

Proper Citation

University of Utah Flow Cytometry Core Facility (RRID:SCR_012210)

Resource Information

URL: <http://www.scienceexchange.com/facilities/flow-cytometry-core-facility-utah>

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Description: Serving 120+ investigators with a 5 laser high speed sorter and 3 benchtop analyzers with 13+ color capabilities, the Utah Flow Cytometry Core Facility has developed a paradigm to balance productivity with quality to minimize the cost per research project, keep overall costs contained, and provide the necessary scientific support. Central to this paradigm is a close working relationship with investigators to define their projects in the early stages of development to make optimal and efficient use of flow cytometry. In addition to high quality and well maintained instrumentation, success of this approach requires focused efforts in three major areas: 1) education of ALL users (faculty and staff) in the science and technology of flow cytometry, 2) active involvement in the scientific development of the project, 3) continuing education for core staff.

Abbreviations: UUtah Flow Cytometry Core Facility

Synonyms: Utah Flow Cytometry Core Facility

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

Resource Name: University of Utah Flow Cytometry Core Facility

Resource ID: SCR_012210

Alternate IDs: SciEx_10681

Record Creation Time: 20220129T080309+0000

Record Last Update: 20260731T042842+0000

Ratings and Alerts

No rating or validation information has been found for University of Utah Flow Cytometry Core Facility.

No alerts have been found for University of Utah Flow Cytometry 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 [PRECISE-TBI](#) .

Fang P, et al. (2025) Differential control of growth and identity by HNF4?? isoforms in pancreatic ductal adenocarcinoma. Molecular cancer research : MCR.