

Resource Summary Report

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University of Colorado Anschutz Medical Campus Cancer Center Drug Discovery and Development Shared Resource Core Facility

RRID:SCR_021986

Type: Tool

Proper Citation

University of Colorado Anschutz Medical Campus Cancer Center Drug Discovery and Development Shared Resource Core Facility (RRID:SCR_021986)

Resource Information

URL:

https://cu.corefacilities.org/service_center/show_external/5637/drug_discovery_and_development_s

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Description: Provides expertise and infrastructure to facilitate drug discovery at different stages of development pipeline, including assay development and validation, high throughput screening and image based high content screening discovery, validation and optimization, and in vitro and in vivo pharmacology including pharmacokinetics and pharmacodynamics as well as toxicology. Sign in to iLab using University of Colorado credentials.

Abbreviations: D3SR

Synonyms: Drug Discovery and Development Shared Resource

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

Keywords: ABRF, USEDit

Resource Name: University of Colorado Anschutz Medical Campus Cancer Center Drug Discovery and Development Shared Resource Core Facility

Resource ID: SCR_021986

Alternate IDs: ABRF_1308

Alternate URLs: <https://coremarketplace.org/?FacilityID=1308>

Record Creation Time: 20220421T050138+0000

Record Last Update: 20260815T183143+0000

Ratings and Alerts

No rating or validation information has been found for University of Colorado Anschutz Medical Campus Cancer Center Drug Discovery and Development Shared Resource Core Facility.

No alerts have been found for University of Colorado Anschutz Medical Campus Cancer Center Drug Discovery and Development Shared Resource 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](#) .

Klosowski ML, et al. (2025) Breast cancer cell coculture induces normal lung fibroblast transition to CAFs, promoting tumor cell dormancy and therapy resistance. Proceedings of the National Academy of Sciences of the United States of America, 122(47), e2423894122.

Rinaldetti S, et al. (2022) High-Content Drug Discovery Targeting Molecular Bladder Cancer Subtypes. International journal of molecular sciences, 23(18).