

Resource Summary Report

Generated by SPARC SAWG on Sep 12, 2026

University of California at Los Angeles Translational Pathology Core Facility

RRID:SCR_012201

Type: Tool

Proper Citation

University of California at Los Angeles Translational Pathology Core Facility
(RRID:SCR_012201)

Resource Information

URL: <https://www.uclahealth.org/departments/pathology/research-services/translational-pathology-core-laboratory-tpcl>

Proper Citation: University of California at Los Angeles Translational Pathology Core Facility (RRID:SCR_012201)

Description: Facility in the UCLA Department of Pathology and Laboratory Medicine and UCLA Jonsson Comprehensive Cancer Center Shared Facility. Provides pathology related services in support of basic, translational and clinical research at UCLA. Provides consultative services to investigators in pathology-related study design, tissue selection, microscopic interpretation, immunohistochemistry, laser capture microdissection, digital image analysis, and IRB-related tissue questions.

Abbreviations: UCLA TPCL, TPCL

Synonyms: , UCLA Translational Pathology Core Laboratory, Translational Pathology Core Laboratory

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

Resource Name: University of California at Los Angeles Translational Pathology Core Facility

Resource ID: SCR_012201

Alternate IDs: SciEx_10615

Old URLs: <http://www.scienceexchange.com/facilities/translational-pathology-core-laboratory-ucla>

Record Creation Time: 20220129T080309+0000

Ratings and Alerts

No rating or validation information has been found for University of California at Los Angeles Translational Pathology Core Facility.

No alerts have been found for University of California at Los Angeles Translational Pathology 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 [SPARC SAWG](#) .

Campbell KM, et al. (2026) Cellular neighborhoods govern antitumor T-cell infiltration following anti-CTLA-4 in melanoma with primary resistance to anti-PD-1. Cancer discovery.