Resource Summary Report

Generated by ASWG on Apr 29, 2025

Louisiana State University in Shreveport Animal Models and Histology Core Facility

RRID:SCR_024776
Type: Tool

Proper Citation

Louisiana State University in Shreveport Animal Models and Histology Core Facility (RRID:SCR_024776)

Resource Information

URL: https://www.lsuhs.edu/centers/cardiovascular-diseases-and-sciences/cobre/cobre-core-facilities

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Description: Core provides services for mouse genotyping and tissue histology, access to tools for analyzing cardiovascular function, data analysis, expertise and training. Services include genotyping using PCR or RT PCR, histology services with tissue processing, paraffin-embedding and sectioning, along with several cardiovascular relevant histological stains such as Picrosirius Red and Masson Trichrome.

Synonyms: Animal Models and Histology Core, Louisiana State University in Shreveport Animal Models and Histology Core

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

Keywords: ABRF, mouse genotyping, tissue histology, analyzing cardiovascular function, genotyping, PCR, RT PCR, histology services,

Funding: NIGMS COBRE

Resource Name: Louisiana State University in Shreveport Animal Models and Histology Core Facility

Resource ID: SCR_024776

Alternate IDs: ABRF_2568

Alternate URLs: https://coremarketplace.org/?FacilityID=2568&citation=1

Record Creation Time: 20231212T050231+0000

Record Last Update: 20250429T060348+0000

Ratings and Alerts

No rating or validation information has been found for Louisiana State University in Shreveport Animal Models and Histology Core Facility.

No alerts have been found for Louisiana State University in Shreveport Animal Models and Histology 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 ASWG.

Aishwarya R, et al. (2024) Diastolic dysfunction in Alzheimer's disease model mice is associated with A?-amyloid aggregate formation and mitochondrial dysfunction. Scientific reports, 14(1), 16715.