## **Resource Summary Report**

Generated by RRID on May 15, 2025

# University of Pennsylvania Perelman School of Medicine Penn Medicine BioBank Core Facility

RRID:SCR\_022415 Type: Tool

### **Proper Citation**

University of Pennsylvania Perelman School of Medicine Penn Medicine BioBank Core Facility (RRID:SCR\_022415)

## **Resource Information**

URL: https://pmbb.med.upenn.edu/

**Proper Citation:** University of Pennsylvania Perelman School of Medicine Penn Medicine BioBank Core Facility (RRID:SCR\_022415)

**Description:** BioBank supports researchers by providing centralized access to large number of annotated blood and tissue samples.

Abbreviations: PMBB

**Synonyms:** University of Pennsylvania Perelman School of Medicine Penn Medicine BioBank

**Resource Type:** material storage repository, core facility, biobank, service resource, storage service resource, access service resource

Keywords: USEDit, ABRF, annotated blood and tissue samples

#### Funding:

**Resource Name:** University of Pennsylvania Perelman School of Medicine Penn Medicine BioBank Core Facility

Resource ID: SCR\_022415

Alternate IDs: ARBF\_1421

Alternate URLs: https://coremarketplace.org?citation=1&FacilityID=1421

Record Creation Time: 20220602T050140+0000

Record Last Update: 20250514T061935+0000

## **Ratings and Alerts**

No rating or validation information has been found for University of Pennsylvania Perelman School of Medicine Penn Medicine BioBank Core Facility.

No alerts have been found for University of Pennsylvania Perelman School of Medicine Penn Medicine BioBank Core Facility.

## Data and Source Information

Source: SciCrunch Registry

## **Usage and Citation Metrics**

We found 5 mentions in open access literature.

Listed below are recent publications. The full list is available at <u>RRID</u>.

Hui D, et al. (2025) Risk factors affecting polygenic score performance across diverse cohorts. eLife, 12.

Zhang DY, et al. (2024) Protein-truncating variant in APOL3 increases chronic kidney disease risk in epistasis with APOL1 risk alleles. JCI insight, 9(19).

Cappadocia J, et al. (2024) PMS2CL interference leading to erroneous identification of a pathogenic PMS2 variant in Black patients. Genetics in medicine open, 2, 101858.

DePaolo J, et al. (2024) Titin-Truncating variants Predispose to Dilated Cardiomyopathy in Diverse Populations. medRxiv : the preprint server for health sciences.

Klarin D, et al. (2023) Genome-wide association study of thoracic aortic aneurysm and dissection in the Million Veteran Program. Nature genetics, 55(7), 1106.