Resource Summary Report

Generated by PRECISE-TBI on Sep 7, 2026

Augusta University Centre for Biotechnology and Genomic Medicine Bioinformatics Core Facility

RRID:SCR_027092
Type: Tool

Proper Citation

Augusta University Centre for Biotechnology and Genomic Medicine Bioinformatics Core Facility (RRID:SCR_027092)

Resource Information

URL: http://augusta.edu/research/core/bioinformatics.php

Proper Citation: Augusta University Centre for Biotechnology and Genomic Medicine Bioinformatics Core Facility (RRID:SCR_027092)

Description: Core provides bioinformatics support for genomics, transcriptomics, epigenomics, and other high-throughput data. Offers data analysis, pipeline development, and consultation services to researchers across disciplines, with focus on reproducibility, scalability, and collaboration.

Synonyms: CBGM Bioinformatics Core, , Augusta University Centre for Biotechnology and Genomic Medicine Bioinformatics Core

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

Keywords: Bioinformatics, genomics, transcriptomics, proteomics, high-throughput data, data analysis, consultation services, training

Resource Name: Augusta University Centre for Biotechnology and Genomic Medicine Bioinformatics Core Facility

Resource ID: SCR_027092

Alternate IDs: ABRF_3268

Alternate URLs: https://coremarketplace.org/RRID:SCR_027092/?citation=1

Record Creation Time: 20250624T054719+0000

Record Last Update: 20260905T133538+0000

Ratings and Alerts

No rating or validation information has been found for Augusta University Centre for Biotechnology and Genomic Medicine Bioinformatics Core Facility.

No alerts have been found for Augusta University Centre for Biotechnology and Genomic Medicine Bioinformatics 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 .

Murphy SA, et al. (2026) CD73 blockade enhances antitumor efficacy of oHSV in solid tumors by increasing macrophage-mediated antigen presentation. Journal for immunotherapy of cancer, 14(1).