Resource Summary Report

Generated by PRECISE-TBI on Aug 5, 2026

Tufts University; Massachusetts; USA

RRID:SCR_000994
Type: Tool

Proper Citation

Tufts University; Massachusetts; USA (RRID:SCR_000994)

Resource Information

URL: http://www.tufts.edu/

Proper Citation: Tufts University; Massachusetts; USA (RRID:SCR_000994)

Description: Private research university in Medford and Somerville, Massachusetts, United States, with additional facilities in Boston and Grafton, as well as Talloires, France. Provides undergraduate, graduate, and professional degree programs.

Synonyms: Tufts University, Tufts

Resource Type: university

Keywords: tufts, university, undergraduate, graduate, professional

Resource Name: Tufts University; Massachusetts; USA

Resource ID: SCR_000994

Alternate IDs: Crossref funder ID:100008090, ISNI:0000 0004 1936 7531, Wikidata:Q49120, nlx_38314, grid.429997.8

Alternate URLs: https://ror.org/05wvpxv85

Record Creation Time: 20220129T080204+0000

Record Last Update: 20260801T190135+0000

Ratings and Alerts

No rating or validation information has been found for Tufts University; Massachusetts; USA.

No alerts have been found for Tufts University; Massachusetts; USA.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We found 9 mentions in open access literature.

Listed below are recent publications. The full list is available at PRECISE-TBI .

Yin Q, et al. (2018) Long-range haplotype analysis of the malaria parasite receptor gene ACKR1 in an East-African population. Human genome variation, 5, 26.

Al-Naemi AH, et al. (2018) Is the rs1801282 (G/C) Polymorphism of PPAR - Gamma Gene Associated with T2DM in Iraqi People? Open access Macedonian journal of medical sciences, 6(3), 447.

Zurawek M, et al. (2017) MAVS is not a Likely Susceptibility Locus for Addison's Disease and Type 1 Diabetes. Archivum immunologiae et therapiae experimentalis, 65(3), 271.

Al-Bustan SA, et al. (2017) Increased Risk of the APOB rs11279109 Polymorphism for CHD among the Kuwaiti Population. Disease markers, 2017, 6963437.

Dawidowska M, et al. (2016) Association of germline genetic variants in RFC, IL15 and VDR genes with minimal residual disease in pediatric B-cell precursor ALL. Scientific reports, 6, 29427.

Ozbek IC, et al. (2016) Timing of coronary artery bypass surgery in patients with non-ST-segment elevation myocardial infarction and postoperative outcomes. Archives of medical science : AMS, 12(4), 766.

El Khoury L, et al. (2016) MMP3 and TIMP2 gene variants as predisposing factors for Achilles tendon pathologies: Attempted replication study in a British case-control cohort. Meta gene, 9, 52.

Sheahan KL, et al. (2015) Identification of mammalian proteins that collaborate with type III secretion system function: involvement of a chemokine receptor in supporting translocon activity. mBio, 6(1), e02023.

Sen A, et al. (2011) Impact of interleukin-6 promoter polymorphism and serum interleukin-6 level on the acute inflammation and neovascularization stages of patients with Eales' disease. Molecular vision, 17, 2552.