Resource Summary Report

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HmtPhenome

RRID:SCR_017289
Type: Tool

Proper Citation

HmtPhenome (RRID:SCR_017289)

Resource Information

URL: https://www.hmtphenome.uniba.it

Proper Citation: HmtPhenome (RRID:SCR_017289)

Description: Collection of data about variants, genes, phenotypes and diseases involved in mitochondrial functionality. Users can search for variant position, gene, phenotype or disease and retrieve all related information through integrated network of biological entities.

Resource Type: data or information resource, data processing software, data visualization software, software resource, service resource, network graph visualization software, software application, database

Defining Citation: DOI:10.1101/660282

Keywords: mitochondria, variant, gene, function, phenotype, data

Funding:

Availability: Free, Freely available

Resource Name: HmtPhenome

Resource ID: SCR_017289

Record Creation Time: 20220129T080334+0000

Record Last Update: 20250412T060108+0000

Ratings and Alerts

No rating or validation information has been found for HmtPhenome.

No alerts have been found for HmtPhenome.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We have not found any literature mentions for this resource.