

SGD Saccharomyces GENOME DATABASE Analyze Sequence Function Literature Community

The *Saccharomyces* Genome Database (SGD) is the premier genetics & genomics resource for the budding yeast *S. cerevisiae*.

Q varj

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VAR1 / Q0140

varicosity

antigenic variation

Variant Viewer

Vargas, RC, et al. (2004)

Gene

Cellular Component

Biological Process

Resource

Reference

ECM22 / YLR228C Literature

Primary Literature

ECM22 / YLR228C Overview

Standard Name: ECM22

Systematic Name: YLR228C

SGD ID: S00004218

Gene Ontology: OMF, Verified

Description: Sterol regulatory element binding protein, regulates transcription of sterol biosynthetic genes upon sterol depletion, after relocating from intracellular membranes to perinuclear foci, redundant activator of transcription with UPCI, up-regulating the expression of genes involved in biomass growth, controls 200 Cyt5 structural cluster; ECM22 has a paralog, UPCI, that arose from the whole genome duplication 111117

Sequence

ECM22 Location: Chromosome XII 600019..602463

SGD provides free public access to high quality, expertly-curated information on the budding yeast genome and proteome.



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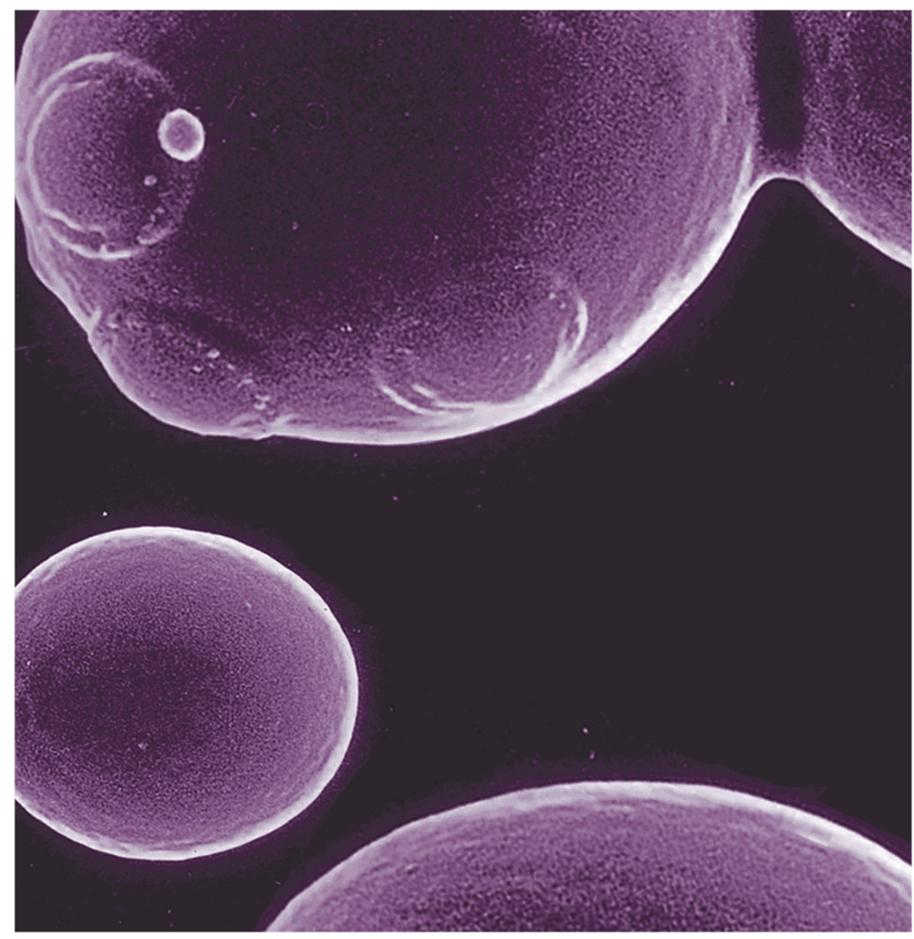
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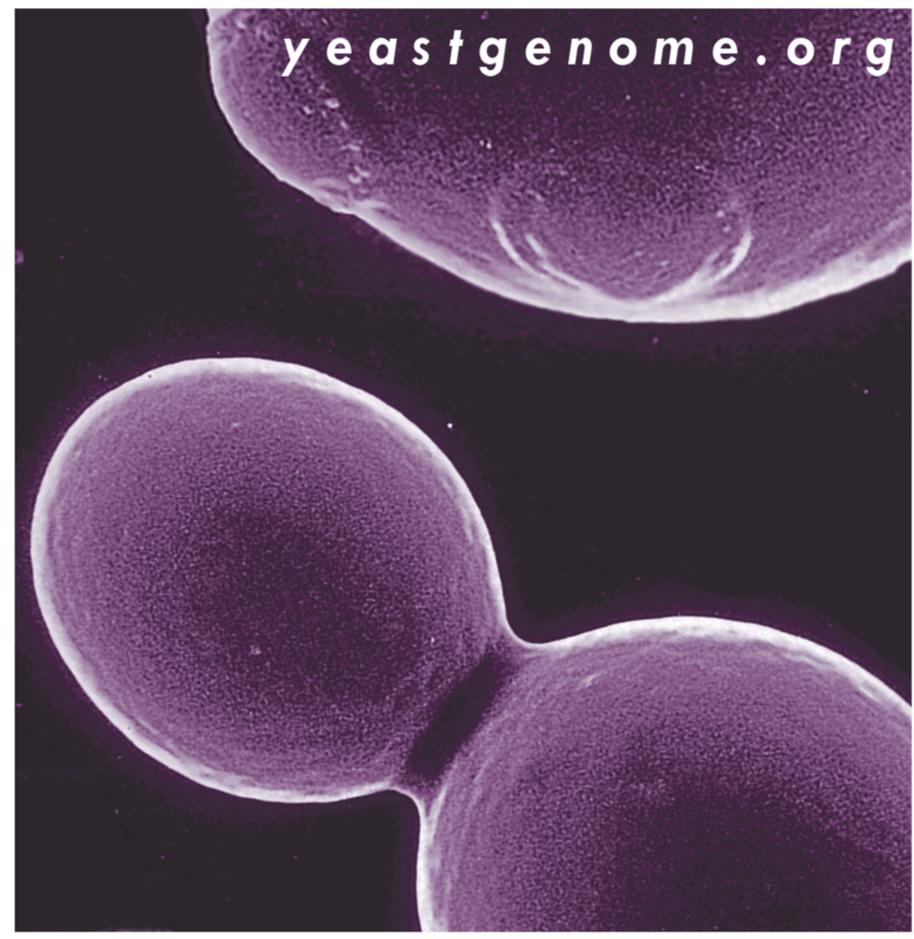
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Stanford University



SGD Saccharomyces GENOME DATABASE



Rich gene summary pages

More data in each tab!

Search for your favorite yeast genes!

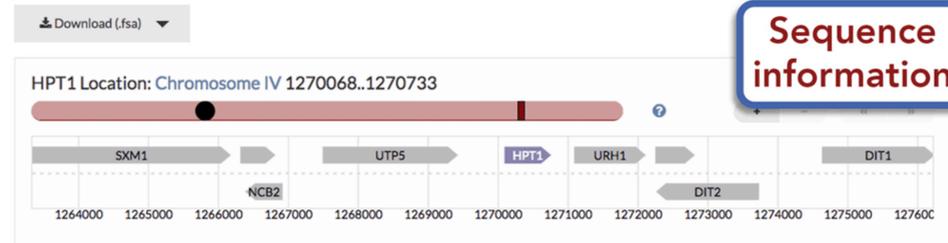
Summary Sequence Protein Gene Ontology Phenotype Interactions Regulation Expression Literature

HPT1 / YDR399W Overview

Standard Name: HPT1¹
 Systematic Name: YDR399W
 SGD ID: S00002807
 Aliases: BRA6², HGPRTase³, HPRT⁴
 Feature Type: ORF, Verified
 Description: Dimeric hypoxanthine-guanine phosphoribosyltransferase; catalyzes the transfer of the phosphoribosyl portion of 5-phosphoribosyl-alpha-1-pyrophosphate to a purine base (either guanine or hypoxanthine) to form pyrophosphate and a purine nucleotide (either guanosine monophosphate or inosine monophosphate); mutations in the human homolog HPRT1 can cause Lesch-Nyhan syndrome and Kelley-Seegmiller syndrome^{3,4,5}
 Name Description: Hypoxanthine guanine PhosphoribosylTransferase⁶

ORF info & description

Sequence Sequence Details



Sequence information

Subfeatures - S288C 1 subfeature

Feature	Relative Coordinates	Coordinates	Coord. Version	Seq. Version
CDS	1..666	chrIV:1270068..1270733	2011-02-03	1996-07-31

Protein Protein Details

Length (a.a.): 221
 Mol. Weight (Da): 25184.9
 Isoelectric Point: 5.40

Protein information

Gene Ontology Gene Ontology Details

View computational annotations

Molecular Function
 Manually Curated: hypoxanthine phosphoribosyltransferase activity (IMP, IDA, IGI)

Biological Process
 Manually Curated: GMP salvage (IDA, IMP, IGI), IMP salvage (IDA)

Cellular Component
 High-Throughput: cytoplasm (IDA), nucleus (IDA)

Molecular functions, biological processes, & cellular components

Pathways

- salvage pathways of guanine, xanthine, and their nucleosides
- salvage pathways of adenine, hypoxanthine, and their nucleosides
- salvage pathways of purines and their nucleosides
- superpathway of purine biosynthesis and salvage pathways

Biochemical pathways

Phenotype

- Classical Genetics
- resistance to cisplatin: increased
- unspecified:
- resistance to 8-azaguanine: decreased
 - viable

Mutant phenotypes & alleles

- Large-scale Survey
- alpha-amino acid accumulation: abnormal
 - auxotrophy
 - resistance to cisplatin: increased
 - resistance to mycophenolic acid: decreased
 - vacuolar morphology: abnormal
 - viable

Interaction Interaction Details

149 total interactions for 98 unique genes

- Physical Interactions
- Affinity Capture-RNA: 4
 - Co-crystal Structure: 1
 - Co-purification: 1
 - Two-hybrid: 1

Genetic & physical interactions

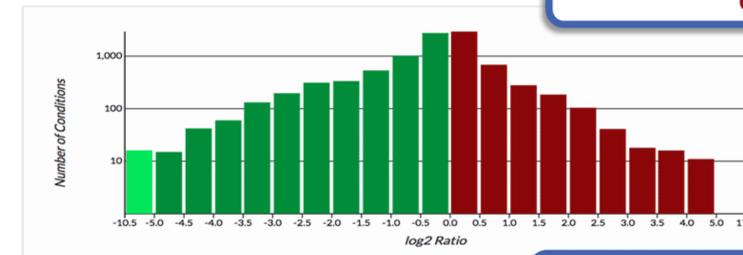
- Genetic Interactions
- Dosage Rescue: 1
 - Negative Genetic: 130
 - Phenotypic Suppression: 5
 - Positive Genetic: 6

Regulation Regulation Details

Regulators: 7
 Targets: 0

Regulation & expression data

Expression



Summary Paragraph

HPT1 encodes hypoxanthine-guanine phosphoribosyltransferase, an enzyme involved in the salvage pathway of purine nucleotide biosynthesis (1). Hpt1p catalyzes the conversion of the purine bases hypoxanthine and guanine to the nucleotides IMP and GMP (5). The enzyme functions as a dimer and can be inhibited *in vitro* by its end-product GMP (4, 6). In *hpt1* null mutants, if the *de novo* pathway of guanine nucleotide biosynthesis is blocked, either through mutation of *ade2* or by the addition of mycophenolic acid, cells are unable to grow even with the addition of guanine to the media (1, 2). Null mutants are also resistant to the hypoxanthine/guanine analog 8-azaguanine as well as to the anti-cancer drug cisplatin (1, 7). In humans, partial and complete deficiencies of the HPT1 ortholog HPRT1 (OMIM) are associated with the genetic disorders HPRT-related gout/Kelley-Seegmiller syndrome (OMIM) and Lesch-Nyhan syndrome (OMIM), respectively (2 and references therein).

Detailed gene summaries

Q var

Show all results ...

VAR1 / Q0140

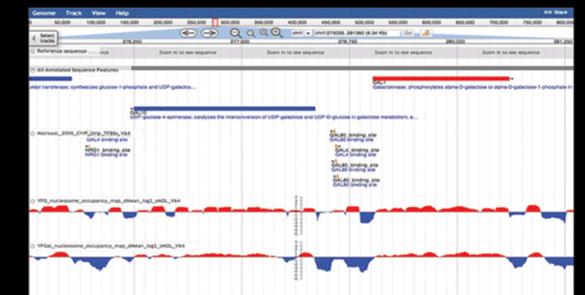
varicosity

antigenic variation

Variant Viewer

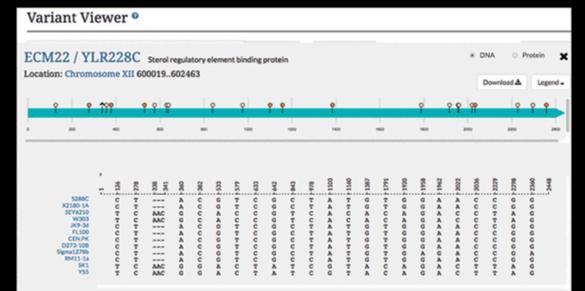
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yeastgenome.org/variant-viewer