

TSA2 (https://www.gephebase.org/search-criteria?/and+GeneGephebase=~TSA2~#gephebase-summary-title)	Gephebase Gene	GP00001831	GepheID
Published	Entry Status	Duveau	Main curator

	Trait Category
Physiology (https://www.gephebase.org/search-criteria?/and+Trait+Category=^Physiology^#gephebase-summary-title)	
	Trait
Xenobiotic resistance (hydrogen peroxide) (https://www.gephebase.org/search-criteria?/and+Trait=^Xenobiotic resistance (hydrogen peroxide)^#gephebase-summary-title)	
	Trait State in Taxon A
Sensitive	
	Trait State in Taxon B
Tolerant	
	Ancestral State
Taxon A	
	Taxonomic Status
Experimental Evolution (https://www.gephebase.org/search-criteria?/and+Taxonomic+Status=^Experimental Evolution^#gephebase-summary-title)	

GENOTYPIC CHANGE	
TSA2	Generic Gene Name UniProtKB <i>Saccharomyces cerevisiae</i> (strain ATCC 204508 / S288c)
YDR453C; D9461.38	Q04120 (http://www.uniprot.org/uniprot/Q04120) GenebankID or UniProtKB
4932.YDR453C	Synonyms <i>Saccharomyces cerevisiae</i> (strain ATCC 204508 / S288c)
(http://string-db.org/newstring.cgi/show_network_section.pl?identifier=4932.YDR453C)	String Q04120 (https://www.ncbi.nlm.nih.gov/nucore/Q04120)
Sequence Similarities	
Belongs to the peroxiredoxin family. AhpC/Prx1 subfamily.	
GO - Molecular Function	
GO:0051920 : peroxiredoxin activity (https://www.ebi.ac.uk/QuickGO/term/GO:0051920)	
GO:0008379 : thioredoxin peroxidase activity (https://www.ebi.ac.uk/QuickGO/term/GO:0008379)	
GO - Biological Process	
GO:0006457 : protein folding (https://www.ebi.ac.uk/QuickGO/term/GO:0006457)	
GO:0045454 : cell redox homeostasis	

(<https://www.ebi.ac.uk/QuickGO/term/GO:0045454>)
GO:0034599 : cellular response to oxidative stress
(<https://www.ebi.ac.uk/QuickGO/term/GO:0034599>)
GO:0006979 : response to oxidative stress
(<https://www.ebi.ac.uk/QuickGO/term/GO:0006979>)
GO:0042744 : hydrogen peroxide catabolic process
(<https://www.ebi.ac.uk/QuickGO/term/GO:0042744>)
GO:0050821 : protein stabilization (<https://www.ebi.ac.uk/QuickGO/term/GO:0050821>)
GO - Cellular Component
GO:0005737 : cytoplasm (<https://www.ebi.ac.uk/QuickGO/term/GO:0005737>)
GO:0005829 : cytosol (<https://www.ebi.ac.uk/QuickGO/term/GO:0005829>)

No	(https://www.gephebase.org/search-criteria?/and+Presumptive Null=^No^#gephebase-summary-title)	Presumptive Null
Gene Amplification	(https://www.gephebase.org/search-criteria?/and+Molecular Type=^Gene Amplification^#gephebase-summary-title)	Molecular Type
Complex Change	(https://www.gephebase.org/search-criteria?/and+Aberration Type=^Complex Change^#gephebase-summary-title)	Aberration Type
Chromosome 4 whole duplication. Using a genetic mapping strategy that involves systematically deleting segments of a duplicated chromosome; the authors show that the chromosome IV’s duplication effect is largely due to the generation of a second copy of the stress-inducible cytoplasmic thioredoxin peroxidase TSA2.		Molecular Details of the Mutation
Linkage Mapping	(https://www.gephebase.org/search-criteria?/and+Experimental Evidence=^Linkage Mapping^#gephebase-summary-title)	Experimental Evidence
The Stress-Inducible Peroxidase TSA2 Underlies a Conditionally Beneficial Chromosomal Duplication in Saccharomyces cerevisiae. (2017) (https://pubmed.ncbi.nlm.nih.gov/28743806)		Main Reference
Linder RA; Greco JP; Seidl F; Matsui T; Ehrenreich IM		Authors

Although chromosomal duplications are often deleterious, in some cases they enhance cells’ abilities to tolerate specific genetic or environmental challenges. Identifying the genes that confer these conditionally beneficial effects to particular chromosomal duplications can improve our understanding of the genetic and molecular mechanisms that enable certain aneuploidies to persist in cell populations and contribute to disease and evolution. Here, we perform a screen for spontaneous mutations that improve the tolerance of haploid Saccharomyces cerevisiae to hydrogen peroxide. Chromosome IV duplication is the most frequent mutation, as well as the only change in chromosomal copy number seen in the screen. Using a genetic mapping strategy that involves systematically deleting segments of a duplicated chromosome, we show that the chromosome IV’s duplication effect is largely due to the generation of a second copy of the stress-inducible cytoplasmic thioredoxin peroxidase TSA2 Our findings add to a growing body of literature that shows the conditionally beneficial effects of chromosomal duplication are typically mediated by a small number of genes that enhance tolerance to specific stresses when their copy numbers are increased.	Abstract
Copyright © 2017 Linder et al.	Additional References

RELATED GEPHE

15 (APJ1, ERG3, ERG5, ERG6, ERG7, LEU2, PHO84, RAD5, SWS2, CIS1, FRM2, GPX2, RTA1, cytochrome b, MKT1) (https://www.gephebase.org/search-criteria?/or+Taxon ID=^4932^/and+Trait=Xenobiotic resistance/and+groupHaplotypes=true#gephebase-summary-title)	Related Genes
No matches found.	Related Haplotypes

EXTERNAL LINKS

COMMENTS

@Fitness @GeneDuplication ; @& Experimental Evidence should be “Gene editing”.