

Variant: NM_000277.2(PAH):c.1242C>T (p.Tyr414=)

CA200893 [↗](#)

102577 (ClinVar) [↗](#)

Gene: PAH ([HGNC:5053](#))

Condition: phenylketonuria ([MONDO:0009861](#))

Inheritance Mode: Autosomal recessive inheritance

UUID: 349736d7-05a8-4ec0-9b30-04fa24dd5529

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HGVS expressions

NM_000277.2(PAH):c.1242C>T (p.Tyr414=)

NM_000277.2:c.1242C>T

NC_000012.12:g.102840473G>A

CM000674.2:g.102840473G>A

NC_000012.11:g.103234251G>A

CM000674.1:g.103234251G>A

NC_000012.10:g.101758381G>A

NG_008690.1:g.82130C>T

NG_008690.2:g.122938C>T

NM_000277.1:c.1242C>T

NM_001354304.1:c.1242C>T

NM_000277.3:c.1242C>T

ENST00000307000.7:c.1227C>T

ENST00000551114.2:n.904C>T

ENST00000553106.5:c.1242C>T

ENST00000635477.1:n.346C>T

ENST00000635528.1:n.757C>T

Benign

Met criteria codes **2**

BS2

BS1

Not Met criteria codes **1**

BP4

Evidence Links **0**

Expert Panel

Phenylketonuria VCEP [↗](#)

Criteria Specification Information **!**

[↗](#) Criteria Specifications for this VCEP

Evidence submitted by expert panel

Phenylketonuria VCEP

PAH-specific ACMG/AMP criteria applied: BS1: MAF=0.01361 in ENF from gnomAD; BS2: 19 homozygotes in gnomAD. In summary this variant meets criteria to be classified as benign for phenylketonuria in an autosomal recessive manner based on the ACMG/AMP criteria applied as specified by the PAH Expert Panel: (BS1, BS2).

Met criteria codes

BS2			19 homozygotes in gnomAD
BS1			MAF=0.01361 in ENF from gnomAD

Not Met criteria codes

BP4			Tolerated and benign in SIFT, Polyphen2. Disease causing in MutationTaster.
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Curation History



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