

## Overview

### Useful For

Diagnosis of primary coenzyme Q10 (CoQ10) deficiencies in some patients who are not supplemented with CoQ10

Monitoring patients receiving statin therapy

Monitoring CoQ10 status during treatment of various degenerative conditions, including Parkinson and Alzheimer diseases

Providing accurate quantitation of total CoQ10 when specimens are hemolyzed

This test is **not useful** for distinguishing primary CoQ10 deficiencies from acquired CoQ10 deficiencies.

### Genetics Test Information

This test is used for the diagnosis of coenzyme Q10 (CoQ10) deficiency in mitochondrial disorders. It is also used to monitor CoQ10 status in patients with mitochondrial cytopathies, patients receiving statin therapy, or during treatment of various degenerative conditions including Parkinson and Alzheimer diseases.

### Method Name

High-Performance Liquid Chromatography (HPLC) with Electrochemical Detection

### NY State Available

Yes

## Specimen

### Specimen Type

Plasma Heparin

### Ordering Guidance

This test does not quantitate reduced coenzyme Q10. For reduced coenzyme Q10 (CoQ10), order Q10 / Coenzyme Q10, Reduced and Total, Plasma.

The most reliable test for the diagnosis of primary defects in ubiquinone (ie, CoQ10) biosynthesis is direct measurement of CoQ10 in muscle.

### Necessary Information

**Patient's age is required.**

### Specimen Required

**Patient Preparation:** Fasting (8 hours)

**Collection Container/Tube:** Green top (lithium or sodium heparin)

**Submission Container/Tube:** Plastic vial

**Specimen Volume:** 0.5 mL

**Collection Instructions:**

1. Immediately after collection, place specimen on wet ice. Maintain on wet ice and process within 3 hours of collection.
2. Centrifuge, aliquot plasma into plastic vial, and freeze immediately.

**Forms**

[If not ordering electronically, complete, print, and send a Biochemical Genetics Test Request \(T798\)](#) with the specimen.

**Specimen Minimum Volume**

0.3 mL

**Reject Due To**

Gross hemolysis	OK
Gross lipemia	Reject
Gross icterus	OK

**Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
Plasma Heparin	Frozen (preferred)	14 days	
	Refrigerated	10 days	

**Clinical & Interpretive**

**Clinical Information**

Coenzyme Q10 (CoQ10) is an essential cofactor in the mitochondrial respiratory chain responsible for oxidative phosphorylation, where it functions as an electron carrier and acts as an antioxidant. It is found in all cell membranes and is carried by lipoproteins in the circulation. Approximately 60% of CoQ10 is associated with low-density lipoprotein (LDL), 25% with high-density lipoprotein, and 15% with other lipoproteins. CoQ10 is present in the body in both the reduced and oxidized forms, with the antioxidant activity of CoQ10 dependent on both its concentration and reduction-oxidation (redox) status.

CoQ10 deficiencies, which are clinically and genetically diverse, can occur due to defects in genes involved in the biosynthesis of ubiquinone (primary CoQ10 deficiency) or due to other causes, such as mitochondrial disorders (secondary CoQ10 deficiency).

Five major clinical phenotypes of CoQ10 deficiency have been described:

- Encephalomyopathy (elevated serum creatine kinase [CK], recurrent myoglobinuria, lactic acidosis)
- Cerebellar ataxia and atrophy (neuropathy, hypogonadism)
- Severe multisystemic infant form (nystagmus, optic atrophy, sensorineural hearing loss, dystonia, rapidly progressing)

nephropathy)

- Nephropathy, steroid resistant nephrotic syndrome leading to end stage kidney disease
- Isolated myopathy (exercise intolerance, fatigue, elevated serum CK)

Treatment with CoQ10 in patients with mitochondrial cytopathies can improve mitochondrial respiration in both brain and skeletal muscle.

CoQ10 has been implicated in other disease processes, including diabetes, neurodegenerative conditions such as Parkinson and Alzheimer diseases, as well as in aging and oxidative stress. CoQ10 may also play a role in hydroxymethylglutaryl-CoA reductase inhibitor (statin) therapy and may be relevant to statin-induced myalgia. Additionally, the redox status of CoQ10 may be a useful early marker for the detection of oxidative LDL modification.

### Reference Values

<18 years: 320-1558 mcg/L

> or =18 years: 433-1532 mcg/L

Miles MV, Horn PS, Tang PH, et al. Age-related changes in plasma coenzyme Q10 concentrations and redox state in apparently healthy children and adults. *Clin Chim Acta*. 2004;34:139-144

### Interpretation

Abnormal results are reported with a detailed interpretation including an overview of the results and their significance, a correlation to available clinical information provided with the specimen, differential diagnosis, and recommendations for additional testing when indicated and available.

### Cautions

No significant cautionary statements

### Clinical Reference

1. Salviati L, Trevisson E, Agosto C, Doimo M, Navas P. Primary coenzyme Q10 deficiency overview. In: Adam MP, Mirzaa GM, Pagon RA, et al. eds. *GeneReviews* [Internet]. University of Washington, Seattle; 2017. Updated June 8, 2023. Accessed November 1, 2023. Available at [www.ncbi.nlm.nih.gov/books/NBK410087/](http://www.ncbi.nlm.nih.gov/books/NBK410087/)
2. Desbats MA, Lunardi G, Doimo M, Trevisson E, Salviati L. Genetic bases and clinical manifestations of coenzyme Q10 (CoQ 10) deficiency. *J Inherit Metab Dis*. 2015;38(1):145-56. doi:10.1007/s10545-014-9749-9
3. Littarru GP, Tiano L. Clinical aspects of coenzyme Q10: An update. *Nutrition*. 2010;26:250-254
4. Hargreaves I, Heaton RA, Mantle D. Disorders of human coenzyme Q10 metabolism: An overview. *Int J Mol Sci*. 2020;21(18):6695. doi:10.3390/ijms21186695
5. Banach M, Serban C, Ursoniu S, et al. Statin therapy and plasma coenzyme Q10 concentrations-A systematic review and meta-analysis of placebo-controlled trials. *Pharmacol Res*. 2015;99:329-336. doi:10.1016/j.phrs.2015.07.008
6. Emmanuele V, Lopez LC, Berardo A, et al. Heterogeneity of coenzyme Q10 deficiency: patient study and literature review. *Arch Neurol*. 2012;69(8):978-983. doi:10.1001/archneurol.2012.206

### Performance

### Method Description

Coenzyme Q10 (CoQ10) is extracted from plasma with cold 1-propanol containing coenzyme Q9 as an internal standard. An aliquot of the lipid extract is fractionated by high-performance liquid chromatography (HPLC). (Tang PH, Miles MV, DeGrauw A, Hershey A, Pesce A. HPLC analysis of reduced and oxidized coenzyme Q[10] in human plasma. Clin Chem. 2001;47[2]:256-265; Claessens AJ, Yeung CK, Risler LJ, Phillips BR, Himmelfarb J, Shen DD. Rapid and sensitive analysis of reduced and oxidized coenzyme Q10 in human plasma by ultra performance liquid chromatography-tandem mass spectrometry and application to studies in healthy human subjects. Ann Clin Biochem. 2016;53[Pt 2]:265-273. doi:10.1177/0004563215593097)

**PDF Report**

No

**Day(s) Performed**

Monday through Friday

**Report Available**

3 to 5 days

**Specimen Retention Time**

1 month

**Performing Laboratory Location**

Rochester

**Fees & Codes****Fees**

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact [Customer Service](#).

**Test Classification**

This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. It has not been cleared or approved by the US Food and Drug Administration.

**CPT Code Information**

82542

**LOINC® Information**

Test ID	Test Order Name	Order LOINC® Value
TQ10	Coenzyme Q10, Total, P	27923-2

Result ID	Test Result Name	Result LOINC® Value
36764	CoQ10 Total	27923-2
36765	Interpretation (TQ10)	59462-2

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