



Serum coenzyme Q10 and risk of disabling dementia: The Circulatory Risk in Communities Study (CIRCS)



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ABSTRACT

Objective: To examine whether coenzyme Q10, a potent antioxidant, is associated with risk of dementia, which has not yet been elucidated. **Approach and results:** We performed a case–control study nested in a community-based cohort of approximately 6000 Japanese aged 40–69 years at baseline (1984–1994). Serum coenzyme Q10 was measured in 65 incident cases of disabling dementia with dementia-related behavioral disturbance or cognitive impairment incident between 1999 and 2004, and in 130 age-, sex- and baseline year-matched controls. Serum coenzyme Q10 was inversely associated with dementia: the multivariate odds ratios (95% confidence intervals) were 0.68 (0.26–1.78), 0.92 (0.33–2.56), and 0.23 (0.06–0.86) for individuals with the second, third, and highest quartiles of coenzyme Q10, respectively, as compared with the lowest quartile (P for trend = 0.05). A similar association was found for the coenzyme Q10/total cholesterol ratio: the respective ORs were 0.67 (0.25–1.78), 0.73 (0.28–1.92), and 0.21 (0.05–0.90) (P for trend = 0.04). **Conclusions:** Serum coenzyme Q10 levels were inversely associated with risk of disabling dementia.

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1. Introduction

Coenzyme Q10 (CoQ10), or ubiquinone, is a vitamin-like substance synthesized by animal cells. CoQ10 largely exists in the myocardium and plays a role in mitochondrial energy production. It also has an antioxidant function and is widely consumed as a supplement in the United States [1]. In addition to the prescription of CoQ10 as an orphan drug for mitochondrial encephalomyopathy, some evidence exists for a beneficial effect of CoQ10 on several neurologic diseases such as Parkinson disease [2], Huntington disease [3], and Friedreich ataxia [4,5] as well as on improved

physical exercise capacity [6] and lowered blood pressure [6,7]. Animal studies have shown a potential benefit of CoQ10 on cognitive function [8–11]. A randomized controlled trial, however, has shown that the supplementation of CoQ10 did not influence cerebrospinal fluid biomarkers in patients with mild-to-moderate Alzheimer disease [12]. Yet, evidence on this issue is still limited, and no prospective study has been performed on the preventive effect of CoQ10 on risk of incident dementia in the general population.

In the present study, we hypothesized that because of its antioxidant effect, serum level of CoQ10 is inversely associated with disabling dementia. To test this hypothesis, we conducted a nested case–control study in the Circulatory Risk in Communities Study (CIRCS), a large community-based cohort study of Japanese population.

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2. Materials and methods

The CIRCS is an ongoing dynamic community cohort study involving 5 communities in Japan. Details of the CIRCS protocol have been described elsewhere [13]. In this study, we included only 1 community, Kyowa, where disabling dementia surveillance is being carried out, and serum CoQ10 values have been measured. A total of 65 cases of patients who were diagnosed between 1999 and 2004 as having disabling dementia, participated in annual health checkups (baseline) at least 5 years before receiving the dementia diagnosis (ie, between 1984 and 1994), were aged 40–69 years at baseline, and provided sera for storage at baseline were identified. The criteria of disabling dementia were the same as those of our previous study [14], which also includes the validation of the criteria and the details of the study protocol. As supplemental analysis, we further classified the dementia cases into cases with and without history of stroke on the basis of the systematic stroke registration system described elsewhere [14]. We excluded individuals aged 70 years or older at baseline, because serum CoQ10 would be likely affected by the baseline age, and residents aged 70 years and over were not invited systematically to the baseline examination. One hundred thirty randomly selected controls whose age (± 3 years), sex, and follow-up time were matched at a ratio of 2:1 with the cases were also identified from the risk set. Venous blood was collected at baseline, and sera were prepared from the blood samples as soon as possible after the blood collection at the checkup sites. The serum samples were collected in 0.3 mL tubes and stored at -80°C until measurement in 2005. Serum CoQ10 was measured using high-performance liquid chromatography (HPLC) at the Public Health Institute of Kochi Prefecture with modification of the reported methods [15]. Briefly, a 10 μL serum sample was pretreated with 380 μL of 2-propanol. After centrifugation at 7000 g for 5 min, 10 μL of supernatant was applied to the HPLC system (L-7000 series; Hitachi High-Technologies Corporation, Tokyo, Japan) using a SUPELCOSIL LC-8 HPLC Column (Sigma–Aldrich Japan, Tokyo, Japan), an RC-10 reduction column (Shiseido Company, Tokyo, Japan), and an electrochemical detector (SI-2; Shiseido Company). The oxidation potential for the electrochemical detector was 600 mV. The mobile phase consisted of 50 mmol/L sodium perchlorate in methanol/2-propanol (100/10, v/v) with a flow rate of 1 mL/min. Serum total cholesterol was measured at baseline using the Liebermann-Burchard direct method at the Osaka Medical Center for Cancer and Cardiovascular Disease, an international member of the US National Cholesterol Reference Method Laboratory Network (CRMLN) [16].

For statistical analyses, we conducted conditional logistic analyses using SAS 9.1.3. Service Pack 4 (SAS Institute, Cary, NC, USA) with adjustments for body mass index, smoking status, alcohol consumption, diastolic blood pressure, total cholesterol, diabetes mellitus, and use of medication for hypertension or hypercholesterolemia. Since CoQ10 and total cholesterol were strongly correlated (Spearman $r = 0.34$), we also examined the association of the CoQ10/total cholesterol ratio with risk of disabling dementia. All probability values for the statistical tests were 2-tailed, and probability values below 0.05 were considered significant. Informed consent was obtained from community leaders and, verbally, from individual participants according to the guidelines of the Council for International Organizations of Medical Science [17]. The study was approved by the institutional review boards of the Osaka Center for Cancer and Cardiovascular Disease Prevention and of the University of Tsukuba.

3. Results

As shown in Table 1, the baseline characteristics did not differ materially between the cases and the non-cases, although diastolic

Table 1

Baseline characteristics of dementia cases and non-cases, CIRCS aged 40–69 years.

	Dementia cases	Non-cases	P for difference
Number	65	130	
Age, y	64.5	64.1	0.64
Male gender, %	34	34	1.00
Body mass index, kg/m ²	24.3	24.1	0.77
Current smokers, %	26	22	0.47
Current drinkers, %	28	28	1.00
Systolic blood pressure, mm Hg	137	136	0.43
Diastolic blood pressure, mm Hg	81	79	0.08
Antihypertensive medication, %	28	33	0.45
Diabetes mellitus, %	14	7	0.12
Serum total cholesterol, mg/dL	205	201	0.53
Cholesterol-lowering medication, %	3	10	0.09
Serum coenzyme Q10, ^a nmol/L	731	762	0.32
Coenzyme Q10/total cholesterol ratio, ^a 10 ⁻⁶	138.5	151.4	0.15

^a Median values and P values for differences tested by the Wilcoxon rank sum test.

blood pressure levels tended to be higher, and prevalence of cholesterol-lowering medication use, lower for cases than for non-cases.

Serum coenzyme Q10 was inversely associated with risk of incident dementia (Table 2). The multivariate odds ratios and 95% confidence intervals were 0.68 (0.26–1.78), 0.92 (0.33–2.56), and 0.23 (0.06–0.86), for individuals with the second, third, and highest quartiles of CoQ10, respectively, compared with the lowest quartile (P for trend = 0.05). A similar association was observed for the CoQ10/total cholesterol ratio. The respective odds ratios were 0.67 (0.25–1.78), 0.73 (0.28–1.92), and 0.21 (0.05–0.90) (P for trend = 0.04).

As supplemental analysis, we stratified dementia cases into those with history of stroke and those without it. Although the number of cases was limited, the associations were stronger for dementia without stroke for both serum CoQ10 and the CoQ10/total cholesterol ratio. The unadjusted odds ratios and 95% confidence intervals of dementia without stroke were 1.18 (0.37–3.75), 0.50 (0.15–1.70), and 0.46 (0.12–1.74) for individuals with the second, third, and highest quartiles of CoQ10 (P for trend = 0.13), and were 0.47 (0.15–1.48), 0.34 (0.11–1.09), and 0.24 (0.06–0.98) for respective quartiles of CoQ10/total cholesterol ratio (P for trend = 0.04). The corresponding odds ratios and 95% confidence intervals of dementia with stroke were 0.82 (0.26–2.57), 2.24 (0.63–7.96) and 0.35 (0.06–2.05) according to quartiles of CoQ10 (P for trend = 0.45), and were 1.68 (0.45–6.29), 2.05 (0.51–8.30) and 0.43 (0.06–2.97) according to quartiles of CoQ10/total cholesterol ratio (P for trend = 0.43).

4. Discussion

We found a strong inverse association between serum CoQ10 concentration and risk of disabling dementia in the Japanese population. To date, this is the first prospective study to examine the association between CoQ10 and incident dementia in a general setting.

Several animal studies have shown that CoQ10 may have a beneficial effect on dementia progression. In transgenic mice, dietary supplementation with CoQ10 reduced brain oxidative stress and deposition of amyloid plaque or amyloid- β and improved behavioral performance [11]. Rats with damaged hippocampi and cerebral cortices showed an adverse alteration in the markers of oxidative damage, but with supplementation with CoQ10, this alteration was reversed [8]. Thus, CoQ10 has been considered a

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