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Increased selenium intake in elderly high fish consumers may account for health benefits previously ascribed to omega-3 fatty acids

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Abstract

Objective: to examine relationships between fish consumption and plasma selenium (Se) and red blood-cell fatty acid (RBC FA) profile in aged subjects. We hypothesised that the importance of Se has been underestimated when interpreting the beneficial effect of fish consumption on health.

Design: cross-sectional analysis of data from a prospective cohort study

Setting: the EVA study in Nantes, France (1991-2002)

Subjects: 200 subjects aged ≥ 69 y with information on RBC FAs, plasma Se and completed food frequency questionnaires.

Methods: we examined correlations between the most abundant FAs, Se and number of fish meals per week. Linear regression models were used.

Results: Plasma Se was negatively correlated with RBC ω6 poly-unsaturated FA (PUFAs) and positively with ω3 PUFAs. Plasma Se, RBC ω3 PUFAs, docosahexaenoic acid (DHA) and eicosapentaenoic acid (EPA) increased with fish consumption. Conversely, levels of ω6 PUFAs were lower in the highest fish consumption group. All associations between plasma Se and fish consumption remained significant when adjusting for ω6 PUFAs alone or additionally for age, sex, education, diabetes, hypertension, dyslipidemia, cardiovascular diseases, and broad food categories (meat, eggs, dairy products, cereals, fruit and vegetable). Associations between ω3 PUFAs and fish also remained significant in the same model independently of Se. In linear regression models adjusted for demographic indicators, fish consumption explained only 2.6% of the variance in RBC ω3 FAs (6.2% for ω6) but as much as 15% of the variance in plasma selenium.

Conclusions: The observed health benefits of fish consumption in the elderly could be related not only to the increase in ω3 FA intake but also to other nutrients such as selenium. It is important to consider this observation when interpreting associations between fish consumption and health status in the elderly, particularly with regard to brain function.

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Introduction

The beneficial effect of fish consumption on elderly health and particularly on brain aging has been extensively studied (1). As fatty fish is the main dietary source of long-chain ω3 PUFA, this protective effect has been attributed to the long-chain ω3 PUFA content of fish (2, 3). In observational studies, high consumption of fish long-chain ω3 PUFA has been reported to reduce cognitive decline (4-6), risk of dementia (7-11), and to protect against stroke (12), cardiovascular diseases(13), metabolic syndrome (14), diabetes(15, 16) and cancer(17). However, these effects on health are controversial (18) as fish contains other nutrients than ω3 PUFA (19) that could be involved in the health benefits related to its consumption.

In the EVA (Epidemiology of Vascular Aging) study, we previously reported that high levels of red blood cell (RBC) ω6 PUFA and low levels of RBC ω3 PUFA were associated with greater risk of cognitive decline (20). We also reported that lower plasma selenium (Se) concentrations, were associated with higher risk of cognitive decline (21, 22). Fish is an important contributor to both long-chain ω3 PUFA and Se intake in older persons (23) as a number of commonly consumed fish species contain high concentrations of Se.(24) Long-chain ω3 PUFA (in particular eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA)) are synthesized from the essential FA alpha-linolenic acid, by elongating and desaturase enzymes. This enzymatic activity decreases with aging (25) and FA status is then, in ageing, more dependent on dietary supply of EPA and DHA.

We hypothesize that the role of Se has been underestimated when interpreting the beneficial effect of fish consumption on health. Thus, the present secondary analysis aimed to examine relationships between fish consumption, plasma Se concentration, and erythrocyte FA composition in older people.

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Methods

Study population

The aims, design and funding of the 9-year longitudinal EVA study have been previously described (22, 26). The study protocol was approved by the ethics committee of Hospital of Kremlin-Bicêtre, (Paris). Signed informed consent was obtained from all participants at enrolment in 1991-1993. Follow-up examinations were conducted every 18 months to 2 years until 2001-2002. In 1995-1997 (wave 3), due to financial limitations, only 342 blood samples, selected in chronological order, were analyzed for determination of erythrocyte FAs (20) (see figure 1). During the last follow-up of the EVA study in 2001-2002 (wave 6), volunteers answered the self-administered food frequency questionnaire (FFQ) and had blood sampled for plasma selenium determination. Out of the 342 selected subjects in 1995-1997, 200 had completed FFQs and blood sampling. These subjects did not differ from the initial subgroup for any of their demographic or health characteristics (data not shown).

As shown in figure 1, determinations of RBC-membrane FAs were performed more than 4 years before FFQ and plasma Se determination. We hypothesized that nutritional habits remained stable and we partially tested this hypothesis by examining the associations between baseline plasma Se determinations(1991-1993) and wave 6 Se determinations (2000-2001). The two determinations are highly correlated ($r=0.43$, $p<0.001$). Furthermore even 9 years before the FFQ administration, the association between plasma Se concentration and fish consumption (in quartiles) was clear. Mean baseline selenium value (sd) was 1.00 µmol/l (0.19) in quartile,1.07 (0.17) in quartile 2, 1.11 (0.17) in quartile 3 and 1.14 (0.17) in quartile 4 ($p=0.0007$).

Data collection

Self-administered food frequency questionnaire

A self-administered food frequency questionnaire assessed dietary habits, based on each meal (breakfast, lunch and dinner) and on consumption of each of the following foods: fish, eggs, milk and dairy products, cereals and starchy foods, fruit and vegetable, sugar and chocolate products.

Selenium and fatty acid determination

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Plasma Se (µmol/l) was determined as previously reported(27). FA composition of red blood cells (RBC) was determined as described previously (20).

General information

During face-to-face interviews conducted by trained psychologists, we gathered information on standard demographic variables and health status. Educational achievement was divided into two levels: no school or primary school (primary school) versus high school or university (high school or above). BMI was calculated. Subjects who reported a medical history of diabetes or who used anti-diabetic drugs or those who had a plasma glucose level ≥ 7.80 mmol/l (1.40g/l) were considered to be diabetic. Hyperlipidemia was defined as having LDL cholesterol ≥ 4.1 mmol/l (2.80 g/l) or using lipid-lowering drugs or reporting a hyperlipidemia medical history. Subjects with systolic blood pressure ≥ 140 mm Hg or diastolic blood pressure ≥ 90mm Hg, or using hypertensive drugs or those reporting a medical history of hypertension were considered to be hypertensive. History of vascular diseases was defined as self-reported history of myocardial infarction, angina pectoris and stroke or use of vascular drugs.

A global cognitive test, the Mini Mental Status Examination (MMSE) (28) and the Digit Symbol Substitution (DSS) from the Wechsler Adult Intelligence Scale-Revised (WAIS-R) that measures sustained attention and logical reasoning (29) were analysed in the current study. Cognitive assessment at wave 6 is available only on a sample of 143 subjects due mostly to refusal of cognitive evaluation.

Statistical analysis

From the food frequency questionnaire, we first examined the numbers of meals with fish consumption per week. In this population living on the Atlantic coast of France, the range was large [0-12] and we constructed dummy variables in 4 classes according to the quartile distribution: one meal with fish consumption or fewer per week (n=45), 2 meals per week (n=51), 3 meals per week (n=34) and 4 or more meals per week (n=70). Few subjects had a low consumption, only 3 never ate fish and only ten less than once a week.

The number of FA species identified was large. We restricted analyses to a limited number of chemical families and the most abundant FAs: saturated fatty

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acids (SFAs), mono-unsaturated fatty acids (MUFAs) and PUFAs (ω3 and ω6), DHA (22:6 ω3), EPA (20:5 ω3), arachidonic acid (AA) (20:4 ω6) and linoleic acid (18:2 ω6).

Qualitative and quantitative characteristics were compared between the four classes of fish consumers respectively by analysis of variance and chi-square test. We examined correlation between the selenium determinations and fatty acid membrane composition using Pearson's correlation coefficient.

In order to examine the respective link between fish consumption (dependant variable in 4 classes) and biological measurements (Se and FA), linear regression models were used (Proc REG from SAS software). The lowest intake level of fish (one or fewer fish meals per week) was the referent category. Covariates systematically included in the models were age, sex and education.

Moreover, we included as potential confounding variables factors that had previously been associated with fish consumption in a previous study (30), even though some were not statistically significantly associated with it in this sample: diabetes, hypertension, dyslipidemia and a history of cardiovascular disease. We further added the consumption of other broad food categories, each considered in two classes, with respect to median values in the population.

Statistical significance was defined by a p value <0.05. Statistical analyses were performed using SAS software version 9.1 (SAS Institute, Inc. Cary, North Carolina).

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Results

As indicated in table 1, 35% of subjects were heavy fish consumers whereas 22.5% ate fish less than once a week. Compared to moderate fish consumption (2 or 3 fish meals a week), low and high fish consumption is more frequently associated with hypertension. MMSE score did not differ between the 4 groups but there was an association between DSS performance and fish consumption, lower fish consumers having the poorest DSS performances. This association, described on a small sub-sample (n=143), did not remain significant when adjusting for age, sex and education.

Se concentration increased linearly with fish consumption from 0.92 µmol/l in the lowest consumption group to 1.10 µmol/l in the highest group. Similarly, increases were observed for total ω3 PUFAs (from 9.2% to 10.3 %), both for DHA and EPA. Conversely, levels of total ω6 PUFAs were lower in the highest fish consumption group (27.3%) compared to subjects with the lowest consumption per week (28.6%).

Se was negatively correlated with total ω6 PUFA ($r=-0.21$, $p\leq0.01$) and its major component AA ($r=0.15$, $p\leq0.05$). Conversely, correlation between plasma Se and total ω3 PUFA was positive ($r=0.22$, $p\leq0.01$). This positive association was found both for DHA ($r=0.22$, $p\leq0.01$) and for EPA ($r=0.21$, $p\leq0.01$). No correlation was observed between Se and SFAs ($r=0.05$, $P>0.05$) or MUFAs ($r=0.025$, $P>0.05$). Therefore, these two FAs groups were not considered in further analyses.

Fish consumption remained positively associated with plasma Se as well as with ω3 PUFAs and, negatively with ω6 PUFAs after adjusting for demographic indicators (table 2) The model explained 15% of the variance of selenium whereas the proportion of variance explained was much lower for PUFAs, 6.2% for ω6, 2.6% for ω3. These effects were mostly observed in the two highest classes of consumption. These associations were not modified when further adjusted for diabetes, hypertension, dyslipidemia, history of cardiovascular diseases and other food categories (data not shown).

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Discussion

In this secondary cross-sectional analysis of a sample of elderly subjects, fish consumption was independently positively associated with both plasma Se and RBC-membranes ω3 PUFAs. The significant linear increase of both plasma Se and RBC-membrane ω3 PUFA levels with increasing fish consumption suggests that the beneficial effect of fish consumption on health may not be exclusively attributed to long-chain ω3 PUFAs.

These independent associations remained significant even when controlling for a large number of demographic and health factors and broad food categories. These relationships are in agreement with previous reports (32-34). In a Chinese rural population, plasma Se and glutathione peroxidase activity were positively associated with RBC DHA. In that same population, ω3 PUFAs were negatively associated with coronary and hypertensive heart disease mortality whereas Se levels were negatively associated with chronic bronchitis, emphysema and respiratory heart diseases (32). In Greenland Inuits, Se intake has always been high and is closely linked to the consumption of traditional foods of marine origin. In addition, Se blood concentrations are highly correlated with long-chain marine FAs (34). In a Finnish population, the comparison of high fish consumers to low fish consumers (41 pairs, mean age 54.3 y) showed higher plasma ω3 PUFAs and lower ω6 in high fish consumers (33).

Long-chain n-3 PUFAs as found in some fish are susceptible to lipid peroxidation forming lipid hydroperoxides. Se, as the selenoenzymes GPx1 and GPx4, is required for the removal of these lipid hydroperoxides before they can be broken down to undesirable products (35). Thus a high oily fish intake probably requires a high Se intake to avoid deleterious health effects. Luckily, fish have a good Se content. The other role of Se in relation to fatty acid metabolism relates to peroxide tone which is lowered by the GPxs thereby down-regulating eicosanoid metabolism and associated physiological events(35).

Our results show that the highest fish consumers who also had the highest plasma selenium and RBC-membrane ω3 PUFA concentrations had the best cognitive performances in crude analyses. Therefore apparent protection against dementia in fish consumers, documented by previous epidemiological studies in

large cohorts (19, 31, 32) should not be exclusively attributed to long-chain ω-3 PUFAs but might involve a possible synergistic effect between selenium and long chain ω3 FA intakes.

The present study has however a number of limitations. First, The FFQ was completed retrospectively by the participants and we used a global indicator of fish consumption in this population of elderly subjects. We estimated the frequency of fish consumption but not the quantities, nor the type of fish consumed. The amount of PUFAs and Se vary largely according to fish species(24). PUFA content of fish is highly dependent on its fat content (fatty fish such as salmon, mackrel, herring, trout, tuna having a higher content than lean fish such as plaice or cod) and on whether it is farmed or wild. In addition, the cooking process (steamed, baked, or fried) may also considerably influence the final FA content of the dish. As regards Se, fish processing (cooking or enzyme and salt treatment) has not been reported to affect Se bioavailability (36).

Second, our sample size was limited by attrition during follow-up which limits the study power for comparing characteristics and health status of fish consumers. Multivariate results for cognition are not presented due to limited sample size and poor statistical power. Third, evaluation of the reported parameters was not performed simultaneously. Nevertheless, despite these limitations, the statistical link between fish consumption and biological indicators is powerful enough to allow us to hypothesize that the beneficial effect of fish consumption may not be limited to ω3 PUFAs alone but can also be attributed to Se. Besides Se, other micronutrients (for example iodine) may also be involved as already suggested (19, 31, 32). It is important to take our findings into account when discussing and interpreting associations between fish consumption and health status, particularly brain function, in the elderly.

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Table 1a: Demographic characteristics, health indicators and biological measurements according to fish consumption frequency (meals/ week)

	≤ 1 fish meal/w n=45	2 fish meals/w n=51	3 fish meals/w n=34	≥4 fish meals/w n=70	P value
Sex (% female)	53.3	58.8	67.6	54.3	0.55
Age (m ± sd)	73.3 ±3.0	73.8 ±3.1	72.6 ± 2.6	73.8 ± 3.3	0.24
High school education (%)	33.3	43.1	55.9	51.4	0.33
Diabetes (%)	4.4	7.8	2.9	10.0	0.50
Dyslipidémia (%)	33.3	31.4	52.9	40.0	0.20
Hypertension (%)	86.7	64.7	70.6	87.1	0.008
Cardiovascular diseases (%)	20.0	21.6	26.5	22.9	0.91
BMI (m ± sd) (n=162)	24,7 ± 3.6	24,8 ± 4.2	24,7 ± 3.0	25.1 ± 3.3	0.96
MMSE (m ± sd) (n=151)	28.6 ±1.2	28.4 ± 1.5	29.1 ± 0.8	28.5 ± 1.7	0.16
DSS score < 41 % (n=143)	43.7	37.8	19.2	18.8	0.04
Selenium (µmol/l)	0.92 ± 0.17	0.97 ± 0.16	1.01 ± 0.13	1.10 ± 0.19	<10-4
Total SFA (%)	43.6 ± 1.6	43.4 ± 1.2	44.3 ± 3.8	43.6 ± 1.6	0.31
Total MUFA %	16.8 ± 1.3	16.5 ± 1.1	16.9 ± 1.5	16.7 ± 1.1	0.33
Total ω6 PUFA (%)	28.6 ± 2.3	28.3 ± 1.9	26.8 ± 2.4	27.3 ± 2.5	0.0005
AA (20:4 ω6) (%)	13.8 ± 1.3	14.1± 1.2	13.3 ± 1.5	13.2 ± 1.5	0.0017
Linoleic Acid (18:2 ω6) (%)	9.42 ± 1.7	9.05 ± 1.5	8.82 ± 1.2	9.17 ± 1.6	0.36
Total ω3 PUFA (%)	9.20 ± 1.9	9.91 ± 1.4	9.95 ± 1.4	10.3 ± 2.0	0.01
DHA (22:6 ω3) (%)	5.78 ± 1.27	6.40 ± 1.05	6.33 ± 1.0	6.52 ± 1.1	0.008
EPA (20:5 ω3) (%)	0.98 ± 0.44	1.08 ± 0.33	1.24 ± 0.39	1.29 ± 0.72	0.014

BMI = Body Mass Index = weight / heigh^2; MMSE = Mini Mental Status Examination; DSS = Digit Symbol Substitution

FA expressed as percent of total fatty acids; SFA= Saturated Fatty Acids; MUFA= Monounsaturated Fatty Acids; PUFA= Polyunsaturated Fatty Acids; AA= Arachidonic Acid; DHA= Docosa Hexaenoic Acid; EPA= Eicosa Pentaenoic Acid

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Table 1b: Dietary habits associated to fish consumption frequency (% above the median of the sample)

	≤ 1 fish meal/w n=45	2 fish meals/w n=51	3 fish meals/w n=34	≥4 fish meals/w n=70	P value
Meat	44.4	52.9	50.0	57.1	0.61
Dairy products	46.7	54.9	50.0	50.0	0.88
Eggs	42.2	39.2	41.2	60.0	0.07
Cereals and starchy foods	33.0	33.0	50.	33.0	0.32
Fruits &Vegetables	51.1	51.0	38.2	55.7	0.42

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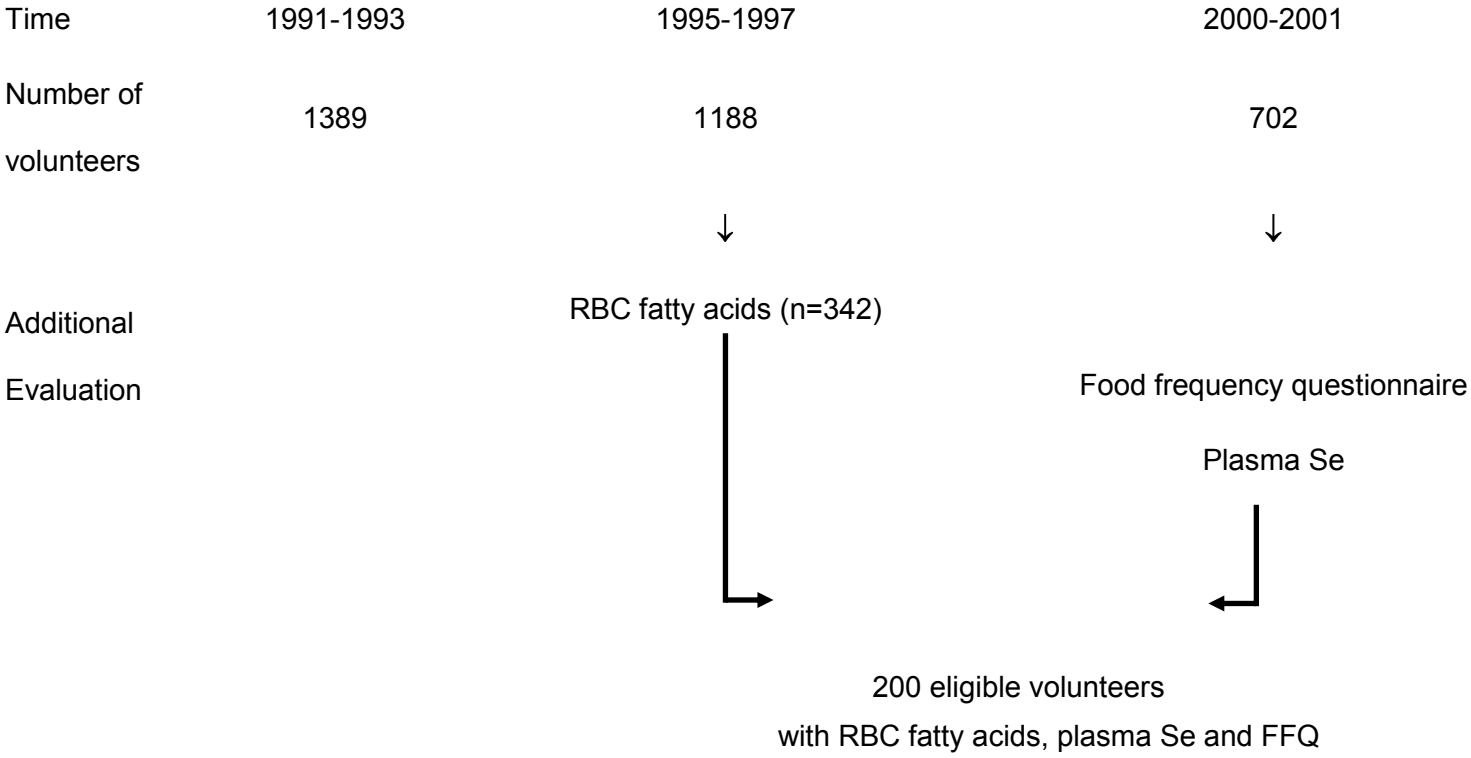
Table 2: Fish consumption (≤ 1, 2, 3, ≥ 4 meals per week): association with selenium and PUFA levels in an elderly population (n=200): results of linear regression models including systematically age, sex and education (Reference ≤ 1meal/week, n=45)

	2 meals/week n=51		3 meals/week n=34		≥4 meals/week n= 70		R^2
	β* ± sd	p	β* ± sd	p	β* ± sd	p	
Selenium	0.046 ± 0.035	0.19	0.076 ± 0.040	0.06	0.17 ± 0.03	<0.0001	0.149
Total PUFA ω6	-0.30 ± 0.47	0.52	-1.88 ± 0.53	0.0005	-1.34 ± 0.44	0.003	0.062
Total PUFA ω3	0.74 ± 0.37	0.04	0.77 ± 0.41	0.06	1.14 ± 0.34	0.001	0.026
DHA (22:6 ω3)	0.60 ± 0.23	0.01	0.51 ± 0.26	0.06	0.73 ± 0.21	0.001	0.040
EPA(20:5 ω3)	0.10 ± 0.11	0.37	0.28 ± 0.12	0.03	0.31 ± 0.10	0.003	0.027

* β regression coefficient is expressed for 1 unit

Figure 1 : Selection of the studied population



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