

Review Article

Is vitamin C supplementation beneficial? Lessons learned from randomised controlled trials

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In contrast to the promised ‘antioxidant miracle’ of the 1980s, several randomised controlled trials have shown no effect of antioxidant supplements on hard endpoints such as morbidity and mortality. The former over-optimistic attitude has clearly called for a more realistic assessment of the benefit:harm ratio of antioxidant supplements. We have examined the literature on vitamin C intervention with the intention of drawing a conclusion on its possible beneficial or deleterious effect on health and the result is discouraging. One of several important issues is that vitamin C uptake is tightly controlled, resulting in a wide-ranging bioavailability depending on the current vitamin C status. Lack of proper selection criteria dominates the currently available literature. Thus, while supplementation with vitamin C is likely to be without effect for the majority of the Western population due to saturation through their normal diet, there could be a large subpopulation with a potential health problem that remains uninvestigated. The present review discusses the relevance of the available literature on vitamin C supplementation and proposes guidelines for future randomised intervention trials.

Vitamin C: Supplementation: Randomised controlled trials

Vitamin C plays a role in numerous biological reactions, many of which are only known in little detail. Over the years, it has been suggested that vitamin C be used as a remedy against many diseases as different as common colds and cancers. Even today, there is considerable controversy about the exact role of the vitamin in human health and no agreement has been reached on the amount needed to be consumed for optimum wellbeing. Thus, as little as 10 mg/d will largely prevent development of the most well-known clinical and ultimately mortal manifestation of severe vitamin C deficiency: scurvy⁽¹⁾. Nevertheless, the RDA for vitamin C was recently increased from 60 mg/d to 75 mg/d for women and 90 mg/d for men in the US, primarily based on biochemical evidence⁽²⁾. Others have argued that the optimum plasma concentration is about the level of saturation (70 µmol/l), which would require a daily intake of about 200 mg^(2–4), and still hypotheses on new specific roles of vitamin C in health and disease are being put forward^(5–7).

The potential benefit of vitamin C supplementation has been fueled in part by a considerable body of epidemiological literature suggesting a positive association between vitamin C status and health. Thus, several large cohort studies have shown an inverse relationship between plasma vitamin C

status and risk of CVD and/or all-cause mortality^(8–15). In contrast, large randomised controlled trials using antioxidant supplements have been less promising. None of the major clinical studies using mortality or morbidity as endpoints has found significant positive effects of supplementation with vitamin C^(16–19). However, the vast majority of these trials have examined the effect a multi-component supplement and consequently not the effect of vitamin C itself.

The results from clinical trials in the last decades have shifted public opinion and that of health authorities towards antioxidants, including vitamin C, being generally unimportant. This development is likely to obscure a public health risk from deficiency, as several large cross-sectional population studies have shown that a considerable proportion (up to 50%) of subpopulations of the Western world can have hypovitaminosis C, defined as a plasma concentration less than 23 µmol/l^(20,21). While the clinical significance of this condition remains to be clarified – beyond the increased risk of developing scurvy – it is obvious that large subpopulations, for example, smokers, do not achieve the RDA of vitamin C⁽²²⁾.

It has been shown that those individuals most likely to benefit from supplements also are those least likely to

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get them^(23–25). So far, this discouraging finding is unfortunately also valid for most of the randomised controlled trials using vitamin C in their intervention. One frequently overlooked problem is that vitamin C uptake is highly dose dependent^(3,26). Thus, subjects already saturated with vitamin C through their daily diet will efficiently excrete any surplus and are therefore highly unlikely to benefit from further vitamin C supplementation. This and several other issues should be taken into account when designing and drawing conclusions from randomised controlled trials with the purpose of studying the effects of vitamin C. In view of the pharmacology and kinetics of vitamin C, the present review examines the current knowledge of the effect of vitamin C supplementation, evaluates the lessons to be learned from the many trials that have been conducted, and provides guidelines for future randomised trials.

Clinical significance and prevalence of vitamin C deficiency in observational studies

The definition of optimal vitamin C status remains a matter of controversy. However, current opinions appear to agree on a dose that gives saturated uptake, i.e. a dietary intake resulting in a plasma concentration of approximately 70 $\mu\text{mol/l}$ ^(3,27–30). Defining vitamin C deficiency is also complex since considerable individual variation apparently exists in the relationship between the plasma concentration of vitamin C and the development of scurvy, the classic hallmark of severe vitamin C deficiency^(31,32). Moreover, the clinical significance of vitamin C deficiency – beyond that of scurvy – has not been clearly defined. Guidelines developed by the National Survey of Canada suggested categories of severe vitamin C deficiency (serum level < 11 $\mu\text{mol/l}$) and marginal vitamin C deficiency (serum levels between 11 and 23 $\mu\text{mol/l}$) and have largely been adopted⁽²¹⁾. Since these categories were put forward in 1987, the RDA for vitamin C has been increased in an attempt to reflect the now-believed optimal vitamin C level in plasma of 70 $\mu\text{mol/l}$. Therefore a new category (for serum levels between 23 and, for example, 50 $\mu\text{mol/l}$) is needed, and we suggest it to be termed suboptimal vitamin C status.

Severe vitamin C deficiency

Scurvy typically constitutes the ultimate clinical manifestation of prolonged and severe vitamin C deficiency. In non-smokers, scurvy is prevented by a daily intake of as little as 10 mg of vitamin C⁽¹⁾. Clinical symptoms include follicular hyperkeratosis, petechiae, ecchymoses, coiled hairs, inflamed and bleeding gums, perifollicular haemorrhages, joint effusions, arthralgia and impaired wound healing⁽³³⁾. Other early symptoms include dyspnoea, weakness, fatigue and depression. Cases of scurvy are usually limited to the group of individuals with plasma concentrations lower than 11 $\mu\text{mol/l}$, i.e. those diagnosed with severe vitamin C deficiency. However, far from all individuals with plasma levels < 11 $\mu\text{mol/l}$ develop clinical scurvy^(31,32). Thus, other factors seem to be of importance and the relationship between plasma vitamin C status and scurvy is not entirely clear, when the diet is not totally depleted from the vitamin. However,

older reports indicate that total deficiency over a prolonged time invariably leads to scurvy⁽²⁾.

While the basic symptoms and cure of the disease have been known for centuries⁽³⁴⁾, a significant part of the population in developed countries continues to suffer from severe vitamin C deficiency and thus have increased risk of experiencing scurvy-like symptoms (Table 1). But the clinical significance of severe vitamin C deficiency may extend beyond that of scurvy. In clinical studies in which subjects were made vitamin C deficient, common complaints such as gingival inflammation, fatigue and depression were among the most sensitive markers of deficiency^(3,35). In a prospective population study, Nyssönen *et al.* found a higher risk of myocardial infarction (relative risk 3.5) among men with severe vitamin C deficiency, constituting about 6% of their Finnish cohort (1605 subjects)⁽¹²⁾. Moreover, Langlois *et al.* recently showed that 14% of patients with peripheral arterial disease suffered from severe vitamin C deficiency compared with none of the healthy controls and suggested a relationship between vitamin C status and severity of atherosclerosis⁽³⁶⁾. In a study with advanced cancer patients, 30% had severe vitamin C deficiency and these patients had shorter survival⁽³⁷⁾.

Marginal vitamin C deficiency

As defined above, a plasma concentration between 11 and 23 $\mu\text{mol/l}$ is termed marginal vitamin C deficiency. Hypovitaminosis C has been characterised as having a plasma concentration of vitamin C < 23 $\mu\text{mol/l}$ ⁽³⁸⁾, i.e. encompassing both severe and marginal vitamin C deficiency. As with severe vitamin C deficiency, smokers also have a markedly increased risk of marginal vitamin C deficiency (Table 1).

The clinical significance of marginal vitamin C deficiency – as different from severe vitamin C deficiency – has not been thoroughly investigated. In most studies, upper and lower tertiles, quartiles or quintiles are compared, making it difficult to compare groups between studies. Consequently, the category of marginal vitamin C deficiency can rarely be singled out from vitamin C deficiency or hypovitaminosis C. With respect to scurvy, clinical cases among individuals with marginal vitamin C deficiency are rare, but do occur^(39,40). Probably more important though, considerable epidemiological evidence suggests that there may be other clinical consequences of marginal vitamin C deficiency. Thus, in a recent re-examination of the Second National Health and Nutrition Examination Survey (NHANES II) data combined with a follow up on vital status 12–16 years later, Loria *et al.* found that men in the lowest (<28.4 $\mu\text{mol/l}$) compared with the highest (>73.8 $\mu\text{mol/l}$) serum ascorbate quartile had a 57% higher risk of death from any cause and a 62% higher risk of dying from cancer⁽¹¹⁾. A similar conclusion was reached by Simon *et al.* who also found that severe or marginal vitamin C deficiency was significantly associated with all-cause mortality while being weakly associated with death from CVD⁽⁴¹⁾. In a 20-year follow-up study in Britain (730 subjects), a significantly higher risk of mortality from stroke was observed in elderly men and women with severe and marginal vitamin C deficiency separately compared with those with plasma concentrations of vitamin C > 28 $\mu\text{mol/l}$ ⁽⁹⁾. The authors concluded that vitamin C status was as strong a predictor of death from

Table 1. Prevalence of vitamin C deficiency in larger cross sectional population studies

Study	Subjects (n)	Severe vitamin C deficiency (< 11 µmol/l)	Marginal vitamin C deficiency (11–23 µmol/l)	Suboptimal vitamin C status (23–50 µmol/l)	Comment
NHANES III ⁽⁵⁰⁾	15 769	14 % (M) 10 % (F)	20 % (M) 17 % (F)	NR	31 % (M) and 25 % (F) of the smokers alone were diagnosed with severe vitamin C deficiency
NHANES II ⁽⁵¹⁾	11 592	2 %	8 %	NR	7 and 20 %, respectively, in smokers alone
NHANES II ⁽⁴¹⁾	8453	NR	9 % (NS) 30 % (S)	31 % (NS)* 35 % (S)	Subpopulation aged 30 years or older
CARDIA ⁽⁵²⁾	2637	NR	8 % (NS) 26 % (S)	33 % (NS)† 40 % (S)	Numbers with marginal vitamin C deficiency include those with severe vitamin C deficiency
Third Glasgow MONICA population survey ⁽⁵³⁾	1267	26 % (M) 14 % (F)	22 % (NSM) 16 % (NSF) 30 % (SM) 30 % (SF)	NR	36 % (M) and 23 % (F) of the smokers alone were diagnosed with severe vitamin C deficiency
French population study ⁽⁵⁴⁾	1039	7–12 % (M) 3–5 % (F)	10–46 % (M)‡ 3–15 % (F)	NR	Values are ranges of various age groups

NHANES III, Third National Health and Nutrition Examination Survey; M, males; F, females; NR, not reported; NHANES II, Second National Health and Nutrition Examination Survey; NS, non-smokers; S, smokers; CARDIA, Coronary Artery Risk Development in Young Adults Study; MONICA, Monitoring of Trends and Determinants in Cardiovascular Disease; NSM, non-smoking males; NSF, non-smoking females; SM, smoking males; SF, smoking females.

* Range used: 23 to 55 µmol/l.

† Range used: 23 to 45 µmol/l.

‡ Range used: 11 to 19 µmol/l.

stroke as diastolic blood pressure⁽⁹⁾. An inverse correlation between vitamin C status and stroke was also reported from a study (2121 subjects) in a rural Japanese population aged 40 years or older⁽⁴²⁾. In the 12-year follow up on the Basel Prospective Study, a significantly increased risk of IHD and stroke was found in individuals with plasma ascorbate < 22.7 µmol/l, corresponding to severe or marginal vitamin C deficiency^(43–45).

Suboptimal vitamin C status

Based on the increased RDA for vitamin C as well as the indication that a plasma concentration of vitamin C of about 70 µmol/l is currently considered optimal for health, we suggest a new category of suboptimal vitamin C status for those individuals with plasma concentrations between 23 and 50 µmol/l. An obvious rationale for this additional category could be that if 70 µmol/l is optimal, for example, 35 µmol/l is probably not, and therefore investigations into the clinical significance of a suboptimal vitamin C status are warranted. Moreover, a proper control group should be selected from individuals with optimal vitamin C status, i.e. excluding those with suboptimal status. However, limited data are available and need to be extracted from studies discriminating between the concentrations of suboptimal and optimal vitamin C status (Table 1).

As discussed above, several large prospective studies have shown an inverse relationship between plasma vitamin C status and risk of CVD and/or all-cause mortality^(8–15). However, no studies have investigated the specific clinical significance of suboptimal vitamin C status as compared with optimal. Thus, it remains to be established if the biochemical evidence pointing towards an optimal plasma level of about 70 µmol/l can be backed up in larger epidemiological studies or ultimately in clinical trials. Clearly, the effects of suboptimal compared with optimal vitamin C

status are likely to be at most moderate and presumably relevant only in the long term if at all. Thus, it is debatable if studies aimed at clarifying such a limited risk are feasible from a cost perspective. On the other hand, the problems potentially associated with suboptimal vitamin C status affect a large percentage of the population and can be readily and inexpensively cured⁽²²⁾.

Randomised controlled trials with vitamin C

Randomised clinical trials have evolved and been refined for testing drug effects. Their strength lies in eliminating or reducing bias by randomisation, blinding and control. In the case of drugs, this design is superior and regarded as the 'gold standard'. The design is particularly strong for testing the effects of a chemical that is normally not present in the organism, has a relatively short pharmacokinetic and pharmacodynamic half-life, and is used in a relatively short dose regimen. In the case of intervention with dietary components in prevention trials, a number of problems arise that are not prominent in drug testing.

In the present context, we will particularly discuss the testing of proper biological hypotheses in relevant cohorts.

In the 1980s and 1990s, the epidemiological evidence pointed towards the importance of antioxidant intake (vitamin C, vitamin E and β-carotene) in the prevention of, for example, cancer and arteriosclerosis. This led to the initiation of a large number of clinical intervention studies. The first large study published was the Linxian study that showed an inspiring preventive effect⁽¹⁶⁾. The subsequent studies were all negative. At that time, the prevailing hypothesis was that dietary components were beneficial, without side effects, and the larger dose the better. Implicitly, it was also believed that cancer and arteriosclerosis were the result of 'high-level deficiency' of these substances. As a consequence, trials were mainly designed for the broad

Table 2. Randomised, controlled trials with vitamin C

Reference	Subjects (n)	Design	Risk of bias	Supplementation period (years)	Follow up (years)	Mean age (years)	Sex (% women)	Intervention used				
								Vitamin C (mg)	β-Carotene (mg)	Vitamin A (IU)*	Vitamin E (IU)†	Se (μg)
McKeown-Eyssen <i>et al.</i> (1988) ⁽⁵⁵⁾	185	Parallel	H	2	2	58	32	400			400	
Penn <i>et al.</i> (1991) ⁽⁵⁶⁾	30	Parallel	H	0.077	0.077	84	80	100		8000	50	
Chandra (1992) ⁽⁵⁷⁾	96	Parallel	H	1	1	74	55	80	16	1333	44	20
Blot <i>et al.</i> (1993) ⁽¹⁶⁾	29584	1/2(2 × 2 × 2 × 2)	H	5.25	5.25	NA	55	120	15	5000	33	50
Wenzel <i>et al.</i> (1993) ⁽⁵⁸⁾	56	Parallel	H	0.082	0.082	48	20	180	12		894	200
ter Riet <i>et al.</i> (1995) ⁽⁵⁹⁾	88	2 × 2	H	0.23	0.23	NA	NA	1000				
Hogarth <i>et al.</i> (1996) ⁽⁶⁰⁾	106	2 × 2	H	0.083	0.083	83	56	500		8000		
Girodon <i>et al.</i> (1997) ⁽⁶¹⁾	81	2 × 2	H	2	2	84	75	120	6		15	100
You <i>et al.</i> (2001) ⁽⁶²⁾	3411	2 × 2 × 2	H	3.25	3.25	NA	49	500	15		200	75
Sasazuki <i>et al.</i> (2003) ⁽⁶³⁾	439	2 × 2	H	5	5	57	65	500	15			
Bonelli <i>et al.</i> (1998) ⁽⁶⁴⁾	304	Parallel	H	5	5	NA	NA	180		6000	30	200
Li <i>et al.</i> (1993) ⁽⁶⁵⁾	3318	Parallel	L	6	6	54	56	180	15	10 000	60	50
Greenberg <i>et al.</i> (1994) ⁽⁶⁶⁾	864	2 × 2	L	4	4	61	21	1000	25		440	
Pike & Chandra (1995) ⁽⁶⁷⁾	47	Parallel	L	1	1	69	72	90		2666	45	
Richer (1996) ⁽⁶⁸⁾	71	Parallel	L	1.5	1.5	72	7	750		20 000	200	50
Girodon <i>et al.</i> (1999) ⁽⁶⁹⁾	725	2 × 2	L	2	2	84	74	120	6		16.5	100
Correa <i>et al.</i> (2000) ⁽⁷⁰⁾	976	2 × 2 × 2	L	6	6	51	54	2000	30			
Jacobson <i>et al.</i> (2000) ⁽⁷¹⁾	112	Parallel	L	0.5	0.5	42	42	500	12		400	
Age Related Eye Disease Study Research Group (2001) ⁽⁷²⁾	4757	2 × 2	L	6.3	6.3	68	56	500	15		400	
Brown <i>et al.</i> (2001) ⁽⁷³⁾	160	2 × 2	L	3	3	53	13	1000	25		800	100
Chylack <i>et al.</i> (2002) ⁽⁷⁴⁾	297	Parallel	L	3	3	68	59	750	18		660	
Graat <i>et al.</i> (2002) ⁽⁷⁵⁾	652	2 × 2	L	1	1	NA	50	60	1.2	2000	272	25
Heart Protection Study Collaborative Group (2002) ⁽¹⁸⁾	20536	2 × 2	L	5	5	NA	25	250	20		660	
Waters <i>et al.</i> (2002) ⁽⁷⁶⁾	423	2 × 2	L	3	3	65	100	1000			800	
White <i>et al.</i> (2002) ⁽⁷⁷⁾	100	Parallel	L	0.23	0.23	63	42	1000			223	
Prince <i>et al.</i> (2003) ⁽⁷⁸⁾	61	Cross-over	L	0.25	0.25	58	92	150	3		74.5	75
Salonen <i>et al.</i> (2003) ⁽⁷⁹⁾	520	2 × 2	L	6	6	NA	51	250			272	
Allsup <i>et al.</i> (2004) ⁽⁸⁰⁾	164	Parallel	L	0.15	0.5	83	63	120		2666	60	60
Hercberg <i>et al.</i> (2004) ⁽¹⁹⁾	13017	Parallel	L	7.54	7.54	49	61	120	6		33	100
Richer <i>et al.</i> (2004) ⁽⁸¹⁾	61	Parallel	L	1	1	75	4	1500	10	2500	500	200
Avenell <i>et al.</i> (2005) ⁽⁸²⁾	910	Parallel	L	1	1	72	47	60		2666	10	
Mooney <i>et al.</i> (2005) ⁽⁸³⁾	284	Parallel	L	1.25	1.25	37	45	500			400	
Tam <i>et al.</i> (2005) ⁽⁸⁴⁾	39	Parallel	L	0.23	2.67	46	100	500			800	
Witte <i>et al.</i> (2005) ⁽⁸⁵⁾	32	Parallel	L	0.75	0.75	NA	NA	500		2666	400	50
Cook <i>et al.</i> (2007) ⁽¹⁷⁾	8171	2 × 2 × 2	L	9.4	9.4	61	100	500	50/2nd d		600/2nd d	

H, high-bias design; NA, not available; L, low-bias design.

* 1 IU vitamin A = 0.3 μg.

† 1 IU vitamin E = 0.667 mg.

Table 3. Compliance of randomised, controlled trials with the present set of guidelines*

Reference	Low- or high-bias design†	Hypovitaminosis C at entry	Entry-level inclusion criterion	Plasma concentration measured at baseline and during study	Single supplement	Dietary intake controlled during study	Smoking status reported	Valid hypothesis or molecular mechanism proposed	Mechanism-related hard clinical endpoint	Inclusion and exclusion criteria reported
McKeown-Eyssen <i>et al.</i> (1988) ⁽⁵⁵⁾	H	–	–	(+)	–	–	+	+	+	–
Penn <i>et al.</i> (1991) ⁽⁵⁶⁾	H	–	–	+	–	–	–	+	–	+
Chandra (1992) ⁽⁵⁷⁾	H	–	–	–	–	–	–	+	–	(+)
Blot <i>et al.</i> (1993) ⁽¹⁶⁾	H	+	–	+	–	–	+	+	+	+
Wenzel <i>et al.</i> (1993) ⁽⁵⁸⁾	H	–	–	–	–	–	–	–	+	+
ter Riet <i>et al.</i> (1995) ⁽⁵⁹⁾	H	–	–	+	+	–	–	–	–	+
Hogarth <i>et al.</i> (1996) ⁽⁶⁰⁾	H	–	–	–	–	–	–	–	–	(+)
Girodon <i>et al.</i> (1997) ⁽⁶¹⁾	H	–	–	+	–	+	–	–	–	+
You <i>et al.</i> (2001) ⁽⁶²⁾	H	–	–	+	–	–	–	–	–	+
Sasazuki <i>et al.</i> (2003) ⁽⁶³⁾	H	–	–	+	+	–	+	+	–	+
Bonelli <i>et al.</i> (1998) ⁽⁶⁴⁾	H	–	–	–	–	–	–	+	+	+
Li <i>et al.</i> (1993) ⁽⁶⁵⁾	L	–	–	+	–	–	+	+	+	+
Greenberg <i>et al.</i> (1994) ⁽⁶⁶⁾	L	–	–	–	–	–	–	+	–	+
Pike & Chandra (1995) ⁽⁶⁷⁾	L	–	–	–	–	–	–	+	–	+
Richer (1996) ⁽⁶⁸⁾	L	–	–	–	–	+	–	+	+	(+)
Girodon <i>et al.</i> (1999) ⁽⁶⁹⁾	L	–	–	+	–	–	–	–	–	+
Correa <i>et al.</i> (2000) ⁽⁷⁰⁾	L	–	–	–	+	–	–	+	+	+
Jacobson <i>et al.</i> (2000) ⁽⁷¹⁾	L	–	–	+	–	–	+	+	–	+
Age Related Eye Disease Study Research Group (2001) ⁽⁷²⁾	L	–	–	+	–	–	–	–	–	+
Brown <i>et al.</i> (2001) ⁽⁷³⁾	L	–	–	+	–	+	+	–	–	+
Chylack <i>et al.</i> (2002) ⁽⁷⁴⁾	L	–	–	+	–	–	+	–	–	+
Graat <i>et al.</i> (2002) ⁽⁷⁵⁾	L	–	–	+	–	–	+	–	–	+
Heart Protection Study Collaborative Group (2002) ⁽¹⁸⁾	L	–	–	–	–	–	+	–	–	+
Waters <i>et al.</i> (2002) ⁽⁷⁶⁾	L	–	–	–	–	–	–	–	+	+
White <i>et al.</i> (2002) ⁽⁷⁷⁾	L	–	–	–	–	–	–	+	–	+
Prince <i>et al.</i> (2003) ⁽⁷⁸⁾	L	–	–	–	–	–	–	–	–	+
Salonen <i>et al.</i> (2003) ⁽⁷⁹⁾	L	–	–	+	+	–	+	+	–	+
Allsup <i>et al.</i> (2004) ⁽⁸⁰⁾	L	–	–	+	–	–	–	+	–	+
Herberg <i>et al.</i> (2004) ⁽¹⁹⁾	L	–	–	+	–	–	–	–	+	+
Richer <i>et al.</i> (2004) ⁽⁸¹⁾	L	–	–	–	–	–	+	–	–	(+)
Avenell <i>et al.</i> (2005) ⁽⁸²⁾	L	–	–	–	–	–	–	–	–	+
Mooney <i>et al.</i> (2005) ⁽⁸³⁾	L	–	–	+	–	–	+	+	–	+
Tam <i>et al.</i> (2005) ⁽⁸⁴⁾	L	–	–	+	–	–	+	+	–	+
Witte <i>et al.</i> (2005) ⁽⁸⁵⁾	L	–	–	–	–	–	–	–	–	+
Cook <i>et al.</i> (2007) ⁽¹⁷⁾	L	–	–	+	(+)	–	+	+	+	+

L, low-bias design; H, high-bias design.

* + indicates full compliance, (+) indicates partial compliance and – indicates that the criterion was not met by the study.

† As defined by Bjelakovic *et al.* (48).

Is vitamin C supplementation beneficial?

and little-selected population and doses were very high. Today, basic knowledge of the biological effects of the antioxidants has increased, and more importantly, their functions are no longer considered to be generally antioxidative, but rather as specific cofactors in biological reactions or direct signalling, signalling modifying, or gene-expression modifying compounds⁽⁴⁶⁾.

Epidemiological evidence is sometimes at great variance with the evidence from randomised controlled trials, particularly if control is not extensive⁽⁴⁷⁾. It should be acknowledged that in the epidemiological studies on the relationship between vitamin C concentrations and diseases, there is no evidence that the relationship is due to vitamin C itself. Thus, it is possible that vitamin C concentrations in plasma are a proxy or surrogate for vegetable and/or fruit intake and it may be some other substance in these foods that provides the health benefit. It might even be that the individuals with a high vegetable and fruit intake have no or a reduced intake of other foods with deleterious health effects, in which case vitamin C is a marker of an absence of a negative factor.

Neither epidemiological studies nor randomised clinical intervention trials can test mechanisms, but randomised controlled intervention trials can confirm if the effect is due to a single substance.

Current knowledge based on randomised controlled trials and recommendations for future studies

A large number of randomised clinical studies on antioxidants are now available. They have recently been reviewed and tabulated for effect by Bjelakovic *et al.*⁽⁴⁸⁾. That review, however, was done with the purpose of estimating risk of mortality for any antioxidant treatment, alone or in combination. The authors categorised the studies as high or low risk of bias. Thus, trials with adequate generation of the allocation sequence, adequate allocation concealment, adequate blinding and adequate follow-up were considered low-bias risk trials (high methodological quality), while trials with one or more unclear or inadequate quality components were classified as high-bias risk trials (low methodological quality)⁽⁴⁹⁾.

We searched the literature by using identical criteria to those above⁽⁴⁸⁾ and reviewed the combined number of papers using vitamin C as an intervention (Table 2). We then added a new set of criteria specifically addressing vitamin C (Table 3). Thus, based on the well-established dose dependency of vitamin C pharmacokinetics, we believe that it is imperative that enrolled subjects have hypovitaminosis C at study entry and that this condition is used as an entry-level inclusion criterion in order to ensure a possibility of effect. To verify the vitamin C status at entry and during the study, plasma concentration needs to be measured before and during the study by a validated method. As discussed above, vitamin C needs to be used as a single supplement to be able to determine the effect of this supplement specifically. Major confounders are, for example, dietary vitamin C and smoking status, and these factors need to be taken into account in the study design. A valid hypothesis or molecular mechanism should be proposed involving vitamin C and a mechanism-related hard clinical endpoint used as the

primary outcome. Finally, inclusion and exclusion criteria should be reported.

Reviewing the extracted literature, it is striking that no study has used vitamin C deficiency as an inclusion criterion. In contrast, reviewing those studies that have included a baseline determination of plasma vitamin C, only one of thirty-five studies (3 (95 % CI 0, 5) %) rendered it probable in a small sample that the subjects were in fact insufficient in vitamin C at the study start. Moreover, only five studies out of thirty-five studies (14 (95 % CI 2, 11) %) were available with data on vitamin C as a single substance.

This means that information from clinical trials with defined and verified vitamin C deficiency from a practical point of view is not available. In contrast, large and long-duration trials with β -carotene are available and show that 'hypervitaminosis' of β -carotene carries a risk for adverse effects on mortality (Alpha-Tocopherol, Beta-Carotene cancer prevention study, etc). It must therefore be concluded that at present we do not have the necessary scientific evidence to judge the effect on health – be that beneficial or deleterious – from vitamin C supplementation as a single substance. Dose–concentration relationships are largely available from pharmacokinetic evaluations, but no dose–response relationships for pharmacodynamic evaluation are available. For most of the available studies, the population status at entry with regard to vitamin C is unclear and may have been severely or marginally deficient, suboptimal or optimal. For the evaluation of the possible effect of vitamin C supplementation on human health, these studies are therefore largely irrelevant.

We had hoped that it would be possible to perform a meta-analysis of high-quality trials with vitamin C as a single substance based on the criteria suggested in Table 3, but have found that at this point this is not possible because such trials have not been performed.

In conclusion, we find that from a public health point of view, there is a dire need to examine the effect of vitamin C as a single supplement in populations which have been carefully defined with inclusion criteria of different levels of vitamin C status and with variable demand for vitamin C, for example, smokers *v.* non-smokers. The outcome markers should be defined and achieved targets, including plasma vitamin C concentration and relevant clinical endpoints.

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