

Increased plant sterol and stanol levels in brain of Watanabe rabbits fed rapeseed oil derived plant sterol or stanol esters

Christiane B. Fricke¹, Malene Schrøder², Morten Poulsen², Klaus von Bergmann¹, Ingmar Wester³, Ib Knudsen², Alicja Mortensen² and Dieter Lütjohann^{1*}

¹Department of Clinical Pharmacology, University of Bonn, Sigmund-Freud-Strasse 25, D-53105 Bonn, Germany

²Danish Institute for Food and Veterinary Research, Mørkhøj Bygade 19, DK-2860 Søborg Denmark

³Raisio Group, PO Box 101, FI-21201 Raisio, Finland

(Received 10 November 2006 – Revised 23 March 2007 – Accepted 12 April 2007)

Foods containing plant sterol or stanol esters can be beneficial in lowering LDL-cholesterol concentration, a major risk factor for CVD. The present study examined whether high dietary intake of rapeseed oil (RSO) derived plant sterol and stanol esters is associated with increased levels of these components in brain tissue of homozygous and heterozygous Watanabe rabbits, an animal model for familial hypercholesterolemia. Homozygous animals received either a standard diet, RSO stanol or RSO sterol ester while heterozygous animals were additionally fed with 2 g cholesterol/kg to the respective diet form for 120 d (*n* 9 for each group). Concentrations of cholesterol, its precursor lathosterol, plant sterols and stanols in brain and additionally in liver and plasma were determined by highly sensitive GC–MS. High-dose intake of RSO derived plant sterols and stanols resulted in increased levels of these components in plasma and liver. In brain a limited uptake of plant sterols and stanols was proven, indicating that these compounds passed the blood–brain barrier and may be retained in the brain tissue of Watanabe rabbits. Plant stanol ester feeding lowered plant sterol levels in brain, liver, and plasma. Cholesterol synthesis in brain, indicated by lathosterol, a local surrogate cholesterol synthesis marker, does not seem to be affected by plant sterol or stanol ester feeding. We conclude that high dose intake of plant sterol and stanol esters in Watanabe rabbits results in elevated concentrations of these components not only in the periphery but also in the central nervous system.

Phytosterols: Brain cholesterol: LDL receptor: Hypercholesterolemia: Atherosclerosis

Fortification of margarine and dairy products with plant sterol or plant stanol esters is a current dietary approach to lower elevated plasma cholesterol levels^{1–3}. High levels of total and LDL cholesterol are risk factors for atherosclerosis and CHD, the major cause of morbidity and mortality in the Western world⁴. Watanabe heritable hyperlipidemic (Watanabe) rabbits, the animal model used in the present study, have a defect of the LDL receptor and consequently develop hypercholesterolemia, resulting in atherosclerotic lesions very similar to those in human subjects and xantomas^{5–7}.

Comparable to human subjects the heterozygous form of hypercholesterolemia in Watanabe rabbits leads to a 2–3-fold increase in plasma cholesterol, while the homozygous form presents up to 10-fold higher circulating plasma cholesterol levels⁶. The present study was part of the European Union project NOFORISK ('Quantitative risk assessment strategies for novel food')⁸. In this project, a scientific methodology necessary for quantitative risk assessment of 'second generation' novel foods to be marketed in the European Union was developed and validated. The primary goal of the Watanabe rabbit experiments within the project was to evaluate the presumable athero-protective or atherogenic character of plant sterol or stanol ester supplementation in

cholesterol-fed heterozygous or chow-fed homozygous Watanabe rabbits. The present investigation examined, as a secondary goal, whether a high dietary intake of plant sterol or stanol esters is associated with increased levels of these components not only in the periphery (plasma and liver) but also in the central nervous system (brain) of heterozygous and homozygous Watanabe rabbits. It is generally assumed that cholesterol, plant sterols or stanols do not pass the blood–brain barrier⁹. However, the presence of the plant sterols campesterol and sitosterol have been reported in brain tissue of mice and man^{10–15}, while so far, the presence of plant stanols, the 5 α -saturated metabolites of sterols, in brain tissue was only reported in a patient with sitosterolemia¹⁶. Furthermore, we wanted to investigate whether the possible presence of plant sterols and stanols in brain of the Watanabe rabbits could influence local cholesterol synthesis and content.

Materials and methods

Animals and diets

Twenty-seven homozygous and twenty-seven heterozygous Watanabe rabbits of both sexes, six-weeks old, were obtained

Abbreviations: RSO, Rapeseed oil.

* **Corresponding author:** Dr Dieter Lütjohann, fax +49 228 287 11836, e-mail dieter.luetjohann@ukb.uni-bonn.de

from the in-house breeding colony of the Danish Institute for Food and Veterinary Research, Soeborg, Denmark (derived from a parent generation obtained from F.G.J. Janssen, University of Leiden, the Netherlands, with permission from Y. Watanabe). The animals were housed individually in stainless steel cages under controlled environmental conditions (temperature $18 \pm 2^\circ\text{C}$, relative humidity $55 \pm 5\%$, 12:12h light–dark cycle, air changed 10 times/h) and observed at least twice daily for any abnormalities in the clinical condition. Body weight was recorded weekly and feed intake daily. Rapeseed oil (RSO) sterols were sourced and either esterified as such with RSO fatty acids or first saturated to corresponding stanols and thereafter esterified with RSO fatty acids (Raisio Group, Raisio, Finland). The RSO sterol and stanol esters used in this study currently lack the required safety evaluation for their use as such in foods with added phytosterols. The homozygous Watanabe rabbits were randomised into three dietary groups receiving a standard diet (Altromin No. 2123; average content in the diet (g/kg): crude protein 120, crude fat 30, crude fibre 200, ash 90, moisture 120, nitrogen-free extract 440, and metabolisable energy 963 kJ/100 g; Altromin International, Lage, Germany), standard diet with added RSO stanol ester (34 g/kg, equivalent to 20 g plant stanols/kg) or standard diet with added RSO sterol ester (34 g/kg, equivalent to 20 g plant sterols/kg), respectively (nine rabbits in each group; body weight: 0.96 (SD 0.07) kg). The heterozygous animals were randomised into three dietary groups receiving standard diet with added cholesterol (2 g/kg), standard diet with added 2 g cholesterol/kg and 34 g RSO stanol esters/kg, or standard diet with added cholesterol (2 g/kg) and 34 g RSO sterol esters/kg (body weight: 1.05 (SD 0.040) kg). All animals were fed the specific diets (100 g/d) for 120 d and had free access to tap water. In homozygous rabbits, the body weight increased during the experimental period feeding standard diet, or this diet with added stanol or sterol esters to 2623 (SD 140) g, 2626 (SD 178) g and 2632 (SD 111) g, respectively. In heterozygous rabbits, the body weight increased to 2884 (SD 176) g, 2732 (SD 124) g and 2837 (SD 171) g, respectively. The recommended maximal daily intake of phytosterols for human subjects in the European Union-market is 3 g/d or 0.04 g/kg body weight, equivalent to a twenty times lower dose than applied to the rabbits in the present study. These high doses of plant sterols and stanols were chosen to achieve measurable changes. Animal studies were performed under conditions approved by the Danish Agency for Protection of Experimental Animals and by the in-house Animal Welfare Committee.

Plasma, tissue and feed sample preparation

Blood samples were collected into tubes containing sodium heparin from the marginal ear vein of unanaesthetised animals after overnight fasting. The samples were taken prior to the first feeding of the respective diet (baseline) and at termination (endpoint). The blood was placed on a rotation bed and centrifuged at 2500 g for 15 min. Plasma was immediately frozen and kept at -20°C until analysed. The rabbits were euthanised by intravenous injection of pentobarbital (100 mg/kg body weight) into the marginal ear vein, followed by exsanguination, and the central arterial system was perfused with about 300 ml chilled 0.9% NaCl solution through a canula, inserted

into the left ventricle of the heart. From the rostral half of the right side of the brain and from the left lobe of the liver, aliquots were excised, weighed and kept frozen at -20°C . Prior to sterol analysis, brain and liver specimens were spun in a Speedvac AES 100 (Servant Instruments Inc., Farmingdale, NY, USA) at 12 mbar for 48 h until completely dried. Wet and dry weights were noted and sterols and stanols were extracted from the dried tissues in 2 ml chloroform–methanol mixture (2:1; v/v) for 48 h. Feed was pulverised in a mortar and 7 ml chloroform–methanol mixture (2:1; v/v) was added to the pulver (30 mg) for extraction of sterols and stanols.

Sterol extraction and determination

300 μl brain and liver extracts and 2 ml feed chloroform–methanol sterol extracts were evaporated to dryness under a stream of N_2 at 63°C . After alkaline hydrolysis the free sterols and stanols were extracted twice with 3 ml cyclohexane and converted to trimethylsilyl ethers by adding 40 μl trimethylsilylation reagent (pyridine–hexamethyldisilazan–trimethylchlorosilane (9:3:1, by volume); Merck, Darmstadt, Germany). For determination of plasma sterol and stanol concentrations, sterols were extracted and derivatised from 50 μl plasma as described¹⁰. Feed was analysed in triplicate. Cholesterol concentrations were determined by GC-flame ionisation detection¹⁷. Levels of plant sterols (campesterol, sitosterol, brassicasterol and stigmasterol), plant stanols (campestanol, sitostanol) and the cholesterol precursor lathosterol were determined using GC-MS with epicoprostanol (Sigma, Deisenhof, Germany) as internal standard¹⁷. The sterols and stanols were monitored as their trimethylsilyl derivatives in the selected ion monitoring mode using the following masses: epicoprostanol m/z 370 ($M+$), lathosterol at m/z 458 ($M+$), campesterol at m/z 472 ($M+$), sitosterol at m/z 486 ($M+$), stigmasterol at m/z 484 ($M+$), brassicasterol at m/z 470 ($M+$), campestanol at m/z 474 ($M+$) and sitostanol at m/z 488 ($M+$). Peak integration was performed manually and sterols were quantified from selected-ion monitoring analyses against internal standards using standard curves for the listed sterols. Identity of all sterols was proven by comparison with the full-scan mass spectra of authentic compounds.

Statistical methods

The results were expressed as means with their standard deviations. Differences for plasma parameters between baseline and endpoint were calculated by paired Student's two-tailed t test using SPSS 12.0 for Windows (SPSS Inc., Chicago, IL, USA). Differences between the parameters for brain within the different groups were analysed by unpaired two-tailed t test. P values < 0.05 were considered significant.

Results

Sterol and stanol concentration in feed

Cholesterol and plant sterol/stanol concentrations in the experimental diets are given in Table 1. Each diet given to the heterozygous rabbits contained about 2 μg cholesterol/mg solid chow (2 g/kg), while in each diet applied to the homozygous rabbits only a minor amount of cholesterol ($< 0.03 \mu\text{g}/\text{mg}$

Table 1. Cholesterol and plant sterol and stanol concentration in the experimental diets (Mean values of triplicate analyses)

Feed	Cholesterol (µg/mg)	Campesterol (µg/mg)	Sitosterol (µg/mg)	Stigmasterol (µg/mg)	Brassicasterol (µg/mg)	Total plant sterols (µg/mg)	Campestanol (µg/mg)	Sitostanol (µg/mg)	Total plant stanols (µg/mg)
Heterozygous rabbits									
Cholesterol (2 g/kg)	1.97	0.09	0.33	0.019	0.003	0.44	0.05	0.17	0.22
Cholesterol (2 g/kg) + RSO stanol ester(34 g/kg)	2.04	0.21	0.41	0.021	0.022	0.66	9.34	11.4	20.7
Cholesterol (2 g/kg) + RSO sterol ester(34 g/kg)	1.91	7.53	9.69	0.072	1.976	19.3	0.17	0.16	0.33
Homozygous rabbits									
Standard diet	0.01	0.09	0.27	0.031	0.003	0.39	0.04	0.15	0.20
RSO stanol ester(34 g/kg)	0.02	0.24	0.41	0.022	0.019	0.69	9.38	11.7	21.0
RSO sterol ester(34 g/kg)	0.01	7.95	10.2	0.078	2.204	20.5	0.19	0.17	0.36

RSO, rapeseed oil.

dry weight) could be detected. Percentage content of plant sterols and stanols in the different feed for heterozygous and homozygous rabbits, respectively, was comparable.

Plant sterol and stanol concentration in plasma, brain and liver

Heterozygous Watanabe rabbits. Feeding RSO plant stanol esters to heterozygous Watanabe rabbits resulted in a significantly higher total plant stanol concentration in plasma, brain, and liver compared to rabbits fed with cholesterol only (Tables 2 and 3). By contrast, total plant sterol concentrations in plasma, brain, and liver were lower in animals fed with RSO plant stanol esters, while total phytosterol levels (sum of plant sterols and plant stanols) were only lowered in the brains of animals of this group. In RSO sterol ester-fed rabbits not only total plant sterol but also total plant stanol concentration in plasma, brain and liver tissues increased significantly compared to control. Therefore, total phytosterol level in RSO sterol ester-fed rabbits was about 7 times ($P < 0.001$) higher in plasma and brain and 13 times ($P < 0.001$) higher in liver compared with the rabbits fed with cholesterol only (Tables 2 and 3). Total plant stanol and sterol levels were significantly higher in the liver compared with brains from all animals (Fig. 1). Furthermore, in plasma from heterozygous Watanabe rabbits, the level of campesterol in the RSO sterol ester-fed group was about 25 times higher than the plasma level of campestanol in the RSO stanol ester-fed group. In brain, tissue level of campesterol in the RSO sterol ester-fed animals was about 50 times higher than the brain tissue level of campestanol in the RSO stanol ester-fed group while in liver it was about 24 times higher. The campesterol:sitosterol and campestanol:sitostanol ratios in plasma and brain are given in Table 4. In brain these ratios were significantly lower in all groups of heterozygous rabbits compared with the ratios in plasma.

Homozygous Watanabe rabbits. In homozygous rabbits, we also observed lower plasma, brain and liver total plant sterol and total phytosterol levels in the 34 g/kg RSO stanol ester-fed rabbits compared with standard diet-fed animals while total plant stanol levels were higher (Tables 5 and 6). Moreover, as in heterozygous rabbits RSO plant sterol ester feeding increased total plant sterol and total plant stanol concentration in plasma, brain and liver. The concentrations of total phytosterols in RSO sterol ester-fed rabbits was about 4.4 times ($P < 0.001$) higher in plasma and brain and 5.3 times ($P < 0.001$) higher in liver compared with standard diet-fed rabbits (Tables 5 and 6). As already seen in heterozygous animals total plant stanol and sterol levels were significantly higher in the liver compared with brains from all animals (Fig. 2). In plasma of homozygous Watanabe rabbits, the level of campesterol in the RSO sterol ester-fed group was about 11 times higher than the plasma level of campestanol in the RSO stanol ester-fed rabbits. In brain tissue in the RSO sterol ester-fed rabbits campesterol concentration was 22 times and in the liver 17 times higher than campestanol in the RSO stanol ester-fed animals.

It was interesting that in the different groups of homozygous rabbits, plasma levels of plant sterols and plant stanols were approximately 5–15 times higher than in heterozygous rabbits, respectively, but concentrations of these substances

Table 2. Cholesterol, lathosterol, lathosterol:cholesterol ratio and plant sterols and stanols in plasma before and during rapeseed oil derived plant sterol and plant stanol feeding with continuous cholesterol intake (0.2 g/kg) for 120 d in heterozygous Watanabe rabbits (Mean values with standard deviations for nine rabbits)

Plant sterol/stanol	Cholesterol (2 g/kg)				Cholesterol (2 g/kg) + stanol ester (34 g/kg)				Cholesterol (2 g/kg) + sterol ester (34 g/kg)			
	Baseline		Endpoint		Baseline		Endpoint		Baseline		Endpoint	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Cholesterol (mg/dl)	107	29.6	490*	370	139	33.1	158†	85.2	123	36.3	177†	115
Lathosterol (mg/dl)	0.09	0.05	0.22	0.19	0.10	0.03	0.06*†	0.02	0.08	0.04	0.09	0.08
Lathosterol:Cholesterol (μg/dl)	0.84	0.22	0.41***	0.12	0.75	0.14	0.43***	0.15	0.62	0.25	0.50*	0.22
Campesterol (mg/dl)	1.38	0.37	2.22	1.76	1.99	0.55	0.63*** †	0.36	1.82	0.38	20.3*** ††	11.7
Sitosterol (mg/dl)	0.42	0.15	0.65	0.35	0.57	0.18	0.19*** ††	0.07	0.54	0.11	1.39** †	0.72
Brassicasterol (μg/dl)	40.2	12.7	30.8	25.6	53.8	14.3	13.9***	3.15	42.1	9.13	142* †	106
Stigmasterol (μg/dl)	5.92	0.87	9.78*	4.14	4.88	2.17	2.52** †††	2.75	7.24	0.96	5.14** ††	1.06
Total plant sterols (mg/dl)	1.84	0.51	2.91	2.13	2.62	0.72	0.84*** †	0.42	2.41	0.47	21.8*** †††	12.4
Campestanol (μg/dl)	31.8	6.18	68.0	47.1	42.8	15.9	817*** †††	454	40.3	9.69	229*** †††	93.1
Sitostanol (μg/dl)	13.0	2.83	16.0	8.64	17.4	8.17	100**††	59.8	16.2	2.91	25.3	15.3
Total plant stanols (μg/dl)	44.8	8.84	84.0*	55.5	60.1	23.6	917*** †††	513	56.5	12.3	254*** †††	103
Total phytosterols (mg/dl)‡	1.88	0.51	2.99	2.19	2.68	0.74	1.75*	0.93	2.47	0.48	22.1*** †††	12.5

Mean values were significantly different at endpoint from baseline: **P* < 0.05, ***P* < 0.01, ****P* < 0.001.
 Mean values were significantly different at endpoint v. endpoint cholesterol (2 g/kg): †*P* < 0.05, ††*P* < 0.01, †††*P* < 0.001.
 ‡ Sum of plant sterols and plant stanols.

Table 3. Cholesterol, lathosterol, the lathosterol:cholesterol ratio and plant sterols/stanols in brain and liver after rapeseed oil derived plant sterol and plant stanol feeding during continuous cholesterol intake (2 g/kg) for 120 d in heterozygous Watanabe rabbits (Mean values with standard deviations for nine rabbits)

Plant sterol/ stanol	Brain						Liver					
	Cholesterol (2 g/kg)		Cholesterol (2 g/kg) + stanol ester (34 g/kg)		Cholesterol (2 g/kg) + sterol ester (34 g/kg)		Cholesterol (2 g/kg)		Cholesterol (2 g/kg) + stanol ester (34 g/kg)		Cholesterol (2 g/kg) + sterol ester (34 g/kg)	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Cholesterol (μg/mg)	70.7	8.24	63.8*	4.66	59.7*	10.5	14.0	5.89	7.35*	3.02	6.04**	2.64
Lathosterol (μg/mg)	0.13	0.02	0.10**	0.01	0.11	0.02	0.005	0.002	0.005	0.004	0.008	0.010
Lathosterol:Cholesterol (ng/mg)	1.78	0.23	1.61	0.14	1.82	0.20	0.37	0.20	0.83	0.65	1.71	2.82
Campesterol (ng/mg)	17.1	3.69	7.78***	2.22	200***	49.6	50.4	19.7	26.4**	12.2	822***	272
Sitosterol (ng/mg)	11.9	2.07	6.13***	0.69	19.3***	4.07	14.8	5.43	6.00***	3.87	47.1***	17.4
Brassicasterol (ng/mg)	1.51	0.20	1.13***	0.22	4.04***	1.06	1.04	0.67	0.54	0.28	6.19***	2.80
Stigmasterol (ng/mg)	0.67	0.14	0.45***	0.08	0.57	0.16	0.21	0.12	0.14	0.06	0.20	0.06
Total plant sterols (ng/mg)	31.2	5.94	15.5***	2.62	224***	53.9	66.5	23.5	33.3**	15.7	876***	285
Campestanol (ng/mg)	0.71	0.14	4.00***	0.87	3.00	0.67	1.42	0.44	34.1***	18.1	8.36***	3.45
Sitostanol (ng/mg)	0.45	0.12	0.81***	0.24	0.60*	0.13	0.45	0.13	2.79***	1.32	0.55	0.27
Total plant stanols (ng/mg)	1.16	0.23	4.81***	1.01	3.60***	0.77	1.87	0.56	36.9***	19.4	8.91***	3.71
Total phytosterols (ng/mg)†	32.4	6.1	20.3***	3.03	228***	54.5	68.6	24.1	70.2	33.7	885***	289

Mean values were significantly different from those for cholesterol (2 g/kg): **P* < 0.05, ***P* < 0.01, ****P* < 0.001.
 † Sum of plant sterols and plant stanols

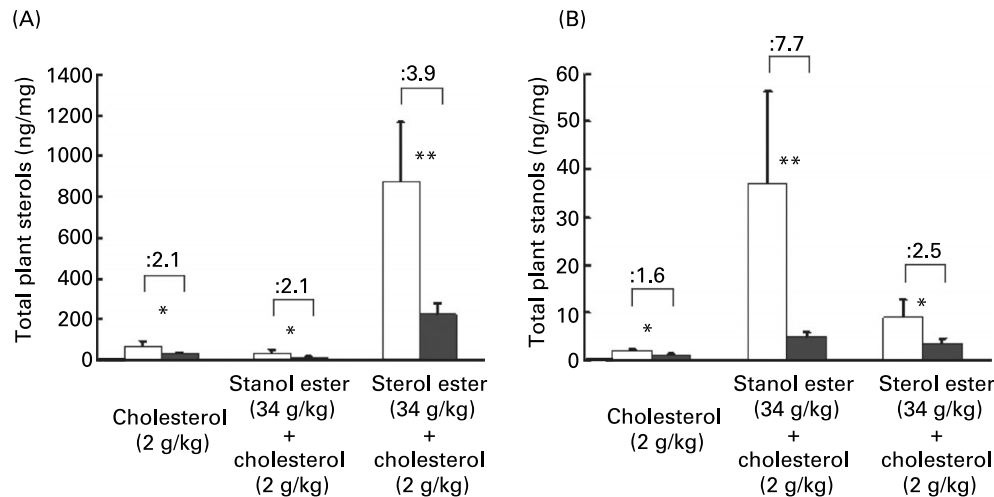


Fig. 1. Total plant sterol (A) and stanol (B) concentrations in brain (■) and liver (□) after rapeseed oil derived plant sterol and plant stanol feeding for 120 d in heterozygous Watanabe rabbits (n 9). Values are means with standard deviations shown by vertical bars. Mean values were significantly different between brain and liver: * P <0.01, ** P <0.001. Values above columns represent the quotient between liver and brain concentrations.

in brain were only 1–2 times higher. The campesterol:sitosterol and campestanol:sitostanol ratios in brains of homozygous animals was significantly lower compared with the ratios in plasma (Table 4).

Cholesterol, lathosterol and the lathosterol: cholesterol ratio in plasma, brain, and liver

Heterozygous Watanabe rabbits. Heterozygous rabbits fed for 120 d with cholesterol (2 g/kg) only, showed on average a 3.6-fold ($P=0.013$) increase in their plasma cholesterol level while pre-feeding (baseline) and terminal (endpoint) plasma cholesterol levels were not significantly different in the animals additionally fed plant stanol or sterol esters (Table 2). Feeding the rabbits with cholesterol (2 g/kg) and RSO sterol or stanol esters for 120 d resulted in lower cholesterol concentrations in plasma, brain, and liver of the animals compared with rabbits fed with cholesterol only (Tables 2 and 3). In plasma the lathosterol:cholesterol ratio, indicator of endogenous cholesterol synthesis, decreased in all groups during the feeding period for 120 d. Differences in endpoint

lathosterol:cholesterol ratios between the groups, in plasma as well as in brain and liver were not detected. Absolute lathosterol concentration was only lower in plasma and brain of the rabbits fed with 2 g cholesterol/kg and RSO stanol esters compared with the rabbits fed with cholesterol only (Tables 2 and 3). Significantly higher cholesterol levels were only found in female (258 (SD 60.5) mg/dl) compared with male (108 (SD 32.3) mg/dl, $P=0.001$) heterozygous rabbits fed with cholesterol and plant stanol ester after 120 d. However, baseline *v.* endpoint comparison of cholesterol in female and male animals of this feeding group revealed no significant difference (data not shown). Thus, the data for cholesterol in male and female rabbits were merged.

Homozygous Watanabe rabbits. In homozygous rabbits, no significant differences were found in the pre-feeding and terminal plasma cholesterol level in any of the three groups (Table 5). Furthermore, no differences in plasma, brain, and liver cholesterol level between the groups could be detected (Tables 5 and 6). As seen in plasma of heterozygous rabbits, feeding with different diets for 120 d resulted in a decrease of the lathosterol:cholesterol ratio in all groups. Differences

Table 4. Campesterol: sitosterol and campestanol:sitostanol ratios in plasma and brain after rapeseed oil (RSO) derived plant sterol and plant stanol ester feeding for 120 d in heterozygous and homozygous Watanabe rabbits (Mean values with standard deviations for nine rabbits)

Feed	Campesterol:sitosterol				Campestanol:sitostanol			
	Plasma		Brain		Plasma		Brain	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Heterozygous rabbits								
Cholesterol (2 g/kg)	2.99	1.14	1.41**	0.14	3.98	0.82	1.65***	0.40
Cholesterol (2 g/kg) + RSO stanol ester (34 g/kg)	3.28	1.06	1.26***	0.31	8.33	0.85	5.10***	1.13
Cholesterol (2 g/kg) + RSO sterol ester (34 g/kg)	15.2	5.44	10.4*	1.33	10.4	3.58	5.04**	0.63
Homozygous rabbits								
Standard diet	2.28	0.33	2.02*	0.22	3.28	0.58	1.42***	0.25
RSO stanol ester (34 g/kg)	2.71	0.40	1.31***	0.40	7.73	1.21	4.82***	0.91
RSO sterol ester (34 g/kg)	10.0	1.29	9.58	0.97	8.33	2.32	3.21***	0.63

Mean values were significantly different between brain and plasma: * P < 0.05, ** P < 0.01, *** P <0.001

Table 5. Cholesterol, lathosterol, the lathosterol:cholesterol ratio and plant sterols and stanols in plasma before and during rapeseed oil derived plant sterol and plant stanol feeding for 120 d in homozygous Watanabe rabbits
(Mean values with standard deviations for nine rabbits)

Plant sterol/stanol	Standard diet				Stanol ester (34 g/kg)				Sterol ester (34 g/kg)			
	Baseline		Endpoint		Baseline		Endpoint		Baseline		Endpoint	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Cholesterol (mg/dl)	960	133	966	139	925	243	1016	271	951	175	884	146
Lathosterol (mg/dl)	1.07	0.30	0.63***	0.31	1.32	0.47	0.62***	0.21	1.16	0.27	0.68***	0.09
Lathosterol:Cholesterol (µg/mg)	1.13	0.35	0.66**	0.26	1.46	0.45	0.67***	0.31	1.23	0.28	0.78***	0.10
Campesterol (mg/dl)	11.9	2.17	16.5***	2.67	11.7	1.61	6.37***†††	1.50	10.4	1.34	99.5***†††	18.6
Sitosterol (mg/dl)	5.18	1.06	7.30**	1.13	4.80	0.73	2.40***†††	0.62	3.49	0.70	9.95***†††	1.60
Brassicasterol (µg/dl)	323	56.0	211**	111	334	44.5	67.6***††	30.8	356	54.0	1022***†††	210
Stigmasterol (µg/dl)	44.0	14.7	64.3**	9.04	41.6	12.2	15.7***†††	4.02	31.6	10.6	12.6***†††	0.96
Total plant sterols (mg/dl)	17.4	3.23	24.1***	3.53	16.9	2.24	8.85***†††	2.06	14.3	2.07	111***†††	19.8
Campestanol (µg/dl)	391	74.2	780***	154	313	124	8657***†††	1879	283	56.9	1141***†††	265
Sitostanol (µg/dl)	188	33.9	243*	52.2	167	82.4	1129***†††	250	140	26.5	140†††	22.0
Total plant stanols (µg/dl)	579	96.4	1023***	189	480	190	9787***†††	2076	423	82.1	1281***†	261
Total phytosterols (mg/dl)‡	18.0	3.24	25.1***	3.63	17.3	2.27	18.7††	3.80	14.7	2.14	112***†††	20.0

Mean values were significantly different at endpoint v. baseline: **P* < 0.05, ***P* < 0.01, ****P* < 0.001.
Mean values were significantly different at endpoint v. endpoint Standard diet: †*P* < 0.05, ††*P* < 0.01, †††*P* < 0.001.
‡ Sum of plant sterols and plant stanols.

Phytosterols in brain of Watanabe rabbits

Table 6. Cholesterol, lathosterol, the lathosterol:cholesterol ratio and plant sterols/stanols in brain and liver after rapeseed oil derived plant sterol and plant stanol feeding for 120 d in homozygous Watanabe rabbits
(Mean values with standard deviations for nine rabbits)

Plant sterol/ stanol	Brain						Liver					
	Standard diet		Stanol ester (34 g/kg)		Sterol ester (34 g/kg)		Standard diet		Stanol ester (34 g/kg)		Sterol ester (34 g/kg)	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Cholesterol (µg/mg)	60.5	10.5	70.3	9.39	62.8	10.1	7.66	1.32	8.61	0.97	7.25	1.48
Lathosterol (µg/mg)	0.11	0.03	0.13	0.02	0.11	0.02	0.014	0.007	0.022	0.014	0.016	0.010
Lathosterol:Cholesterol (ng/mg)	1.89	0.21	1.79	0.22	1.82	0.12	1.81	0.86	2.62	1.81	2.26	1.59
Campesterol (ng/mg)	32.2	9.27	15.8***	5.61	198***	42.0	163	32.0	63.1***	18.0	1102***	376
Sitosterol (ng/mg)	16.0	3.24	11.8**	1.92	20.6*	3.64	56.4	14.5	23.0***	7.91	102**	32.8
Brassicasterol (ng/mg)	1.95	0.53	1.46*	0.21	3.80***	0.72	3.67	1.09	2.15**	0.73	11.5***	3.98
Stigmasterol (ng/mg)	0.86	0.18	0.79	0.23	0.73	0.19	1.75	0.16	1.12***	0.41	0.90***	0.21
Total plant sterols (ng/mg)	51.0	12.9	29.8***	6.63	223***	45.6	225	42.9	89.4***	25.8	1216***	408
Campestanol (ng/mg)	0.96	0.18	8.82***	1.93	2.36***	0.56	4.27	1.32	66.5***	25.2	11.8**	5.78
Sitostanol (ng/mg)	0.69	0.16	1.85***	0.35	0.73	0.11	1.41	0.64	7.77***	2.27	1.20	0.34
Total plant stanols (ng/mg)	1.66	0.31	10.7***	2.14	3.09***	0.64	5.67	1.89	74.3***	27.4	13.0**	6.08
Total phytosterols (ng/mg)†	52.6	13.1	40.6*	8.64	226***	46.1	231	44.3	164**	50.3	1229***	414

Mean values were significantly different from those for the standard diet: **P* < 0.05, ***P* < 0.01, ****P* < 0.001.
† Sum of plant sterols and plant stanols.

895

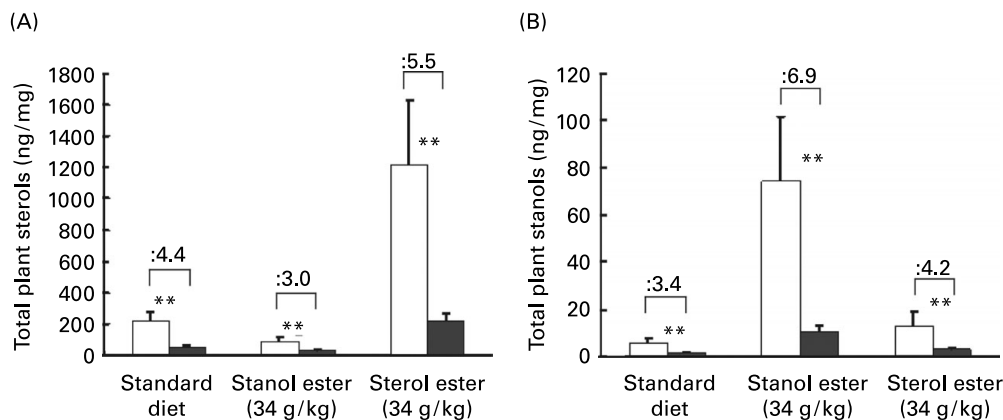


Fig. 2. Total plant sterol (A) and stanol (B) concentrations in brain (■) and liver (□) after rapeseed oil derived plant sterol and plant stanol feeding for 120 d in homozygous Watanabe rabbits (n 9). Values are means with standard deviations shown by vertical bars. Mean values were significantly different between brain and liver: ** $P < 0.001$. Values above columns represent the quotient between liver and brain concentrations.

between the groups in plasma as well as in brain and liver were not observed (Tables 5 and 6). Independent of the diet, baseline lathosterol concentrations in plasma were higher in all homozygous rabbits compared with endpoint lathosterol concentrations (Table 5). Differences in lathosterol levels in plasma, brain, and liver between the dietary groups were not proven. As in heterozygous rabbits, significantly higher cholesterol levels in female (1222 (SD 131) mg/dl) compared with male (758 (SD 113) mg/dl, $P = 0.001$) homozygous rabbits were found in the stanol ester-fed group after 120 d. Since baseline *v.* endpoint comparison of cholesterol in female and male animals of this feeding group also revealed no significant difference (data not shown), the data for cholesterol in male and female homozygous rabbits were merged. In homozygous Watanabe rabbits, pre-feeding cholesterol levels in plasma were approximately up to 10 times higher than in heterozygous rabbits. Although in homozygous Watanabe rabbits cholesterol levels in plasma were much higher, cholesterol concentrations in their brains did not differ from those found in heterozygous rabbits.

Discussion

Feeding a diet fortified with RSO plant sterol or stanol esters to heterozygous and homozygous Watanabe rabbits, resulted in increased levels of plant sterols and stanols not only in peripheral compartments (plasma and liver) but also in the brain. Consequently, plant sterols and stanols seem to be able to cross the blood–brain barrier. Increase in plasma, brain and hepatic plant stanol concentrations in RSO sterol ester-fed rabbits indicated that plant sterols are converted to plant stanols within the mammalian body. Application of plant stanol esters increased plant stanol levels while plant sterol levels decreased in plasma, brain and liver. In homozygous Watanabe rabbits, RSO plant sterols and stanols had no influence on plasma, brain and liver cholesterol. In contrast, in cholesterol-fed heterozygous rabbits, feeding a diet with added RSO plant sterols or stanols decreased cholesterol levels in plasma, brain and liver when compared to animals receiving the diet with only 2 g cholesterol/kg added. In plasma of non-cholesterol-fed homozygous and cholesterol-fed heterozygous rabbits the lathosterol:cholesterol ratio, indicator of

endogenous cholesterol synthesis, decreased in all groups during the feeding period for 120 d while no differences in endpoint lathosterol:cholesterol ratios between the groups, in plasma as well as in brain and liver, were detected.

In the brains of non-sitosterolemic human subjects and wild-type animals low concentrations of plant sterols, ranging from 10–80 ng/mg, have been described^{10,11,14,18–21}. In the brain of a patient with sitosterolemia, a rare, inherited sterol storage disease with a defect in the gene(s) for ABCG5/G8 transporters^{22,23}, campesterol (200 ng/mg) was found in concentrations comparable to sterol ester-fed rabbits of the present study¹⁶. ABCG5 and ABCG8 are half transporters which are expressed in the intestine and the liver where they re-secrete plant sterols and stanols back from the enterocytes into the lumen of the intestine, or mediate efflux of phytosterols and stanols as well as cholesterol from hepatocytes into the bile^{4,24}. Plasma levels of plant sterols in sitosterolemic patients^{16,25} and in the rabbits used in the present study were also comparable. The only difference between our model and patients with sitosterolemia was that these patients had higher concentrations of sitosterol than campesterol in plasma and tissues. Recent studies demonstrated a 7- to 20-fold increase in brain plant sterol levels in chow-fed ABCG5/G8 deficient mice^{11,12} and an increase of about 30% in plant sterol-fed (8.3 g/kg) wild type mice¹².

There are a great number of animal studies using plant stanol and sterol ester fortified food; however, only data on serum and liver but not on brain are available. In the present study, it was demonstrated that an increase in plasma plant sterol and stanol levels may result in increased levels of plant sterols and stanols in liver and in the brain. However, in healthy human subjects serum levels of phytosterols are typically much lower^{26–29} than those obtained in the present study with Watanabe rabbits. As a consequence, brain tissue levels of this component are assumed to be also lower. From previous studies it is well established that patients homozygous and heterozygous for familial hypercholesterolemia not only present increased cholesterol but also plant sterol levels. A plant sterol rich diet in combination with statins in these patients results in even more elevated serum plant sterol levels³⁰. Whether or not this therapeutic approach in these patients has an effect on their plant sterol level in

brain needs to be investigated, possibly in combination with neurological examination using cerebrospinal fluids.

Our findings provide further evidence that not only plant sterols but also plant stanols can cross the blood–brain barrier or that they are formed locally from sterols. Increased amounts of the plant stanols, campestanol and sitostanol, were also found in plasma, brain and liver in the animals fed with RSO sterol esters. Presumably, these stanols could have been produced endogenously from their unsaturated precursors campesterol and sitosterol in the mammalian body, also demonstrated by a previous study in subjects with sitosterolemia¹⁵. In fact, feed enriched with 34 g RSO sterol esters/kg also contained small amounts of plant stanols, probably formed during the production process. However, percentage distribution of campestanol and sitostanol in plasma (approximately 90% and 10%, respectively) and brain (80% and 20%, respectively) compared with feed (50% and 50%, respectively) indicates that the shift is comparable to that of campesterol to sitosterol (about 90% and 10% in brain and plasma, respectively).

The question remains as to how plant sterols and stanols are transported into the brain. In a previous study, chow-fed ABCG5 and ABCG8-deficient and plant sterol-fed (8.3 g/kg) wild type mice showed significantly elevated plant sterol levels, especially of campesterol, in the brain, while in ApoE-deficient mice they were not increased¹². Watanabe rabbits show a 4- to 7-fold increase in plasma levels of ApoE³¹. Therefore, the possibility remains that ApoE is involved in this transport. ABCG5/8 transporter expression at the level of the blood–brain barrier is extremely low and it appears very unlikely that they play a role in plant sterol transport¹².

It is noteworthy, that despite the approximately 10-fold higher plasma and liver plant sterol and stanol levels in homozygous Watanabe rabbits compared to heterozygous Watanabe rabbits, the concentrations of plant sterols and stanols in the brain of these animals did not reveal such differences. This finding indicates a limit for sterol and/or for stanol uptake into the brain. Independent from the genotype and diet of the Watanabe animals, the plant sterol and stanol levels in the liver are significantly higher than in the brain (Figs. 1 and 2). Thus, the blood–brain barrier seems to protect the brain from excessive uptake of circulating phytosterols.

The campesterol:sitosterol ratios in plasma and brain were different (Table 4). Therefore, it seems obvious that high levels of non-cholesterol sterols in brain are not attributed to contaminations from the circulation by microvessels. In agreement with a previous study³², campesterol uptake within the intestine was presumably higher than that of sitosterol, indicated by higher plasma campesterol concentrations. By contrast, the lower campesterol:sitosterol and campestanol:sitostanol ratios in brain compared to these ratios in plasma reflected that sitosterol and sitostanol seem to pass the blood–brain barrier easier than campesterol and campestanol. Either there was relatively more sitosterol and sitostanol taken up into the brain or the excretion of campesterol and campestanol from the brain was higher than that for sitosterol and sitostanol.

In addition to previous reports dietary plant stanol ester reduced tissue levels of plant sterols not only in plasma and liver but also in the brain with recorded increases in levels of campestanol and sitostanol^{26,33}. However, the plasma,

liver and brain levels of campestanol and sitostanol in Watanabe rabbits fed with RSO stanol ester-added diet still remained at low levels compared to campesterol and sitosterol in the rabbits fed with RSO sterol esters.

Feeding cholesterol and RSO plant sterol or stanol ester to heterozygous and homozygous Watanabe rabbits had no influence on cholesterol synthesis in brain. In heterozygous rabbits only we found a lower cholesterol content in the RSO sterol and stanol ester-fed rabbits. In contrast, in plant stanol ester-fed homozygous rabbits cholesterol content in brain was not significantly but about 16% higher compared with standard diet-fed animals. However, we are unable to provide a valid explanation for this. It was previously demonstrated that an increased dietary intake of cholesterol in New Zealand white rabbits was associated with suppression of cholesterol synthesis in many extrahepatic tissues, but not in brain³⁴. Additionally brain cholesterol content did not change either. Increased levels of plant sterols in brain of ABCG5/8 KO mice also showed no influence on cholesterol synthesis¹². In sitosterolemic patients with increased plasma and brain non-cholesterol concentrations, brain cholesterol content did not differ from that of control subjects¹⁶. This confirms strict regulation of cholesterol metabolism in brain.

In homozygous Watanabe rabbits, RSO plant sterols and stanols had no influence on plasma cholesterol concentration, while in the cholesterol-fed heterozygous rabbits, supplementation of the diet with plant sterol or stanol esters decreased plasma cholesterol levels when compared to the animals receiving the diet with added cholesterol but no plant sterol or stanol esters. Plant sterols and stanols are noted for reducing cholesterol absorption by displacing cholesterol from mixed micelles^{35,36}. In the present study, homozygous rabbits were not fed with cholesterol. Hence, no effect on dietary cholesterol absorption of plant sterols and stanols in the rabbits could be expected. Furthermore the extraction of biliary cholesterol may be impaired in homozygous Watanabe rabbits, leading to a situation where very limited amounts of cholesterol enters the digestive tract daily.

Cholesterol synthesis marker, the lathosterol:cholesterol ratio, in plasma, decreased in both homozygous and heterozygous rabbits, independent of the diet. In heterozygous rabbits this was caused by an increase of absolute amounts of cholesterol, though not significantly, in the plant stanol and sterol ester group (Table 2). In the homozygous animals absolute cholesterol levels remained unchanged while lathosterol itself decreased (Table 5). Thus the lathosterol:cholesterol ratio also decreased significantly in each group. As shown in a previous study, endogenous cholesterol synthesis decreased with aging in homozygous Watanabe animals³⁷. This was in agreement with the results in homozygous but not with heterozygous animals in the present study. Unfortunately, we do not have a comparison in heterozygous animals which did not receive cholesterol. Thus, this biochemical marker does not provide an answer to the question whether feeding of cholesterol, RSO plant sterols or stanols has an effect on cholesterol synthesis in these rabbits.

The results with these novel RSO sterol and stanol ester preparations are valid for the Watanabe rabbit model used but whether the use of commercial food with added plant sterols and stanols also cause a similar elevation or reduction of total phytosterols in the brain tissue of man remains to be

elucidated. To clarify the mechanisms by which plant sterols and stanols are transported into the brain and whether they have an influence on brain function(s) requires further investigation.

Acknowledgements

This work was completed on behalf of the NOFORISK Consortium and funded under the EU 6th Framework Food Quality and Safety Programme, project number FP6-506387. The authors thank Silvia Friedrichs, Meryem Gültekin, Merete Lykkegaard, Karen Roswall and the animal technicians for technical assistance.

References

- Devaraj S & Jialal I (2006) The role of dietary supplementation with plant sterols and stanols in the prevention of cardiovascular disease. *Nutr Rev* **64**, 348–354.
- Katan MB, Grundy SM, Jones P, Law M, Miettinen T & Paoletti R (2003) Efficacy and safety of plant stanols and sterols in the management of blood cholesterol levels. *Mayo Clin Proc* **78**, 965–978.
- Plat J & Mensink RP (2005) Plant stanol and sterol esters in the control of blood cholesterol levels: mechanism and safety aspects. *Am J Cardiol* **96**, 15D–22D.
- Grundy SM, Cleeman JI, Merz CN, Brewer HB Jr, Clark LT, Hunninghake DB, Pasternak RC, Smith SC Jr & Stone NJ (2004) Implications of recent clinical trials for the National Cholesterol Education Program Adult Treatment Panel III guidelines. *Circulation* **110**, 227–239.
- Watanabe Y, Ito T & Shiomi M (1985) The effect of selective breeding on the development of coronary atherosclerosis in WHHL rabbits. An animal model for familial hypercholesterolemia. *Atherosclerosis* **56**, 71–79.
- Kita T, Brown MS, Watanabe Y & Goldstein JL (1981) Deficiency of low density lipoprotein receptors in liver and adrenal gland of the WHHL rabbit, an animal model of familial hypercholesterolemia. *Proc Natl Acad Sci U S A* **78**, 2268–2272.
- Watanabe Y (1980) Serial inbreeding of rabbits with hereditary hyperlipidemia (WHHL-rabbit). *Atherosclerosis* **36**, 261–268.
- NOFORISK (2007) Quantitative risk assessment strategies for novel foods. <http://www.NOFORISK.org>.
- Dietschy JM & Turley SD (2004) Thematic review series: brain Lipids. Cholesterol metabolism in the central nervous system during early development and in the mature animal. *J Lipid Res* **45**, 1375–1397.
- Lütjohann D, Brzezinka A, Barth E, Abramowski D, Staufenbiel M, von Bergmann K, Beyreuther K, Multhaup G & Bayer TA (2002) Profile of cholesterol-related sterols in aged amyloid precursor protein transgenic mouse brain. *J Lipid Res* **43**, 1078–1085.
- Yu L, von Bergmann K, Lütjohann D, Hobbs HH & Cohen JC (2004) Selective sterol accumulation in ABCG5/ABCG8-deficient mice. *J Lipid Res* **45**, 301–307.
- Jansen PJ, Lütjohann D, Abildayeva K, *et al.* (2006) Dietary plant sterols accumulate in the brain. *Biochim Biophys Acta* **1761**, 445–453.
- Thelen KM, Rentsch KM, Gutteck U, Heverin M, Olin M, Andersson U, von Eckardstein A, Björkhem I & Lütjohann D (2006) Brain cholesterol synthesis in mice is affected by high dose of simvastatin but not of pravastatin. *J Pharmacol Exp Ther* **316**, 1146–1152.
- Heverin M, Bogdanovic N, Lütjohann D, Bayer T, Pikuleva I, Bretillon L, Diczfalusy U, Winblad B & Björkhem I (2004) Changes in the levels of cerebral and extracerebral sterols in the brain of patients with Alzheimer's disease. *J Lipid Res* **45**, 186–193.
- Salen G, Kwiterovich PO Jr, Shefer S, Tint GS, Horak I, Shore V, Dayal B & Horak E (1985) Increased plasma cholestanol and 5 alpha-saturated plant sterol derivatives in subjects with sitosterolemia and xanthomatosis. *J Lipid Res* **26**, 203–209.
- Salen G, Horak I, Rothkopf M, *et al.* (1985) Lethal atherosclerosis associated with abnormal plasma and tissue sterol composition in sitosterolemia with xanthomatosis. *J Lipid Res* **26**, 1126–1133.
- Lütjohann D, Stroick M, Bertsch T, *et al.* (2004) High doses of simvastatin, pravastatin, and cholesterol reduce brain cholesterol synthesis in guinea pigs. *Steroids* **69**, 431–438.
- Dorszewska J & Adamczewska-Goncerzewicz Z (2000) Patterns of free and esterified sterol fractions of the cerebral white matter in severe and moderate experimental hypoxia. *Med Sci Monit* **6**, 227–231.
- Kritchevsky D & Defendi V (1962) Deposition of tritium labelled sterols (cholesterol, sitosterol, lanosterol) in brain and other organs of the growing chicken. *J Neurochem* **9**, 421–425.
- Prasad VV, el-Maraghy E, Ponticorvo L & Lieberman S (1985) Detection in extracts of bovine brain of lipophilic complexes of sulfate esters of cholesterol and beta-sitosterol. *Proc Natl Acad Sci U S A* **82**, 2657–2659.
- Ratnayake WM, Plouffe L, Hollywood R, L'Abbe MR, Hidiroglou N, Sarwar G & Mueller R (2000) Influence of sources of dietary oils on the life span of stroke-prone spontaneously hypertensive rats. *Lipids* **35**, 409–420.
- Bhattacharyya AK & Connor WE (1974) Beta-sitosterolemia and xanthomatosis. A newly described lipid storage disease in two sisters. *J Clin Invest* **53**, 1033–1043.
- Lütjohann D & von Bergmann K (1997) Phytosterolaemia: diagnosis, characterization and therapeutical approaches. *Ann Med* **29**, 181–184.
- Yu L, Hammer RE, Li-Hawkins J, Von Bergmann K, Lütjohann D, Cohen JC & Hobbs HH (2002) Disruption of Abcg5 and Abcg8 in mice reveals their crucial role in biliary cholesterol secretion. *Proc Natl Acad Sci U S A* **99**, 16237–16242.
- Lütjohann D, Björkhem I, Beil UF & von Bergmann K (1995) Sterol absorption and sterol balance in phytosterolemia evaluated by deuterium-labeled sterols: effect of sitostanol treatment. *J Lipid Res* **36**, 1763–1773.
- Gylling H, Puska P, Vartiainen E & Miettinen TA (1999) Serum sterols during stanol ester feeding in a mildly hypercholesterolemic population. *J Lipid Res* **40**, 593–600.
- Jones PJ, Raeni-Sarjaz M, Ntanos FY, Vanstone CA, Feng JY & Parsons WE (2000) Modulation of plasma lipid levels and cholesterol kinetics by phytosterol versus phytostanol esters. *J Lipid Res* **41**, 697–705.
- Sudhop T, Sahin Y, Lindenthal B, Hahn C, Luers C, Berthold HK & von Bergmann K (2002) Comparison of the hepatic clearances of campesterol, sitosterol, and cholesterol in healthy subjects suggests that efflux transporters controlling intestinal sterol absorption also regulate biliary secretion. *Gut* **51**, 860–863.
- Teunissen CE, De Vente J, von Bergmann K, Bosma H, van Boxel MP, De Bruijn C, Jolles J, Steinbusch HW & Lütjohann D (2003) Serum cholesterol, precursors and metabolites and cognitive performance in an aging population. *Neurobiol Aging* **24**, 147–155.
- Ketomaki A, Gylling H & Miettinen TA (2005) Non-cholesterol sterols in serum, lipoproteins, and red cells in statin-treated FH subjects off and on plant stanol and sterol ester spreads. *Clin Chim Acta* **353**, 75–86.
- Rall CJ, Hoeg JM, Gregg RE, Law SW, Monge JC, Meng MS, Zech LA & Brewer HB Jr (1988) Enhanced apolipoprotein

- E production with normal hepatic mRNA levels in the Watanabe heritable hyperlipidemic rabbit. *Arteriosclerosis* **8**, 804–809.
32. Thelen KM, Falkai P, Bayer TA & Lütjohann D (2006) Cholesterol synthesis rate in human hippocampus declines with aging. *Neurosci Lett* **403**, 15–19.
 33. Miettinen TA, Puska P, Gylling H, Vanhanen H & Vartiainen E (1995) Reduction of serum cholesterol with sitostanol-ester margarine in a mildly hypercholesterolemic population. *N Engl J Med* **333**, 1308–1312.
 34. Andersen JM, Turley SD & Dietschy JM (1982) Relative rates of sterol synthesis in the liver and various extrahepatic tissues of normal and cholesterol-fed rabbits. Relationship to plasma lipoprotein and tissue cholesterol levels. *Biochim Biophys Acta* **711**, 421–430.
 35. Ikeda I, Tanabe Y & Sugano M (1989) Effects of sitosterol and sitostanol on micellar solubility of cholesterol. *J Nutr Sci Vitaminol (Tokyo)* **35**, 361–369.
 36. Mel'nikov SM, Seijen ten Hoorn JW & Eijkelenboom AP (2004) Effect of phytosterols and phytostanols on the solubilization of cholesterol by dietary mixed micelles: an *in vitro* study. *Chem Phys Lipids* **127**, 121–141.
 37. Dietschy JM, Kita T, Suckling KE, Goldstein JL & Brown MS (1983) Cholesterol synthesis *in vivo* and *in vitro* in the WHHL rabbit, an animal with defective low density lipoprotein receptors. *J Lipid Res* **24**, 469–480.