



Universiteit
Leiden
The Netherlands

Risk factors of chronic kidney disease progression: Dutch cohort studies

Esmeijer, K.

Citation

Esmeijer, K. (2020, March 19). *Risk factors of chronic kidney disease progression: Dutch cohort studies*. Retrieved from <https://hdl.handle.net/1887/137184>

Version: Publisher's Version

License: [Licence agreement concerning inclusion of doctoral thesis in the Institutional Repository of the University of Leiden](#)

Downloaded from: <https://hdl.handle.net/1887/137184>

Note: To cite this publication please use the final published version (if applicable).

Cover Page



Universiteit Leiden



The handle <http://hdl.handle.net/1887/137184> holds various files of this Leiden University dissertation.

Author: Esmeijer, K.

Title: Risk factors of chronic kidney disease progression: Dutch cohort studies

Issue Date: 2020-03-19

Chapter 6 –

Effect of different types of statins on kidney function decline and proteinuria: a network meta-analysis

K. Esmeijer, Olaf M. Dekkers, Johan W. de Fijter, Friedo W. Dekker,
Ellen K. Hoogeveen

Sci Rep. 2019, 9: 16632

ABSTRACT

Background: Previous studies showed that statins reduce the progression of kidney function decline and proteinuria, but whether specific types of statins are more beneficial than others remains unclear. We performed a network meta-analysis of randomized controlled trials (RCT) to investigate which statin most effectively reduces kidney function decline and proteinuria.

Methods: We searched MEDLINE, Embase, Web of Science, and the Cochrane database until July 13, 2018, and included 43 RCTs (>110,000 patients). We performed a pairwise random-effects meta-analysis and a network meta-analysis according to a frequentist approach. We assessed network inconsistency, publication bias, and estimated for each statin the probability of being the best treatment.

Results: Considerable heterogeneity was present among the included studies. In pairwise meta-analyses, 1-year use of statins versus control reduced kidney function decline by 0.61 (95%-CI: 0.27; 0.95) mL/min/1.73m² and proteinuria with a standardized mean difference of -0.58 (-0.88; -0.29). The network meta-analysis for the separate endpoints showed broad confidence intervals due to the small number available RCTs for each individual comparison.

Conclusions: 1-year statin use versus control attenuated the progression of kidney function decline and proteinuria. Due to the imprecision of individual comparisons, results were inconclusive as to which statin performs best with regard to renal outcome.

INTRODUCTION

Chronic kidney disease (CKD) is an increasing global health burden owing to population ageing and unhealthier lifestyle.¹ Up to 11% of the European population aged 45y or older has CKD stage 3, defined as an estimated glomerular filtration rate (eGFR) below 60 mL/min/1.73m².² CKD is an independent risk factor for cardiovascular morbidity and mortality.³ Nowadays, the most important causes of CKD are cardiovascular disease, hypertension, diabetes, smoking, and hypercholesterolemia.^{4,5} Generally, patients with symptomatic cardiovascular disease are prescribed cholesterol-lowering medication for secondary cardiovascular prevention. The latest KDIGO guideline on lipid management in CKD, recommends treatment with a statin in all non-dialysis dependent CKD patients ≥ 50 years with an eGFR below 60 mL/min/1.73m² or with at least 30 mg/g albuminuria, independent of serum cholesterol levels, which is also stated by the 2016 ESC/EAS guidelines.^{6,7} Younger patients should use a statin in case of elevated cardiovascular risk, such as diabetes or coronary heart disease. Finally, statins should be continued, but not initiated, in patients on dialysis.⁶ Multiple meta-analyses studied the effect of statins on renal outcomes. Recently, a meta-analysis by Su *et al.* concluded that statin users vs nonusers have a slower rate of kidney function decline and less proteinuria.⁸

Targeted prevention of kidney function decline is important to improve life expectancy and quality of life. However, it remains unclear whether specific types of statins are more beneficial than others regarding slowing down kidney function decline and lowering proteinuria. Various statins have different characteristics in terms of half-life, structure, lipophilicity, and potency.⁹ We therefore performed a network meta-analysis of randomized controlled trials in adults that compare any statin with another statin or control treatment, to investigate which statin most effectively reduces kidney function decline or proteinuria. Network meta-analyses take into account both direct and indirect evidence of multiple comparisons in a treatment network, and provide information on which treatment performs best. These results may inform future guidelines about prevention of CKD and slowing down its progression.

METHODS

Systematic literature review

We performed a systematic review of the literature, searching MEDLINE, Embase, Web of Science, and the Cochrane Library, on July 13th, 2018. Eligible studies were randomized controlled trials (RCT) in adults (patients ≥ 18

years) with a follow-up duration of at least one year, that included at least 10 patients per trial arm, and reported on changes in eGFR and/or proteinuria. The intervention of interest was statin therapy, the comparator either another statin, no intervention, cholesterol lowering diet, or placebo. In the entire manuscript, control treatment refers to any non-statin intervention. Combination therapy of statin with ezetimibe was also considered. A detailed outline of the search strategy is provided in the Supplemental Data, Appendix. Titles and abstracts were screened and relevant articles were read in full by two reviewers (KE and EH). Conference abstracts were excluded. No language restrictions were imposed. Post-hoc analyses of RCTs were only included when outcomes according to the original randomization group could be derived. In case of duplicate publications, we selected the publication that reported the data of interest most completely. References of included studies were additionally screened for relevant RCTs. We reported the results according to the Preferred Reporting Items for Systematic Reviews and Meta-analyses (PRISMA) guidelines for network meta-analyses.¹⁰ The protocol for this meta-analysis was registered at PROSPERO: registration number CRD42018099613.¹¹

Outcome measures

The outcomes of interest were annual change of estimated glomerular filtration rate (eGFR) and proteinuria. Kidney function estimates calculated by the Cockcroft-Gault formula, the Modification of Diet in Renal Disease (MDRD) formula, or Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) equation were pooled. If change of kidney function or proteinuria was not reported, it was calculated by subtracting the baseline value from follow-up. The standard deviation (SD) of change was calculated using the SDs of eGFR or proteinuria at baseline and follow-up, according to the following formula:¹²

$$SD_{change} = \sqrt{SD_0^2 + SD_1^2 - (2 * Corr * SD_0 * SD_1)}$$

Where SD_0 and SD_1 represent the SD of baseline and follow-up, respectively, and $Corr$ represents a correlation coefficient, which describes the similarity between baseline and follow-up measurements. The correlation coefficient was derived from studies that reported both baseline and follow-up eGFR or proteinuria with an SD, and change in eGFR or proteinuria with SD, according to the following formula:¹²

$$Corr = \frac{SD_0^2 + SD_1^2 + SD_{change}^2}{2 * SD_0 * SD_1}$$

Based on data from three intervention studies investigating the effect of statins on kidney function, and data from the Alpha Omega Trial, we assumed a correlation coefficient between baseline and follow-up eGFR of 0.8.¹³⁻¹⁶ In the main analysis we compared change of eGFR or proteinuria after 12 months for statin users vs control treatment. If no data were reported on change in eGFR or proteinuria after one year, we used the available data to calculate an annual change assuming a linear decline in line with the results of a recent study.¹⁷

Data extraction and quality assessment

Data extraction was performed by two independent reviewers (KE and EH) who used a standard form. Discrepancies were resolved by discussion or by consulting a third reviewer (OD). We extracted the following data: study name, study year, trial acronym, duration, population type, treatment arms, sample size, mean age, sex (% males), diabetes (%), hypertension (%), mean systolic and diastolic blood pressure, use of renin-angiotensin system (RAS) blocking drugs (%), low-density lipoprotein (LDL) level at baseline and follow-up, baseline and follow-up eGFR, change in eGFR, baseline and follow-up proteinuria, and change in proteinuria. When the outcome of interest was not reported in a table or text, we extracted the exact numbers from figures.

The Cochrane Collaboration Risk of Bias tool was used to assess potential sources of bias: selection, performance, detection, attrition and reporting bias.¹⁸ We scored per included RCT each type of bias as follows: low, high, or unclear risk of bias. Risk of bias was scored high in case of broken randomization, absent blinding of participants, absence of allocation concealment, and in case of large number of missing outcome data, or exclusion of patients. Since the outcome of interest was based on laboratory measurements, we considered for all RCTs, including the open-label RCTs, the risk of bias “low” with regard to blinding of outcome assessment.

Statistical analysis

First, we performed a pairwise random-effects meta-analysis for the effect of statin vs control on eGFR and proteinuria decline. For eGFR decline we used the weighted mean difference (WMD) as measure for the pooled estimates. For proteinuria we estimated standardized mean differences (SMD) to account for different methods to express proteinuria: urinary albumin to creatinine ratio, urinary protein excretion, urinary albumin excretion, or log-transformed protein excretion. Statistical heterogeneity was assessed by the I^2 -statistic, which quantifies the variation across studies due to heterogeneity rather than chance.¹⁹ We used meta-regression to evaluate whether heterogeneity could be explained by age, sex, diabetes, blood pressure, baseline LDL, change in LDL,

or risk of bias. Finally, we assessed the presence of publication bias visually with a funnel plot and formally by the Egger's test.^{20,21} This rank-based method estimates the number and outcomes of missing unpublished studies, and adjusts the estimate after incorporating these theoretical studies.

Second, we performed a random-effects network meta-analysis, following a frequentist approach. In case multiple dosages were reported, we analyzed high and low statin dosages as separate treatments. We took as outcome the WMD of annual kidney function decline and change of proteinuria expressed as SMD. We checked for transitivity and consistency. Transitivity was judged clinically; consistency was judged formally.²² We tested for possible inconsistency globally using a χ^2 -test, and locally by calculating inconsistency factors for each comparison in closed loops. In case of minor inconsistencies, possible reasons for inconsistency were considered. Furthermore, we estimated for each statin, compared to control, the treatment effect with 95%-confidence intervals and prediction intervals. The prediction interval represents the expected range of true effects in similar (future) studies, and will be broader than the confidence interval in case of high heterogeneity.²³ Finally, for each statin, with or without ezetimibe, we calculated the surface under the cumulative ranking (SUCRA) line. We used the SUCRA to provide a hierarchic overview of treatments, and to give an impression of the most efficacious treatments.²⁴ The SUCRA takes into account for every treatment the cumulative probabilities of all possible rankings. If a treatment always ranks first, the SUCRA is 100% (or 1), and 0% (or 0) if it always ranks last.²⁵

We repeated the analyses excluding RCTs with a total sample size <100 patients or stratified by open-label (yes/no) or post-hoc (yes/no) status. Subgroup analyses were not considered if too few RCTs remained to form a network. All statistical analyses were performed using STATA Statistical Software version 14 (Statacorp, Texas, USA), and the *StataNMA* package.²⁶

RESULTS

Characteristics of included studies

After removing duplicate RCTs, 1303 titles and abstracts were screened for eligibility; 76 full publications were assessed. Finally, 43 RCTs comprising over 110,000 patients reported in 42 publications were included (Figure 1). Of these 42 publications, 40 were in English, one was Russian,²⁷ and one Japanese.²⁸ In total, 40 RCTs reported about the effect of statins on change of eGFR,^{13-15,27,29-63} of which 30 compared a statin to control, and 10 compared two or more statins with each other. The effect of statins on proteinuria was reported in 25

RCTs,^{13,14,28,29,32-34,36,39,45,46,48-54,57,60,62-65} of which 19 compared a statin to control intervention, and six compared two or more statins. Characteristics of included RCTs are shown in Table 1. The included RCTs investigated seven different statins with varying dosages, and in three RCTs a statin was combined with ezetimibe.^{40,46,48} Of all included RCTs, 11 comprised coronary heart disease patients, 11 comprised CKD patients, and 11 comprised diabetes mellitus type 2 patients. The mean age of the enrolled patients in most RCTs was over 50 years and about 66% were men. The unweighted mean (range) of baseline LDL-cholesterol from all individual RCTs was 3.7 (2.2-7.8) mmol/L, and statin compared to control treatment led to a mean (SD) 27% (9%) reduction of the serum LDL level. The majority of RCTs had a low risk of bias (Supplementary Figure S1). However, about a 44% of all RCTs was open-label and about 25% were post-hoc analyses.

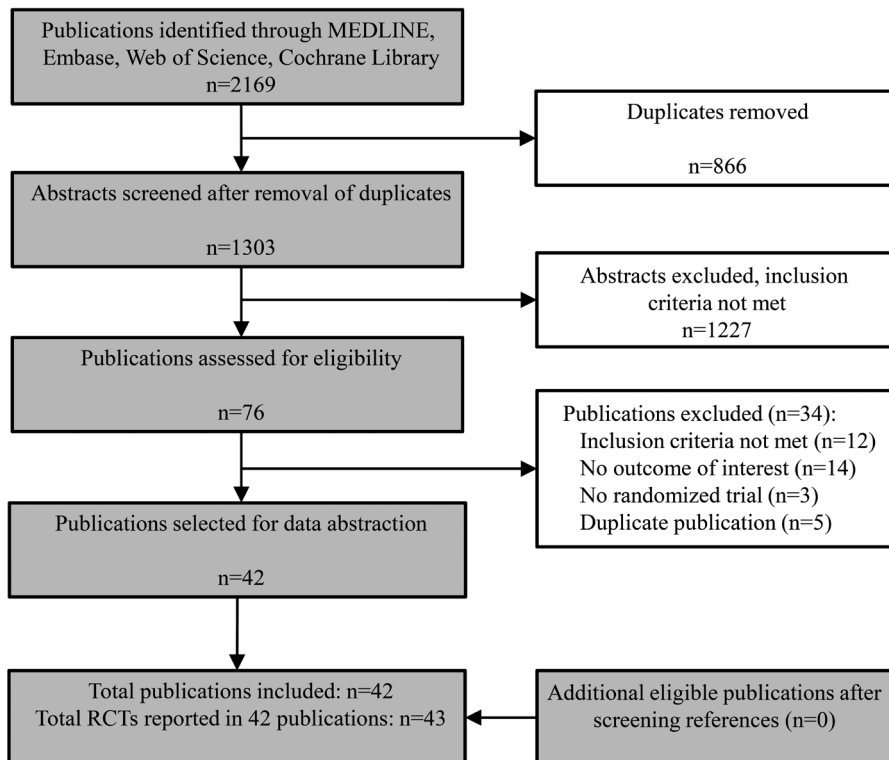


Figure 1: Flow chart of literature search and included full text publications. All included publications were included in quantitative analyses, depending on the reported endpoint(s).

Table 1: characteristics of included studies.

Author, year Study name	Population	Intervention	Sample size (n)	Follow-up Age (y)	Male sex (%)	Mean baseline characteristics per RCT					Outcome [annual change (SD)] per intervention		
						Diabetes (%)	Blood pressure (mmHg)	eGFR (mL/min/1.73m ²)	LDL (mmol/L)	eGFR	Proteinuria (measure)		
Abe, 2015	CKD	Rosuvastatin 2.5 mg Pitavastatin mean1.4 mg	134	1	70	58	44	.	58	3.6	2.80 (12.1)	-392 (802)	-250 (707) ^a (UACR)
Amarenco,2014 SPARCL	Stroke, TIA	Atorvastatin 80 mg Placebo	4719	5	63	60	17	139/82	66	3.4	0.96 (13.1)	.	-0.50 (13.1) ^b
Athyros, 2004 GREACE	CHD	Atorvastatin mean24 mg Control	1600	4	58	78	20	123/75	77	.	2.00 (2.0)	.	-0.75 (1.8) ^a
Atthobari, 2006 PREVEND-IT	General population	Pravastatin 40 mg Placebo	788	4	52	66	3	131/76	76	4.1	0.15 (3.7)	-0.02 (0.07)	-0.03 (0.08) ^c (UAE)
Bianchi, 2003	CKD	Atorvastatin 40 mg Placebo	56	1	56	47	0	133/85	50	5.1	-1.00 (5.9)	-1.0 (0.47)	0.3 (0.47) ^b (UPE)
Castelao, 1993	Transplant	Lovastatin 20 mg Simvastatin 10 mg	51	1	44	69	.	.	52	4.9	-1.00 (16.6)	0.38 (1.9)	0.31 (1.1) (UPE)
Colhoun, 2009 CARDS	DM2	Atorvastatin 20 mg Placebo	2838	4	62	68	100	144/83	64	3.0	-4.60 (15.3)	0.48 (2.7)	.
Dalla Nora, 2003	DM2	Atorvastatin 10 mg Placebo	25	1	65	60	100	.	.	3.5	0.30 (2.6)	.	2.0 (1.9)
Deedwania,2015 SAGE	CAD	Atorvastatin 80 mg Pravastatin 40 mg	868	1	72	69	23	.	62	3.8	2.38 (10.4)	.	6.0 (1.9) ^d (AER)
Fassett, 2010	ADPKD	Pravastatin20 mg Control	60	2	51	39	.	133/86	55	3.3	-0.31 (10.4)	-0.04 (0.20)	0.01 (0.09) (UPE)
Fassett, 2010 LORD	CKD	Atorvastatin 10 mg Placebo	132	3	60	65	8	143/81	31	3.4	-1.04 (3.84)	-0.39 (0.71)	-0.14 (0.85) (UPE)
Fellstrom, 2004 ALERT	Transplant	Fluvastatin 40 mg Placebo	439	5	50	66	19	144/86	52	4.1	-0.93 (8.9)	.	-1.87 (8.3) ^a
Fried, 2001	DM1	Simvastatin 10 mg Placebo	39	1.5	32	56	100	.	.	3.3	.	0.09 (0.44)	0.14 (0.66) ^d (AER)

Table 1: Continued

Author, year Study name	Population	Intervention	Sample size (n)	Follow-up time (y)	Age (y)	Male sex (%)	Mean baseline characteristics per RCT				Outcome [annual change (SD)] per intervention	
							Diabetes (%)	Blood pressure (mmHg)	eGFR (mL/min/1.73m ²)	LDL (mmol/L)	eGFR	Proteinuria (measure)
Gheith, 2002	Nephrotic syndrome	Fluvastatin 20 mg Control	43	1	23	42	.	.	107	7.8	-4.80 (28.8)	-6.0 (2.3)
Haynes, 2014 SHARP	CKD	Simvastatin 20 mg/eze Placebo	5037	4	63	62	23	139/80	27	2.9	-1.66 (3.5)	-2.0 (2.4) ^{de} (UPE)
Holme, 2010 IDEAL	MI	Atorvastatin 80 mg Simvastatin 20 mg	8888	4.8	62	81	12	137/80	68	3.1	-1.83 (3.5)	.
HPS, 2003 HPS	DM	Simvastatin 40 mg Placebo	20536	4.8	64	76	29	144/81	.	3.3	0.01 (2.7)	.
Huskey, 2009 4S	CHD	Simvastatin 20 mg Placebo	3842	5.5	58	80	4	139/83	76	4.9	0.34 (2.7)	-1.23 (1.86)
Kendrick, 2010 AFCAPS/Tex ASUCA	Primary prevention	Lovastatin 20 mg Placebo	4994	5.3	58	85	2	138/78	87	3.8	-1.40 (1.83)	.
Kimura, 2017	CKD	Atorvastatin 5–20 mg Control	334	2	63	64	34	133/77	55	3.7	-0.34 (7.4)	-1.30 (3.5)
Kimura, 2012	DM2	Pitavastatin 2 mg Pravastatin 10 mg	83	1	65	57	100	132/76	74	3.4	-0.41 (7.4) ^f	-1.40 (3.5)
Kinouchi, 2013	Dyslipidemia	Fluvastatin 20 mg Fluvastatin 20 mg/eze	54	1	54	67	6	140/90	71	4.1	-1.15 (4.4)	0.3 (1.3)
Koren, 2009 ALLIANCE	CHD	Atorvastatin mean 41mg Control	2442	4.5	61	82	22	134/79	73	3.8	-1.30 (4.4)	-0.2 (1.3) log(UAE)
Kouvelos, 2015	Vascular surgery	Rosuvastatin 10 mg Rosuvastatin 10 mg/eze	262	1	71	90	30	.	65	3.8	-2.0 (9.0)	-50 (150)
Lam, 1995	NID-DM	Lovastatin 20–40 mg Placebo	34	2	56	56	100	.	84	4.2	-0.5 (9.5) ^b	25 (175) ^b (UACR)
Lee, 2005	Controlled HT	Pravastatin 10 mg Placebo	61	1	49	68	0	121/73	87	3.2	-4.10 (7.7)	22.5 (72.4)
Lemos, 2013	CKD	Rosuvastatin 10 mg Control	77	2	58	61	21	.	40	3.1	4.10 (6.4)	-44.7 (74.5) ^d (UAE)
											-0.18 (6.4)	.
											-0.30 (7.2)	.
											-7.60 (10.1)	0.9 (2.0)
											-6.80 (10.7) ^a	0.5 (1.9) ^{ad} (UPE)
											-1.10 (5.7)	0 (0.1)
											-1.30 (3.6) ^{ab}	0.25 (0.2) ^{ab} (UPE)
											13.0 (13.3)	-673 (44.8)
											4.0 (12.4) ^a	-7 (327) ^{ab} (UPE)
											-1.15 (6.0)	0.08 (0.18)
											-2.50 (5.1) ^a	0.23 (0.26) ^{ad} (UPE)

Table 1: Continued

Author, year Study name	Population	Intervention	Sample size (n)	Follow-up (y)	Age (y)	Male sex (%)	Mean baseline characteristics per RCT				Outcome [annual change (SD)] per intervention	
							Diabetes (%)	Blood pressure (mmHg)	eGFR (mL/min/1.73m ²)	LDL (mmol/L)	eGFR	Proteinuria (measure)
Mori, 1992	NID-DM	Pravastatin 10 mg Control	33	1	63	36	100	134/80	.	2.9	.	-5.4 (71.8) ^a (UACR)
Mou, 2016	Chronic glom. nephritis	Pravastatin 20 mg Control	48	1.8	51	.	8	133/75	75	3.5	-1.08 (12.7) -4.33 (10.6) ^{ab}	-0.33 (0.9) -0.27 (0.9) ^{ab} (UPE)
Nanayakkara, 2007 ATIC	CKD	Pravastatin 40 mg* Placebo	87	2	53	57	0	135/79	34	3.6	0 (4.3) 0.15 (4.3) ^{ab}	-0.1 (0.8) 0.2 (0.8) ^b log(UAE)
Ohsawa, 2015	CKD	Pitavastatin 1-4 mg Control	28	1	62	71	33	130/78	49	3.6	-3.50 (3.21) -4.20 (2.96) ^a	-24.4 (574) -338 (1141) ^a (UACR)
Rahman, 2008 ALLHAT	HT, HCh	Pravastatin 40 mg Control	10355	6	67	51	35	143/83	78	3.8	-1.45 (5.9) -1.65 (5.9) ^a	.
Rutter, 2011 PANDA	DM2	Atorvastatin 80 mg Atorvastatin 10 mg	119	2.5	64	83	100	.	67	3.1	1.0 (13.8) -3.0 (11.8) ^{ab}	.
Sawara, 2008	CKD	Rosuvastatin 2.5 mg Control	38	1	65	0	.	127/78	53	3.3	2.60 (12.3) -2.20 (10.6) ^a	-0.04 (0.19) 0.05 (0.24) ^a (UPE)
Scanferla, 1991	CKD	Sim/pravastatin 10 mg Control	24	1	54	58	.	172/106	40	.	-1.80 (4.2) -3.10 (4.2)	.
Shepherd, 2007 TNT	CAD	Atorvastatin 80 mg Atorvastatin 10 mg	10001	5	61	81	15	131/78	65	2.5	1.5 (9.7) 0.1 (9.7) ^{bc}	.
Takazakura, 2015	DM	Atorvastatin 10 mg Pravastatin 10 mg Control	106	1	62	87	100	129/0	64	3.0	-0.80 (11.4) -2.80 (10.8) -3.10 (9.6) ^a	-0.2 (0.4) -0.1 (0.7) 0.1 (0.5) ^a log(UACR)
Tonelli, 2005 PPP **	CAD	Pravastatin 40 mg Placebo	18569	5	58	90	7	133/81	73	4.2	Effect of pravastatin: 0.10 (0.02; 0.17) mL/ min/1.73m ² *	.

Table 1: Continued

Author, year Study name	Population	Intervention	Sample size (n)	Follow-up time (y)	Age (y)	Male sex (%)	Mean baseline characteristics per RCT				Outcome [annual change (SD)] per intervention	
							Diabetes (%)	Blood pressure (mmHg)	eGFR (mL/min/1.73m ²)	LDL (mmol/L)	eGFR	Proteinuria (measure)
Vidt, 2011 JUPITER	Healthy population	Rosuvastatin 20 mg Placebo	16279	2.3	66	62	31	.	75	.	-7.10 (11.9)	.
Yakusevich, 2013	Stroke	Simvastatin 40 mg Control	210	1	66	45	.	.	76	2.2	7.05 (12.1)	.
Yasuda, 2004	CKD	Fluvastatin 20 mg Control	80	0.9	58	46	43	144/80	60	4.4	-8.67 (3.9) -6.50 (4.0) ^a	0 (0.14) 0 (0.15) ^a (UAE)
De Zeeuw, 2015 PLANET I	DM	Rosuvastatin 10 mg Rosuvastatin 40 mg Atorvastatin 80 mg	325	1	58	70	100	139/79	71	3.9	-3.70 (14.7) -7.29 (20.4) -1.61 (13.0)	2 (79) -4 (77) -13 (57) %change
De Zeeuw, 2015 PLANET II	Non-DM proteinuria	Rosuvastatin 10 mg Rosuvastatin 40 mg Atorvastatin 80 mg	220	1	49	62	0	130/81	75	4.3	-2.71 (13.3) -3.30 (12.5) -1.74 (14.2)	-6 (99) 8 (75) -24 (60)

ACS, acute coronary syndrome; ADPKD, autosomal dominant polycystic kidney disease; CAD, coronary artery disease; CHD, coronary heart disease; CKD, chronic kidney disease; eGFR, estimated glomerular filtration rate; eze, ezetimibe; HT, hypertension; MI, myocardial infarction; TIA, transient ischemic attack; (NID-)DM1/DM2, non-insulin dependent diabetes mellitus 1 or 2, LDL, low-density lipoprotein; prot, proteinuria; UACR, urinary albumin-to-creatinine ratio; UAE, urinary albumin excretion; UPE, urinary protein excretion.

*Intervention was a combination of statin and vitamin E supplementation.

**PPP: Pravastatin Pooling Project, study representing pooled estimates of three RCTs: LIPID, CARE, and WOSCOPS. Individual data on each RCT was not published.

a: based on eGFR (SD) value at baseline and follow-up. SD of eGFR change was calculated according to the formula provided in the Cochrane Handbook.¹¹ b: data extracted from figure.

c: reported geometric mean was log-transformed to achieve normal distribution with symmetrical SD.

d: SD acquired by dividing interquartile range by 1.35.

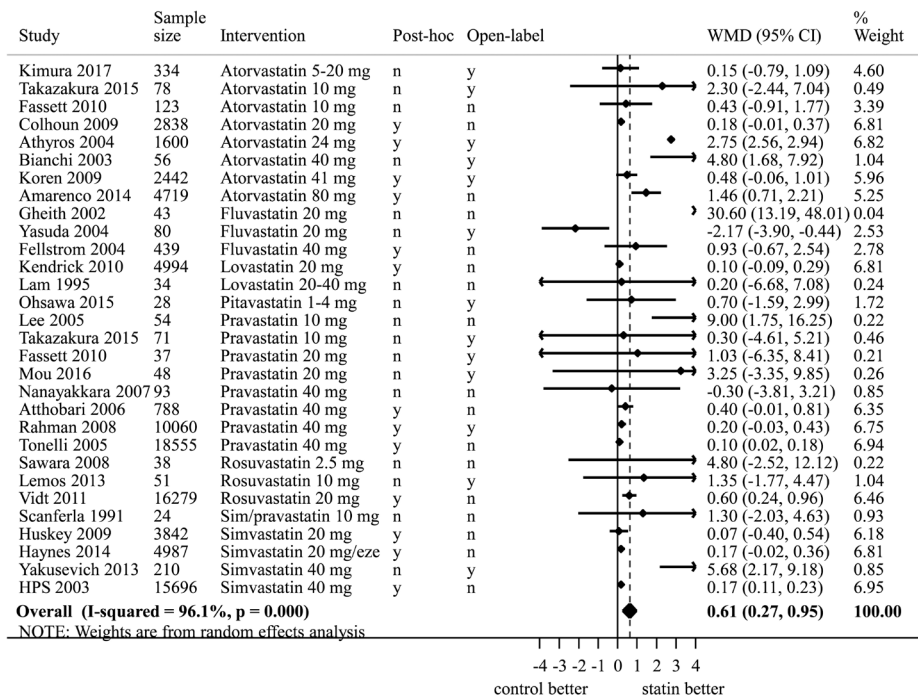
e: no SD or SE reported, these were therefore borrowed from comparable studies.

f: SD of baseline eGFR value used to calculate SD of eGFR change.

g: only effect of treatment vs control reported.

Pairwise comparison: statins and eGFR decline

Except for two medium sized trials (Yasuda *et al.*, and Nanayakkara *et al.*), effect estimates of all RCTs showed a protective effect of statin on eGFR decline.^{53,62} Random-effects meta-analysis showed that statin use, compared to control, led to a 0.61 (95% CI 0.27; 0.95) mL/min/1.73m² slower annual eGFR decline (Figure 2). When only RCTs with a sample size of at least 100 patients (n=16) were analyzed, the beneficial effect of statin treatment on annual eGFR decline was 0.58 (0.23; 0.92) mL/min/1.73m². Heterogeneity between RCTs was high, with an I² of 96%. Meta-regression showed that higher systolic blood pressure at baseline was significantly associated with smaller effects of statins, explaining 40% of the between-study variance. We found no evidence for interaction between diabetes and statins with regard to the beneficial effect on kidney function decline. Age, sex, serum LDL level, or change in LDL, had no significant impact on the effect estimates. In post-hoc RCTs (n=11) the beneficial effect on annual kidney function decline of statins vs control was smaller but more precise than in RCTs in which change in eGFR was the primary outcome (n=17): 0.55 (0.19; 0.92) vs 1.55 (0.26; 2.85) mL/min/1.73m², respectively. In open-label RCTs (n=17, mean sample size 4326) the beneficial effect on eGFR decline of statins vs control was stronger than in blinded RCTs (n=13, mean sample size 1161): 1.25 (0.08; 2.42) vs 0.23 (0.11; 0.34) mL/min/1.73m², respectively. The funnel plot for eGFR decline was slightly asymmetrical (Supplementary Figure S2), but the Egger's test for small study effects was not significant (p= 0.3).



Change in annual eGFR decline, mL/min/1.73m²

Figure 2: Pairwise random effects meta-analysis of randomized controlled trials investigating the effect of statin therapy versus control on the rate of annual eGFR decline. Positive values mean slower eGFR decline for statin users vs non-users, thus favouring statin use. eGFR, estimated glomerular filtration rate; eze, ezetimibe 10 mg; WMD, weighted mean difference.

Pairwise comparison: statins and proteinuria

The two largest RCTs showed that statin treatment vs control did not lower proteinuria: SMD of 0.40 (0.18; 0.61) and 0.18 (0.04; 0.32), respectively.^{32,63} In a meta-analysis, statin use compared to control showed a significant reduction of proteinuria with an SMD -0.58 (-0.88; -0.29) (Figure 3). However, the funnel plot of the effect of statins on proteinuria suggested publication bias (Supplementary Figure S3) and the Egger's test was significant ($p < 0.001$).

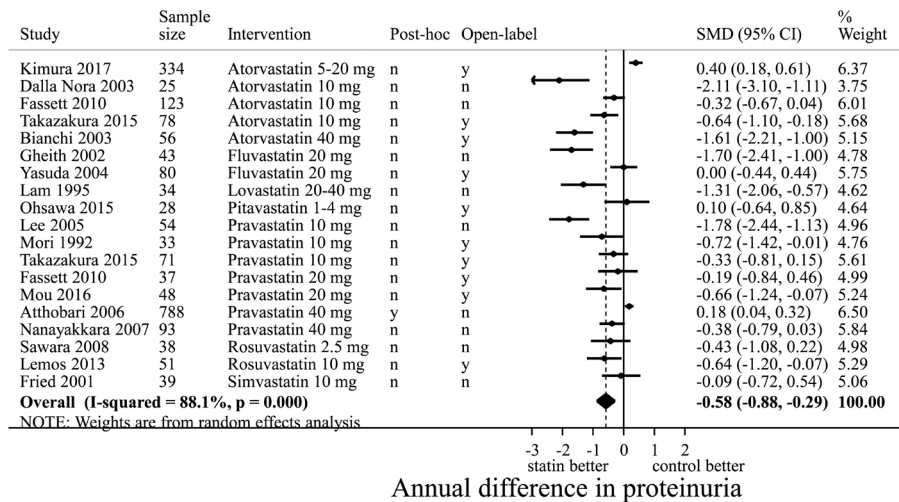


Figure 3: Pairwise random effects meta-analysis of randomized controlled trials investigating the effect of statin therapy versus control on the rate of annual change in proteinuria. Negative values mean a decrease in proteinuria for statin users vs non-users, thus favouring statin use. Effects expressed as SMD (standardized mean difference).

Network meta-analysis

Figure 4 (upper panel) shows the network plot of different statin treatments for change in eGFR. Each connection was formed by maximally 4 RCTs. We found no evidence for inconsistency in the network for eGFR decline and proteinuria using global tests (p-value for inconsistency 0.8) or local tests (p > 0.3 for all loops). We found that almost all statins performed better than control (Figure 5). The most beneficial effect on eGFR decline was caused by fluvastatin 20 mg/ezetimibe 10 mg, rosuvastatin 20 mg/ezetimibe 10 mg, pravastatin 10-20 mg, and atorvastatin 40-80 and 10<40 mg. However, point estimates had broad 95%-confidence intervals and prediction intervals. Except for combined fluvastatin 20 mg/ezetimibe 10 mg and atorvastatin 40-80 mg, all 95%-confidence intervals crossed the line of no effect.

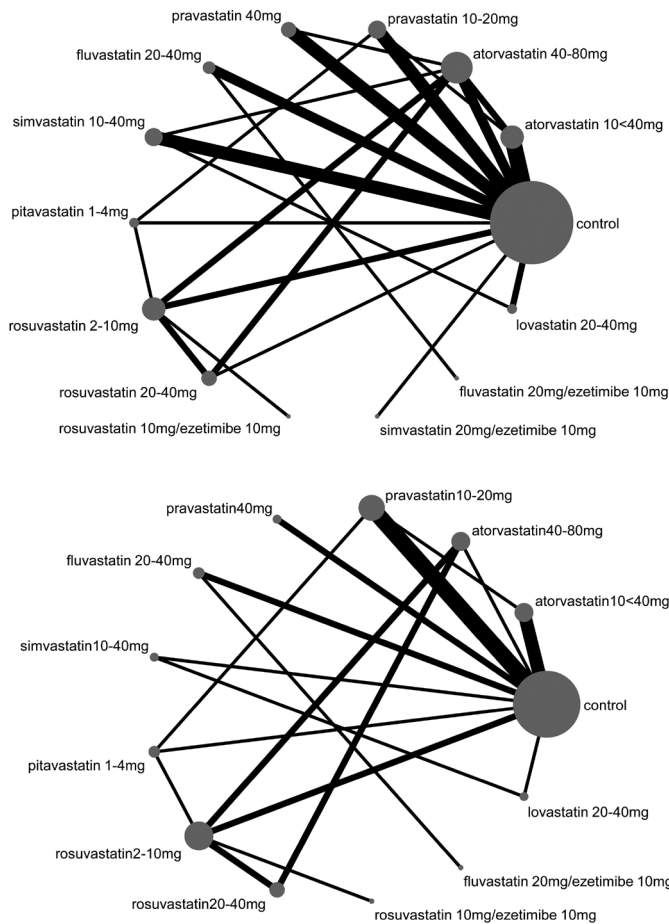


Figure 4: Network plots for outcome eGFR decline (upper panel) and proteinuria (lower panel). The width of the interconnecting lines is proportional to the number of RCTs providing evidence (ranging from 1 to 4). The size of the nodes is proportional to the total number of patients. eGFR, estimated glomerular filtration rate; RCT, randomized controlled trial.

Figure 4 (lower panel) shows the network plot for all statin treatments regarding proteinuria. For proteinuria, no single RCT compared the combination therapy simvastatin/ezetimibe. Globally, there was no evidence for inconsistency (p -value 0.8). However, using local tests, there were 2 inconsistent loops: control, atorvastatin 40–80 mg, rosuvastatin 2–10 mg ($p=0.04$) and control, simvastatin 10–40 mg, lovastatin 20–40 mg ($p=0.03$). The inconsistencies between direct and indirect effects were introduced by the relatively large effect estimates of small studies ($n < 60$). The most efficacious treatments regarding

proteinuria were fluvastatin 20 mg/ezetimibe 10 mg, atorvastatin 40–80 mg, and rosuvastatin 20 mg/ezetimibe 10 mg (Figure 6).

Finally, SUCRA analysis showed that control treatment had the lowest SUCRA. Fluvastatin 20 mg/ezetimibe 10 mg had the highest SUCRA value for eGFR decline (99%) and fluvastatin 20 mg/ezetimibe 10 mg (86%) as well as atorvastatin 40–80 mg (78%) had the highest SUCRA value for change in proteinuria (Figure 7).

Reduction of annual eGFR decline for different statins compared to control

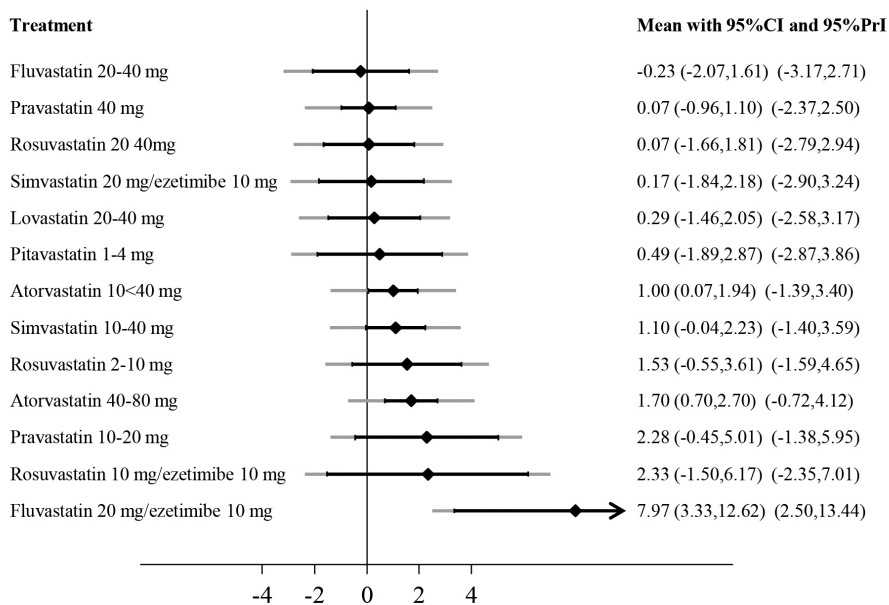


Figure 5: Effect of different statins compared to control treatment, on annual eGFR decline. Effects are presented as weighted mean differences (WMD). Positive values mean a slower eGFR decline. Black lines around point estimates reflect 95%-confidence intervals, grey lines reflect prediction intervals. Prediction intervals represent the expected range of true effects in (future) similar studies and is suitable to assess the variability of effect across different settings. CI, confidence interval; eGFR, estimated glomerular filtration rate; PrI, prediction interval.

Annual change in proteinuria for different statins compared to control

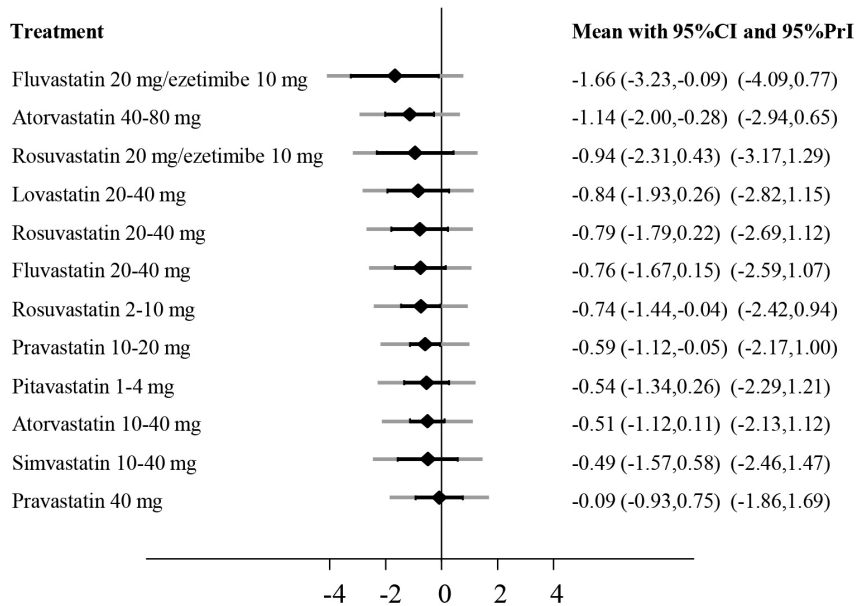


Figure 6: Effect of different statins compared to control treatment, on annual change in proteinuria. Effects are presented as standardized mean differences (SMD). Negative values mean a reduction of proteinuria. Black lines around point estimates reflect 95%-confidence intervals, grey lines reflect prediction intervals. Prediction intervals represent the expected range of true effects in (future) similar studies and is suitable to assess the variability of effect across different settings.
CI, confidence interval; eGFR, estimated glomerular filtration rate; PrI, prediction interval.

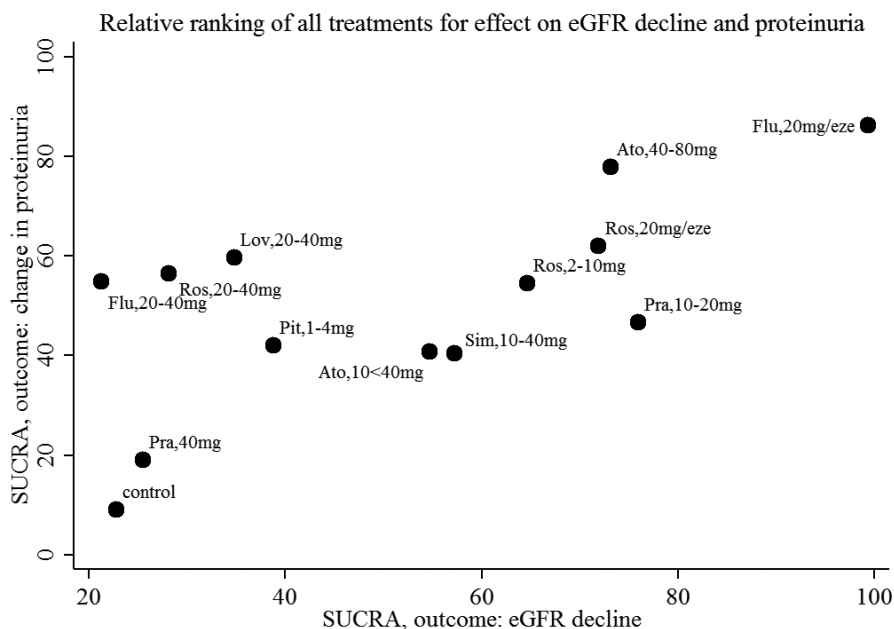


Figure 7: SUCRA analyses. Each dot represents the SUCRA value of each treatment. The SUCRA takes into account for every treatment the cumulative probabilities of all possible rankings. If a treatment always ranks first or last, the SUCRA is 100% or 0%, respectively. The horizontal axis shows SUCRA values with regards to the outcome eGFR decline, the vertical axis shows the SUCRA for the outcome proteinuria.

Ato, atorvastatin; eze, ezetimibe 10 mg; Flu, fluvastatin; Lov, lovastatin; Pit, pitavastatin; Pra, pravastatin; Ros, rosuvastatin; Sim, simvastatin; eGFR, estimated glomerular filtration rate; SUCRA, surface under the cumulative ranking curve.

Sensitivity analyses

Since we included RCTs with seven different types of statin treatments with one or more different dosages, networks of subgroups had only few closed loops. Therefore, estimates were based mostly on either direct or indirect evidence, but not on mixed evidence. Nonetheless, we repeated the network meta-analysis for eGFR decline excluding RCTs with a sample size <100 (n=16), excluding open-label RCTs (n=17), or excluding post-hoc analyses (n=20). Although effect estimates and rankings of individual treatments were variable across the analyses, in general atorvastatin 40–80 mg, fluvastatin 20 mg/ezetimibe 10 mg, pravastatin 10–20 mg, simvastatin 10–40 mg, and fluvastatin 20 mg were the most effective treatments with regard to eGFR decline. However, 95%-confidence intervals had substantial overlap, and individual treatments were rarely statistically significantly different from control. Since only a

small number of RCTs with small sample sizes studied the effect of statins on proteinuria, we could not perform the aforementioned sensitivity analyses.

DISCUSSION

In this network meta-analysis, we showed that there are no substantial differences in the efficacy of seven different statins and dosages, with or without ezetimibe, regarding slowing down eGFR decline or reducing proteinuria. If anything, the combination of fluvastatin 20 mg/ezetimibe 10 mg and atorvastatin 40–80 mg most consistently had the strongest beneficial effect on both renal endpoints, but the differences between treatments were small and confidence intervals were wide. In the pairwise meta-analysis we showed that use of statins lowered the rate of annual kidney function decline by 0.61 mL/min/1.73m² and reduced the amount of proteinuria by -0.58 (-0.88; -0.29) standard deviations per year.

Our results are in line with a recent meta-analysis Su *et al.* which reported that statins compared to control led to a 0.41 (0.11; 0.70) mL/min/1.73m² slower annual eGFR decline and a reduction of -0.65 (-0.94; -0.37) standard deviations in proteinuria.⁸ The small difference in outcomes between the present study and Su *et al.* are explained by different inclusion criteria. In contrast to the study of Su *et al.*, we included three RCTs investigating combinations of statins plus ezetimibe. Including also treatments combining statins with ezetimibe, results in a more complete review of existing literature on lipid-lowering therapy by statins. As a consequence we incorporated in our meta-analysis three extra RCTs, including the SHARP trial (n=5037). Furthermore, we excluded RCTs with a short follow-up (<12 months) or less than 10 patients per study arm, of which Su *et al.* included 19 RCTs. Finally, we found that the beneficial effect of statins on eGFR decline was weaker in RCTs with a higher mean systolic blood pressure. Systolic blood pressure explained 40% of the between-study variance. Taken together, these results suggest that a high systolic blood pressure modifies the effect of statins on eGFR decline. Hypertension is most likely a stronger risk factor for kidney function decline compared to hypercholesterolemia. Therefore, we speculate that the positive effect of statins on kidney function decline is overwhelmed in the presence of high blood pressure.

In our network meta-analysis, we specifically investigated the efficacy of individual statins and different dosages, using both direct and indirect evidence. We showed that each different statin compared to placebo had a beneficial effect on the annual eGFR decline and reduced proteinuria. However, confidence intervals were broad for individual treatment comparisons in our

network, due to the small number of RCTs contributing to each comparison. Su *et al.* showed in subgroup analyses the strongest beneficial effect on change in eGFR decline for atorvastatin, fluvastatin, and rosuvastatin.⁸ However, they pooled for each statin all dosages. The validity of these comparisons may be limited, considering the clear differential effects of different dosages.^{8,66}

We showed that fluvastatin 20 mg/ezetimibe 10 mg was the most efficacious treatment regarding both renal outcomes. However, this result was strongly influenced by the study of Kinouchi *et al.*, comprising 54 patients, reporting an annual eGFR decline of $-4.1 \text{ mL/min/1.73m}^2$ in patients treated with fluvastatin 20 mg compared to an annual eGFR increase of $4.1 \text{ mL/min/1.73m}^2$ in patients treated with fluvastatin 20 mg/ezetimibe 10 mg.⁴⁶ Since the average annual eGFR decline in adults with a history of cardiovascular disease is about 2 mL/min/1.73m^2 , the reported effect of Kinouchi *et al.* of $8.2 \text{ mL/min/1.73m}^2$ is large, and should be interpreted with caution.⁶⁷ We found that the second most efficacious statin on both renal endpoints was high dose atorvastatin, which improved the annual eGFR decline by 1.70 (0.70; 2.70) mL/min/1.73m^2 and reduced proteinuria by 1.14 (0.28; 2.00) standard deviations, compared to control.

Statins included in the present study reduced LDL levels on average by 27%, which is in line with a previous meta-analysis showing an LDL-lowering effect for all statins.⁶⁶ However, there is no clear evidence that high LDL itself increases CKD risk.⁶⁸ Statins also may have pleiotropic effects favourable for reducing CKD progression, such as lowering oxidative stress, reducing inflammation, and stabilizing atherosclerotic plaques.^{7,69} Hence, current guidelines recommend a statin for patients at risk for CKD, independent of LDL levels.^{9,70}

The main strength of the current study is that we performed a network meta-analysis, in addition to a pairwise meta-analysis, to investigate differential effects of different statins with or without ezetimibe. We only included RCTs because they are more likely to provide unbiased information. We excluded small trials (<10 patients per arm) since they are more susceptible to publication bias.

This network meta-analysis has several limitations. First, heterogeneity was high ($I^2 = 96\%$) owing to variation of the included patient populations across RCTs, differences in blinding methods, randomization procedures, sample size, and variability in primary endpoints. The I^2 statistic represents statistical heterogeneity, rather than clinically relevant heterogeneity, and is most strongly affected by the sample size of the individual studies. Upon increasing precision (sample size) of studies within a meta-analysis, the I^2 statistic rapidly approaches 100%.⁷¹ Deciding whether it is valid to pool studies, should be based on the clinical relevance of any present heterogeneity,

rather than solely on the I^2 statistic.⁷¹ We used random effects models to take heterogeneity into account. Second, we found an asymmetric funnel plot regarding proteinuria, which may be an indication of publication bias. On the other hand, larger compared to smaller RCTs showed a weak but opposite effect. Thus, the asymmetry may also be the consequence of inclusion of smaller RCTs with lower quality. Therefore, we cannot rule out that the beneficial effect of statins on proteinuria is an overestimation. Additionally, there were relatively few RCTs investigating the effect of statins on proteinuria, and most of them were small (sample size <100). Small studies therefore had a large impact on the network meta-analysis estimates, introducing inconsistencies especially in loops comprising small numbers of RCTs. The advantage of a network analysis is that it takes both direct and indirect effects into account, reducing the impact of single studies with a small sample size. For the outcome eGFR decline, the sample sizes of the included RCTs were large (24 RCTs with $n > 100$) which improved precision and reduced potential publication bias. The much smaller effect of statins compared to control in double blind compared to open-label RCTs may suggest bias due to the lack of blinding in the open-label RCTs. Since 17 out of 30 RCTs were open-label, we may have overestimated the beneficial effect on eGFR decline of statins compared to control. Third, due to the low number of RCTs contributing to each connection in the network meta-analyses, there was insufficient power to detect differences between statins. Fourth, a large number of the included RCTs used the MDRD formula to estimate eGFR, which is known to underestimate the true eGFR for values reported higher than 60 mL/min/1.73m².⁷² If anything, this may have underestimated the beneficial effect of statin use compared to control in studies with a mean eGFR higher than 60 mL/min/1.73m².

In conclusion, we found a beneficial effect of different statins, with or without ezetimibe, compared to control on progression of eGFR decline, and possibly proteinuria. Due to the imprecision of individual comparisons, results were inconclusive as to which statin performs best with regard to renal outcome.

ACKNOWLEDGEMENTS

The authors would like to thank Jan Schoones (Waleaus Library, Leiden University Medical Center) for his indispensable help with the literature search. We also thank Katerina Papadimitropoulou, for advice regarding network meta-analysis methodology.

DISCLOSURES

The authors have no disclosures to report.

FUNDING

The current study received no external funding.

AUTHORS' CONTRIBUTIONS

Research idea and study design: KE, OD, EH; data acquisition: KE, EH; data analysis/interpretation: EH, OD, KE, JF, FD; statistical analysis: KE, OD; supervision and mentorship: JF, EH, OD, FD. Each author contributed important intellectual content during manuscript drafting or revision and accepts accountability for the overall work by ensuring that questions pertaining to the accuracy or integrity of any portion of the work are appropriately investigated and resolved.

REFERENCES

1. El Nahas AM, Bello AK. Chronic kidney disease: the global challenge. *Lancet*. 2005; 365: 331–340.
2. Brück K, Stel VS, Gambaro G, et al. CKD Prevalence Varies across the European General Population. *J Am Soc Nephrol*. 2015; 27: 2135–2147.
3. Hoogeveen EK, Geleijnse JM, Giltay EJ, et al. Kidney function and specific mortality in 60–80 years old post-myocardial infarction patients: A 10-year follow-up study. *PLoS One*. 2017; 12: e0171868.
4. Obermayr RP, Temml C, Knechtelsdorfer M, et al. Predictors of new-onset decline in kidney function in a general middle-european population. *Nephrol Dial Transplant*. 2008; 23: 1265–1273.
5. Hobeika L, Hunt KJ, Neely BA, et al. Comparison of the Rate of Renal Function Decline in NonProteinuric Patients With and Without Diabetes. *Am J Med Sci*. 2015; 350: 447–452.
6. Kidney Disease: Improving Global Outcomes (KDIGO) Lipid Work Group. KDIGO Clinical Practice Guideline for Lipid Management in Chronic Kidney Disease. *Kidney Int Suppl*. 2013; 3: 259–305.
7. Catapano AL, Graham I, De Backer G, et al. 2016 ESC/EAS Guidelines for the Management of Dyslipidaemias: The Task Force for the Management of Dyslipidaemias of the European Society of Cardiology (ESC) and European Atherosclerosis Society (EAS). *Atherosclerosis*. 2016; 253: 281–344.
8. Su X, Zhang L, Lv J, et al. Effect of Statins on Kidney Disease Outcomes: A Systematic Review and Meta-analysis. *Am J Kidney Dis*. 2016; 67: 881–892.
9. Oesterle A, Laufs U, Liao JK. Pleiotropic Effects of Statins on the Cardiovascular System. *Circ Res*. 2017; 120: 229–243.
10. Hutton B, Salanti G, Caldwell DM, et al. The prisma extension statement for reporting of systematic reviews incorporating network meta-analyses of health care interventions: Checklist and explanations. *Ann Intern Med*. 2015; 162: 777–784.
11. Booth A. PROSPERO's progress and activities 2012/13. *Systematic Reviews*. 2013; 2: 111.
12. Higgins JPT, Green S. Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0 Part 3, chapter 16.13.2. The Cochrane Collaboration; 2011.
13. Fassett RG, Coombes JS, Packham D, et al. Effect of pravastatin on kidney function and urinary protein excretion in autosomal dominant polycystic kidney disease. *Scand J Urol Nephrol*. 2010; 44: 56–61.
14. Fassett RG, Robertson IK, Ball MJ, et al. Effect of atorvastatin on kidney function in chronic kidney disease: a randomised double-blind placebo-controlled trial. *Atherosclerosis*. 2010; 213: 218–224.

15. Vidt DG, Ridker PM, Monyak JT, et al. Longitudinal assessment of estimated glomerular filtration rate in apparently healthy adults: a post hoc analysis from the JUPITER study (justification for the use of statins in prevention: an intervention trial evaluating rosuvastatin). *Clin Ther*. 2011; 33: 717-725.
16. Hoogeveen EK, Geleijnse JM, Kromhout D, et al. Effect of omega-3 fatty acids on kidney function after myocardial infarction: the Alpha Omega Trial. *Clin J Am Soc Nephrol*. 2014; 9: 1676-1683.
17. Van den Brand JAJG, Dijkstra TMJ, Wetzels J, et al. Predicting kidney failure from longitudinal kidney function trajectory: A comparison of models. *PLoS One*. 2019; 14: e0216559.
18. Higgins JPT, Altman DG, Gøtzsche PC, et al. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials. *Br med J*. 2011; 343: d5928.
19. Higgins JP, Thompson SG, Deeks JJ, et al. Measuring inconsistency in meta-analyses. *Br med J*. 2003; 327: 557-560.
20. Egger M, Davey Smith G, Schneider M, et al. Bias in meta-analysis detected by a simple, graphical test. *Br med J*. 1997; 315: 629-634.
21. Noordzij M, Hooft L, Dekker FW, et al. Systematic reviews and meta-analyses: when they are useful and when to be careful. *Kidney Int*. 2009; 76: 1130-1136.
22. Rouse B, Chaimani A, Li T. Network meta-analysis: an introduction for clinicians. *Intern Emerg Med*. 2017; 12: 103-111.
23. In 't Hout J, Ioannidis JPA, Rovers MM, et al. Plea for routinely presenting prediction intervals in meta-analysis. *BMJ Open*. 2016; 6: e010247.
24. Mbuagbaw L, Rochwerg B, Jaeschke R, et al. Approaches to interpreting and choosing the best treatments in network meta-analyses. *Systematic Reviews*. 2017; 6: 79.
25. Salanti G, Ades AE, Ionnidis JPA. Graphical methods and numerical summaries for presenting results from multiple-treatment meta-analysis: an overview and tutorial. *J. Clin. Epidemiol*. 2011; 64: 163-171.
26. Chaimani A, Higgins JPT, Mavridis D, et al. Graphical Tools for Network Meta-Analysis in STATA. *PLoS One*. 2013; 8: e76654.
27. Yakusevich VV, Malygin AY, Kabanov AV. Effect of simvastatin on the prognosis and the changes of the clinical status in patients with acute ischemic stroke. The results of the 12 month randomized, open comparative study. [Russian]. *Rational Pharmacotherapy in Cardiology*. 2013; 9: 379-385.
28. Mori Y, Tsuruoka A. Effect of pravastatin on microalbuminuria in patients with non-insulin-dependent diabetes mellitus. *J Jpn Diabetes Soc*. 1992; 35: 265-268.
29. Abe M, Maruyama N, Maruyama T, et al. A Trial of Pitavastatin Versus Rosuvastatin for Dyslipidemia in Chronic Kidney Disease. *J Atheroscler Thromb*. 2015; 22: 1235-1247.
30. Amarenco P, Callahan A, III, Campese VM, et al. Effect of high-dose atorvastatin on renal function in subjects with stroke or transient ischemic attack in the SPARCL trial. *Stroke*. 2014; 45: 2974-2982.

31. Athyros VG, Mikhailidis DP, Papageorgiou AA, et al. The effect of statins versus untreated dyslipidaemia on renal function in patients with coronary heart disease. A subgroup analysis of the Greek atorvastatin and coronary heart disease evaluation (GREACE) study. *J Clin Pathol.* 2004; 57: 728-734.
32. Atthobari J, Brantsma AH, Gansevoort RT, et al. The effect of statins on urinary albumin excretion and glomerular filtration rate: results from both a randomized clinical trial and an observational cohort study. *Nephrol Dial Transplant.* 2006; 21: 3106-3114.
33. Bianchi S, Bigazzi R, Caiazza A, et al. A controlled, prospective study of the effects of atorvastatin on proteinuria and progression of kidney disease. *Am J Kidney Dis.* 2003; 41: 565-570.
34. Castela AM, Grino JM, Andres E, et al. HMGCoA reductase inhibitors lovastatin and simvastatin in the treatment of hypercholesterolemia after renal transplantation. *Transplant Proc.* 1993; 25: 1043-1046.
35. Colhoun HM, Betteridge DJ, Durrington PN, et al. Effects of atorvastatin on kidney outcomes and cardiovascular disease in patients with diabetes: an analysis from the Collaborative Atorvastatin Diabetes Study (CARDS). *Am J Kidney Dis.* 2009; 54: 810-819.
36. De Zeeuw D, Anzalone DA, Cain VA, et al. Renal effects of atorvastatin and rosuvastatin in patients with diabetes who have progressive renal disease (PLANET I): a randomised clinical trial. *Lancet Diabetes Endocrinol.* 2015; 3: 181-190.
37. Deedwania PC, Stone PH, Fayyad RS, et al. Improvement in Renal Function and Reduction in Serum Uric Acid with Intensive Statin Therapy in Older Patients: A Post Hoc Analysis of the SAGE Trial. *Drugs Aging.* 2015; 32: 1055-1065.
38. Fellstrom B, Holdaas H, Jardine AG, et al. Effect of fluvastatin on renal end points in the Assessment of Lescol in Renal Transplant (ALERT) trial. *Kidney Int.* 2004; 66: 1549-1555.
39. Gheith OA, Sobh MA, Mohamed K, et al. Impact of treatment of dyslipidemia on renal function, fat deposits and scarring in patients with persistent nephrotic syndrome. *Nephron.* 2002; 91: 612-619.
40. Haynes R, Lewis D, Emberson J, et al. Effects of lowering LDL cholesterol on progression of kidney disease. *J Am Soc Nephrol.* 2014; 25: 1825-1833.
41. Heart Protection Study. MRC/BHF Heart Protection Study of cholesterol-lowering with simvastatin in 5963 people with diabetes: a randomised placebo-controlled trial. *Lancet.* 2003; 361: 2005-2016.
42. Holme I, Fayyad R, Faergeman O, et al. Cardiovascular outcomes and their relationships to lipoprotein components in patients with and without chronic kidney disease: results from the IDEAL trial. *J Intern Med.* 2010; 267: 567-575.
43. Huskey J, Lindenfeld J, Cook T, et al. Effect of simvastatin on kidney function loss in patients with coronary heart disease: findings from the Scandinavian Simvastatin Survival Study (4S). *Atherosclerosis.* 2009; 205: 202-206.

44. Kendrick J, Shlipak MG, Targher G, et al. Effect of lovastatin on primary prevention of cardiovascular events in mild CKD and kidney function loss: a post hoc analysis of the Air Force/Texas Coronary Atherosclerosis Prevention Study. *Am J Kidney Dis.* 2010; 55: 42-49.
45. Kimura S, Inoguchi T, Yokomizo H, et al. Randomized comparison of pitavastatin and pravastatin treatment on the reduction of urinary albumin in patients with type 2 diabetic nephropathy. *Diabetes Obes Metab.* 2012; 14: 666-669.
46. Kinouchi K, Ichihara A, Bokuda K, et al. Effects of adding ezetimibe to fluvastatin on kidney function in patients with hypercholesterolemia: a randomized control trial. *J Atheroscler Thromb.* 2013; 20: 245-256.
47. Koren MJ, Davidson MH, Wilson DJ, et al. Focused atorvastatin therapy in managed-care patients with coronary heart disease and CKD. *Am J Kidney Dis.* 2009; 53: 741-750.
48. Kouvelos GN, Arnaoutoglou EM, Millionis HJ, et al. The effect of adding ezetimibe to rosuvastatin on renal function in patients undergoing elective vascular surgery. *Angiology.* 2015; 66: 128-135.
49. Lam KS, Cheng IK, Janus ED, et al. Cholesterol-lowering therapy may retard the progression of diabetic nephropathy. *Diabetologia.* 1995; 38: 604-609.
50. Lee TM, Lin MS, Tsai CH, et al. Add-on and withdrawal effect of pravastatin on proteinuria in hypertensive patients treated with AT receptor blockers. *Kidney Int.* 2005; 68: 779-787.
51. Lemos MM, Watanabe R, Carvalho AB, et al. Effect of rosuvastatin and sevelamer on the progression of coronary artery calcification in chronic kidney disease: a pilot study. *Clin Nephrol.* 2013; 80: 1-8.
52. Mou S, Wang Q, Yu Z, et al. Pravastatin improves renal progression in patients with chronic glomerulonephritis. *Int J Clin Exp Med.* 2016; 9: 1732-1739.
53. Nanayakkara PW, van GC, Ter Wee PM, et al. Effect of a treatment strategy consisting of pravastatin, vitamin E, and homocysteine lowering on carotid intima-media thickness, endothelial function, and renal function in patients with mild to moderate chronic kidney disease: results from the Anti-Oxidant Therapy in Chronic Renal Insufficiency (ATIC) Study. *Arch Intern Med.* 2007; 167: 1262-1270.
54. Ohsawa M, Tamura K, Wakui H, et al. Effects of pitavastatin add-on therapy on chronic kidney disease with albuminuria and dyslipidemia. *Lipids Health Dis.* 2015; 14: 161.
55. Rahman M, Baimbridge C, Davis BR, et al. Progression of kidney disease in moderately hypercholesterolemic, hypertensive patients randomized to pravastatin versus usual care: a report from the Antihypertensive and Lipid-Lowering Treatment to Prevent Heart Attack Trial (ALLHAT). *Am J Kidney Dis.* 2008; 52: 412-424.

56. Rutter MK, Prais HR, Charlton-Menys V, et al. Protection Against Nephropathy in Diabetes with Atorvastatin (PANDA): a randomized double-blind placebo-controlled trial of high- vs. low-dose atorvastatin(1). *Diabet Med.* 2011; 28: 100-108.
57. Sawara Y, Takei T, Uchida K, et al. Effects of lipid-lowering therapy with rosuvastatin on atherosclerotic burden in patients with chronic kidney disease. *Intern Med.* 2008; 47: 1505-1510.
58. Scanferla F, Toffoletto PP, Roncali D, et al. Associated effect of hepatic hydroxymethylglutaryl coenzyme A reductase + angiotensin converting enzyme inhibitors on the progression of renal failure in hypertensive subjects. *Am J Hypertens.* 1991; 4: 868.
59. Shepherd J, Kastelein JJ, Bittner V, et al. Effect of intensive lipid lowering with atorvastatin on renal function in patients with coronary heart disease: the Treating to New Targets (TNT) study. *Clin J Am Soc Nephrol.* 2007; 2: 1131-1139.
60. Takazakura A, Sakurai M, Bando Y, et al. Renoprotective effects of atorvastatin compared with pravastatin on progression of early diabetic nephropathy. *Journal of Diabetes Investigation.* 2015; 6: 346-353.
61. Tonelli M, Isles C, Craven T, et al. Effect of pravastatin on rate of kidney function loss in people with or at risk for coronary disease. *Circulation.* 2005; 112: 171-178.
62. Yasuda G, Kuji T, Hasegawa K, et al. Safety and efficacy of fluvastatin in hyperlipidemic patients with chronic renal disease. *Ren Fail.* 2004; 26: 411-418.
63. Kimura G, Kasahara M, Ueshima K, et al. Effects of atorvastatin on renal function in patients with dyslipidemia and chronic kidney disease: assessment of clinical usefulness in CKD patients with atorvastatin (ASUCA) trial. *Clin Exp Nephrol.* 2017; 21: 417-424.
64. Dalla Nora E, Passaro A, Zamboni PF, et al. Atorvastatin improves metabolic control and endothelial function in type 2 diabetic patients: a placebo-controlled study. *J Endocrinol Invest.* 2003; 26: 73-78.
65. Fried LF, Forrest KY, Ellis D, et al. Lipid modulation in insulin-dependent diabetes mellitus: effect on microvascular outcomes. *J Diabetes Complications.* 2001; 15: 113-119.
66. Naci H, Brugts JJ, Fleurence R, et al. Dose-comparative effects of different statins on serum lipid levels: a network meta-analysis of 256,827 individuals in 181 randomized controlled trials. *Eur J Prev Cardiol.* 2013; 20: 658-670.
67. Zoppini G, Targher G, Chonchol M, et al. Predictors of Estimated GFR Decline in Patients with Type 2 Diabetes and Preserved Kidney Function. *Clin J Am Soc Nephrol.* 2012; 7: 401.
68. Hayward RA, Krumholz HM. Three reasons to abandon low-density lipoprotein targets: an open letter to the Adult Treatment Panel IV of the National Institutes of Health. *Circ Cardiovasc Qual Outcomes.* 2012; 5: 2-5.
69. Zhou Q, Liao JK. Pleiotropic effects of statins. - Basic research and clinical perspectives. *Circ J.* 2010; 74: 818-826.

70. Wanner C, Tonelli M. KDIGO Clinical Practice Guideline for Lipid Management in CKD: summary of recommendation statements and clinical approach to the patient. *Kidney Int.* 2014; 85: 1303-1309.
71. Rucker G, Schwarzer G, Carpenter JR, et al. Undue reliance on I(2) in assessing heterogeneity may mislead. *BMC Med Res Methodol.* 2008; 8: 79.
72. Lin J, Knight EL, Hogan ML, et al. A comparison of prediction equations for estimating glomerular filtration rate in adults without kidney disease. *J Am Soc Nephrol.* 2003; 14: 2573-2580.

SUPPLEMENTARY DATA

	Random sequence generation						Concealed allocation						Selective reporting						Blinding of participants and personnel						Blinding of outcome assessor						Incomplete outcome data					
Abe 2015	?	?	-	+	-	-	?	-	?	-	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Amarenco 2014	?	-	?	-	-	-	?	-	?	-	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Athyros 2004	?	?	+	+	-	-	?	?	+	+	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Atthobari 2006	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Bianchi 2003	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Castelao 1993	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Colhoun 2009	-	-	+	-	-	-	-	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Dalla Nora 2003	?	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Deedwania 2015	?	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Fassett 2010	-	?	-	+	-	-	-	-	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Fassett 2010	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Fellstrom 2004	?	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Fried 2001	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Gheith 2002	?	?	-	?	-	-	?	?	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Haynes 2014	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Holme 2010	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
HPS 2003	?	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Huskey 2009	?	-	?	-	-	-	?	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Kendrick 2010	?	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Kimura 2017	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Kimura 2012	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Kinouchi 2013	?	+	-	+	-	-	?	+	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Koren 2009	-	?	-	+	-	-	-	-	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Kouvelos 2015	-	?	-	+	-	-	-	-	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Lam 1995	?	-	-	+	-	-	?	-	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Lee 2005	?	-	?	-	-	-	?	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Lemos 2013	-	?	-	+	-	-	-	-	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Mori 1992	?	?	-	?	-	-	?	?	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Mou 2016	-	-	-	+	-	-	-	-	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Nanayakkara 2007	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Ohsawa 2015	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Rahman 2008	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Rutter 2011	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Sawara 2008	?	?	-	?	-	-	?	?	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Scanferla 1991	?	?	-	?	-	-	?	?	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Shepherd 2007	?	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Takazakura 2015	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Tonelli 2005	?	-	?	-	-	-	?	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Vidt 2011	?	-	-	-	-	-	?	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Yakusevich 2013	?	?	?	+	-	-	?	?	?	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Yasuda 2004	?	?	-	+	-	-	?	?	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Zeeuw, de 2015 PI I	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Zeeuw, de 2015 PI II	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	

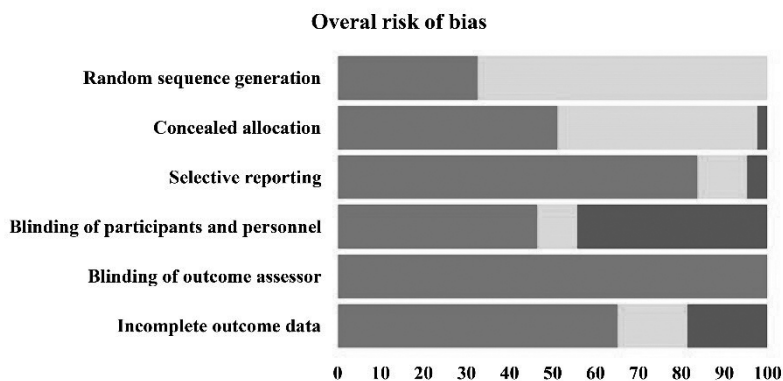


Figure S1: Risk of bias assessment per study (upper panel, page 157) and summarized over all studies (lower panel), according to the Cochrane Risk of Bias tool. Red, green and yellow cells mean high, low, and unclear risk of bias, respectively. PI I and PI II refer to PLANET I and II trials, respectively.

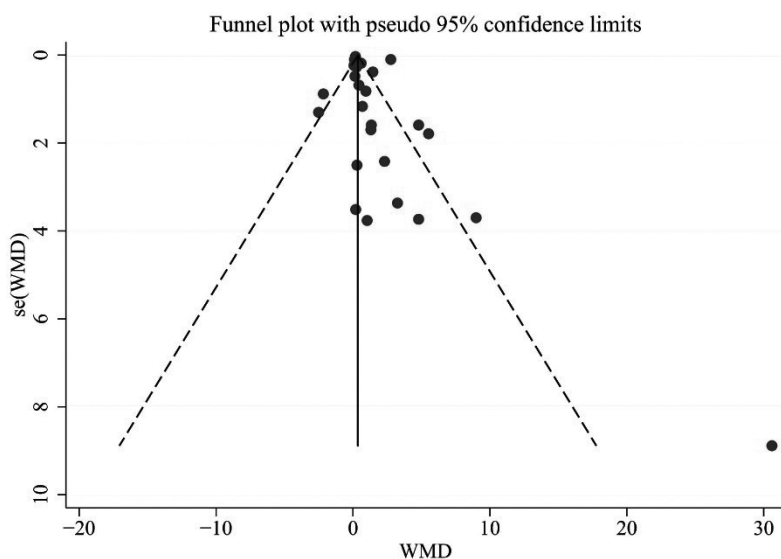


Figure S2: Funnel plot of included randomized controlled trials investigating the effect of statin therapy on annual eGFR decline. According to Egger's test there was no evidence for publication bias ($p = 0.3$). eGFR, estimated glomerular filtration rate; WMD, weighted mean difference.

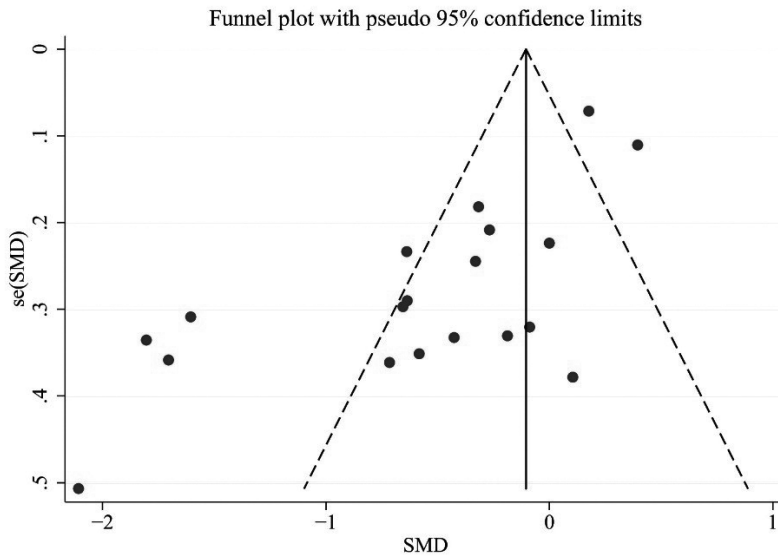


Figure S3: Funnel plot of included randomized controlled trials investigating the effect of statin therapy on change in proteinuria. According to Egger's test there was significant evidence for publication bias ($p < 0.001$). SMD, standardized mean difference.

