

Use of Statins and Risk of Hospitalization With Dementia

A Danish Population-based Case-control Study

Henriette Thisted Horsdal, MSc,*† Anne Vingård Olesen, MSc,‡§
 Christiane Gasse, Dr.rer.med.,*|| Henrik Toft Sørensen, MD, DMSc,*¶
 Robert C. Green, MD, MPH,¶¶ and Søren Paaske Johnsen, MD, PhD*

Abstract: Several epidemiologic studies have indicated reduced risk of dementia among users of statins. We assessed the risk of hospitalization with dementia associated with use of statins in a population-based case-control study in 4 Northern Danish counties in the period 1991 to 2005. We identified 11,039 cases with dementia and 110,340 age- matched and sex-matched population controls using data from the National Patient Registry, the Danish Psychiatric Central Register, and the Civil Registration System. Prescriptions for statins filled before the admission for dementia were identified using population-based prescription databases. We used conditional logistic regression analysis to compute relative risk of hospitalization with dementia associated with use of statins using nonusers as reference group. We found an overall reduced risk of hospitalization with dementia among statin users (adjusted odds ratio: 0.67, 95% confidence intervals: 0.60-0.75). The reduced risk associated with statin use remained robust in various subanalyses, however, we found no clear dose-response pattern between the number of filled prescriptions for statin and the risk of hospitalization with dementia. In conclusion, we found a reduced risk of hospitalization with dementia among users of statins, however, whether this association is causal remains to be clarified.

Key Words: statins, dementia, risk, epidemiology, case-control study

(*Alzheimer Dis Assoc Disord* 2008;00:000–000)

Received for publication November 20, 2006; accepted April 24, 2008.
 From the *Department of Clinical Epidemiology; †Unit for Psychiatric Research, Aalborg Psychiatric Hospital; Departments of ‡Pharmacology; ‡Epidemiology, Institute of Public Health; ||National Center for Registry-based Research, University of Aarhus, Aarhus, Denmark; and ¶Departments of Neurology, Medicine (Genetics), and Epidemiology, Boston University Schools of Medicine and Public Health, Boston, MA.

Supported by the Western Danish Research Forum for Health Sciences and NIH Grants AG13846, HG02213, and AG09029.

Reprints: Henriette Thisted Horsdal, MSc, Department of Clinical Epidemiology, Aarhus University Hospital, Olof Palmes Allé 43-45, DK-8200 Aarhus N, Denmark (e-mail: ht@dce.au.dk).

Copyright © 2008 by Lippincott Williams & Wilkins

Vascular and lipid-related mechanisms are thought to have a role in the pathogenesis of both Alzheimer disease and vascular dementia.^{1,2} It is, therefore, possible that treatment with lipid-lowering drugs, including statins, could lessen the risk of dementia and slow the progression of the disease. In addition to their beneficial effects on cholesterol levels, statins have other pharmacologic effects that may influence the pathogenesis of Alzheimer disease and dementia, including anti-inflammatory, antioxidant, and neuroprotective effects.³ However, the epidemiologic evidence is contradictory. Several observational studies, primarily case-control and cross-sectional studies, have shown a protective role of statins,^{4–16} but a number of cohort studies have not found a clear association between statins and reduced risk of dementia.^{6,10,14} Most randomized controlled trials have not demonstrated any protective effect of statins, but it should be noted that many of those published so far were primarily designed to evaluate the vascular effects, using cognitive function as a surrogate measure for dementia only as a secondary end point.^{17,18} Results from a recent randomized controlled trial with a 1-year exposure to either atorvastatin or placebo found that statins may have a positive effect on the progressive deterioration of cognitive function and behaviors anticipated in mild to moderate Alzheimer disease.¹⁹

The lack of consistency in the existing observational studies may reflect differences in the study design and analytic methods. Also, the potential effects of indication bias may influence the results as patients prescribed statins may have a lower baseline risk of dementia compared with other patients. This hypothesis is supported by the fact that some physicians might be less motivated to prescribe long-term preventive agents to patients who shows signs of cognitively impairment.^{20,21} Further, unaccounted confounding, such as socioeconomic factors, could also affect the results of the observational studies. Physicians may be more likely to prescribe statins to patients who are highly educated, who have knowledge about the drugs, who are concerned about their future health, and who actually ask for the prescriptions/treatment.²²

Here we conducted a large population-based case-control study based on prospectively collected data to

further explore the association between use of statins and the risk of hospitalization with dementia.

METHODS

Study Population

We conducted the study within the population of North Jutland, Aarhus, Viborg, and Ringkøbing Counties, Denmark from January 1, 1991 to December 31, 2005.

The National Health Service provides tax-supported healthcare for all inhabitants in Denmark, that is, guaranteeing free access to general practitioners and hospitals, and refunds a variable proportion of costs of medication prescribed by physicians. Since 1968, the Danish Civil Registration System has maintained electronic records of the entire Danish population and assigned a unique identification number (ie, the civil registry number) to all Danish residents at birth. All services are registered by use of the civil registry number, and it is used in all public Danish registers. Use of the civil registry number secures valid linkage between various population-based registers.

Cases of Dementia

We used computerized data from the National Patient Registry²³ and the Danish Psychiatric Central Register²⁴ to identify cases hospitalized with dementia in North Jutland (January 1992 to December 2005), Aarhus (January 1997 to December 2005), and Viborg and Ringkøbing (January 1999 to December 2005) corresponding to the availability of computerized prescription data with a minimum length of prescription history for all cases and controls of 1 year.

The National Patient Registry, established in the year 1977, store data on all discharges from nonpsychiatric hospitals in the county, including the civil registry number, dates of admission and discharge, surgical procedure(s) performed, and up to 20 discharge diagnoses assigned by the treating physician, whereas the Danish Psychiatric Central Register, established in the year 1969, store data on all discharges from all psychiatric hospitals and psychiatric departments in general hospitals, including the civil registry number, dates of admission and discharge, and all diagnoses assigned by the treating physician.

The diagnoses were classified according to the Danish version of the International Classification of Diseases (ICD), 8th until the end of 1993 and the 10th revision thereafter. On the basis of the data going back to 1977 or 1969, we constructed the hospital history for all persons, who were residents in the counties, and hereafter identified all patients, who were registered with a diagnosis of dementia (ICD-8: 29009, 29010, 29018, 29019, 29309, 29319; ICD-10: F00, F01, F03, G30) during the study period. Among these cases, we further identified patients with Alzheimer disease (ICD-8: 29010; ICD-10: F00, G30). After excluding cases that were not residents in the counties at study start and cases with unconfirmed diagnoses of dementia during the hospital

stay (modification code 1 or 2 in the ICD-8 period), a total of 11,039 cases were available for analysis.

Population Controls

For each dementia case, we aimed to identify 10 age-matched and sex-matched controls from the general population of the county through the Civil Registration System using risk set sampling,²⁵ that is, the controls had to be alive and at risk of hospitalization with dementia at the time the corresponding case was diagnosed (index date). The Civil Registration System is updated daily, keeps electronic records on vital status (dead or alive), date of death, and residence of all Danish inhabitants. We identified a total of 110,340 population controls.

Data on Statin Use

We used the population-based prescription databases²⁶ of North Jutland, Aarhus, Viborg, and Ringkøbing to obtain data on prescriptions for statins, as statins are only available by prescription in Denmark. The databases were initiated on January 1, 1991 in North Jutland, 1996 in Aarhus, and 1998 in Viborg and Ringkøbing.

The counties are served by pharmacies equipped with electronic accounting systems that are primarily used to secure reimbursement from the Danish National Health Service. The registered data include type of drug and the date the prescription is filled.

We identified all prescriptions for statins [fluvastatin, simvastatin, atorvastatin, pravastatin, lovastatin, cerivastatin, and rosuvastatin (ATC codes: C10AA01-07, B04AB01-04)] filled by cases and controls before the index date. The cases and controls were then classified according to their use of statins: either having filled no prescriptions of statins in the last 5 years or having filled 1 or more prescriptions of statins within the last 5 years. Users of statins within the last 5 years were further classified according to their total number of filled prescriptions within the 5 years, that is, 1 to 4 prescriptions, 5 to 10 prescriptions, 11 to 19 prescriptions, and > 19 prescriptions.

Data on Possible Confounding Factors

Data on possible confounding factors were obtained from the National Patient Registry, the prescription databases, and the Prevention Registry at Statistics Denmark. We obtained information on prior discharge diagnoses of hypertension, stroke, ischemic heart disease, invasive coronary revascularizations, alcohol-related diseases, diabetes mellitus, and chronic bronchitis and emphysema (as a proxy for severe smoking) from the National Patient Registry. Data from the prescription databases included all prescriptions for other lipid-lowering drugs than statins, antiplatelets, antihypertensives, high-dose aspirin, nonaspirin nonsteroidal anti-inflammatory drugs, peroral anticoagulants, hormone replacement therapy, and insulin and other antidiabetic drugs filled within 5 years before hospitalization with dementia or the corresponding date for controls.

Data on socioeconomic status of both cases and controls were obtained from the Prevention Registry at

Statistics Denmark.²⁷ This registry collects data from several other registers, that is, health-related registers, registers on living conditions, and registers of population statistics. The cases and controls were classified according to marital status (single, married, or cohabiting), employment status (old-age pensioner, self-employed, or salaried employed), gross income (below 20th percentile, 20th to 40th, 40th to 60th, 60th to 80th, or above 80th percentile), and educational level (university degree, short/medium-term formal education, basic vocational education, or unspecified).

Statistical Analysis

We formed contingency tables and used conditional logistic regression analysis to compute crude and adjusted odds ratios (ORs) of hospitalization with dementia according to use of statins. As we used risk set sampling of controls, these ORs are unbiased estimates of the corresponding incidence rate ratio.²⁵ Nonusers were used as the reference group in all analyses. We adjusted for a previous history of hypertension, stroke, ischemic heart disease, invasive coronary revascularizations, alcohol-related diseases, diabetes mellitus, chronic bronchitis, and emphysema, and for current use of other lipid-lowering drugs than statins, antiplatelets, antihypertensives, high-dose aspirin, nonaspirin nonsteroidal anti-inflammatory drugs, peroral anticoagulants, hormone replacement therapy, and socioeconomic status in the logistic regression analyses. Information on socioeconomic status was missing for a part of the study population, and these subjects were given a separate “missing” classification for those variables for inclusion in the adjusted models. We also did the analyses excluding these subjects. We further restricted the analyses to cases with Alzheimer disease and their controls. Stratified analyses were performed by age, sex, calendar time period (1992 to 2000 and 2001 to 2005), and prior cardiovascular disease (stroke, ischemic heart disease, and invasive coronary revascularizations). All analyses were performed using SAS version 8.02 (SAS Institute Inc, Cary, NC).

RESULTS

Table 1 shows the descriptive data for the 11,039 dementia cases and 110,340 controls.

Among cases and controls, 389 (3.5%) and 4855 (4.4%), respectively, filled at least 1 prescription for statins in the last 5 years before a hospital diagnosis of dementia or index date among controls. Hypertension, stroke, ischemic heart disease, alcohol-related disease, diabetes mellitus, chronic bronchitis, and emphysema were more prevalent among cases than controls, and more cases than controls had filled prescriptions for antiplatelets, antihypertensives, high-dose aspirin, insulin, and oral antidiabetic drugs. Cases were also more likely to be single and old-age pensioner.

After adjusting for possible confounding factors and further adjustment for socioeconomic status, use of statins was associated with a reduced risk of hospitalization with dementia [adjusted OR: 0.63 and 0.67, 95%

TABLE 1. Descriptive Characteristics of Cases Hospitalized With Dementia and Age-matched and Sex-matched Population Controls

	Cases (%)	Controls (%)
Total	11,039 (100.0)	110,340 (100.0)
Age (y)		
0-59	239 (2.2)	2412 (2.2)
60-79	4194 (38.0)	41,983 (38.0)
≥ 80	6606 (59.8)	65,945 (59.8)
Sex		
Male	4217 (38.2)	42,149 (38.2)
Female	6822 (61.8)	68,191 (61.8)
Discharge diagnosis of		
Hypertension	1624 (14.7)	10,789 (9.8)
Stroke	1520 (13.8)	6907 (6.3)
Ischemic heart disease	1769 (16.0)	13,705 (12.4)
Invasive coronary revascularizations	191 (1.7)	2403 (2.2)
Alcohol-related diseases	266 (2.4)	462 (0.4)
Diabetes mellitus	947 (8.6)	4946 (4.5)
Chronic bronchitis and emphysema	721 (6.5)	6072 (5.5)
Liver cirrhosis	2 (< 0.1)	53 (0.1)
Prescription for		
Statins	389 (3.5)	4855 (4.4)
Other lipid-lowering drugs than statins	25 (0.2)	358 (0.3)
Antiplatelets	3,137 (28.4)	23,847 (21.6)
Antihypertensives	7017 (63.6)	67,037 (60.8)
High-dose aspirin	2885 (26.1)	21,875 (19.8)
Nonaspirin NSAIDs	5069 (45.9)	50,234 (45.5)
Peroral anticoagulants	649 (5.9)	5837 (5.3)
Hormone replacement therapy	1157 (10.5)	12,013 (10.9)
Insulin or oral antidiabetic drugs	1053 (9.5)	6945 (6.3)
Marital status		
Single	6850 (62.1)	65,573 (59.4)
Married or cohabiting	2433 (22.0)	39,734 (36.0)
Missing	1756 (15.9)	5033 (4.6)
Employment status		
Old-age pensioner	9162 (83.0)	101,103 (91.6)
Self-employed or salaried employed	121 (1.1)	4204 (3.8)
Missing	1756 (15.9)	5033 (4.6)
Gross income		
Below 20th perc	1625 (14.7)	20,532 (18.6)
20th-40th perc	2253 (20.4)	23,071 (20.9)
40th-60th perc	2131 (29.3)	22,949 (20.8)
60th-80th perc	1837 (16.7)	20,185 (18.3)
Above 80th perc	1437 (13.0)	18,570 (16.8)
Missing	1756 (15.9)	5033 (4.6)
Educational level		
University degree	79 (0.7)	1050 (0.9)
Short/medium-term formal education	925 (8.4)	10,730 (9.7)
Basic vocational education	293 (2.7)	3755 (3.4)
Unspecified	7986 (72.3)	89,772 (81.4)
Missing	1756 (15.9)	5033 (4.6)

NSAID indicates nonsteroidal anti-inflammatory drug; perc, percentile.

confidence interval (CI): 0.56-0.70 and 0.60-0.75, respectively] (Table 2). The result was similar when excluding cases and controls with missing information on socioeconomic status (adjusted OR: 0.68, 95% CI: 0.60-0.78).

We found no clear association between the number of filled prescriptions for statins and risk of

TABLE 2. Crude and Adjusted Odds Ratios With 95% Confidence Intervals for Hospitalization With Dementia According to Use of Statins and the Total Number of Filled Prescriptions for Statins

	Cases (%) (N = 11,039)	Controls (%) (N = 110,340)	ORs (95% CI)	Adjusted ORs* (95% CI)	Adjusted ORs† (95% CI)
Nonusers in the last 5 years	10,650 (96.48)	105,485 (95.60)	1.00 (reference)	1.00 (reference)	1.00 (reference)
Use of statins within the last 5 years	389 (3.52)	4855 (4.40)	0.78 (0.70-0.87)	0.63 (0.56-0.70)	0.67 (0.60-0.75)
No. prescriptions					
1-4 prescriptions	86 (0.78)	1301 (1.18)	0.49 (0.39-0.62)	0.49 (0.39-0.62)	0.53 (0.43-0.67)
5-10 prescriptions	111 (1.01)	1168 (1.06)	0.73 (0.59-0.89)	0.73 (0.59-0.89)	0.78 (0.63-0.96)
11-19 prescriptions	88 (0.80)	1366 (1.24)	0.53 (0.42-0.67)	0.53 (0.42-0.67)	0.57 (0.45-0.71)
> 19 prescriptions	104 (0.94)	1020 (0.92)	0.81 (0.66-1.00)	0.81 (0.66-1.00)	0.86 (0.69-1.06)

*Adjusted for use of antiplatelets, antihypertensives, high-dose aspirin, nonaspirin NSAIDs, peroral anticoagulants, lipid-lowering drugs, and hormone replacement therapy, and previous diagnoses of hypertension, stroke, ischemic heart disease, invasive coronary revascularization, alcohol-related diseases, diabetes mellitus, and chronic bronchitis or emphysema (as a proxy for smoking).

†Adjusted for marital status, employment status, gross income, educational level, use of antiplatelets, antihypertensives, high-dose aspirin, nonaspirin NSAIDs, peroral anticoagulants, lipid-lowering drugs, and hormone replacement therapy, and previous diagnoses of hypertension, stroke, ischemic heart disease, invasive coronary revascularization, alcohol-related diseases, diabetes mellitus, and chronic bronchitis or emphysema (as a proxy for smoking).

CI indicates confidence interval; NSAID, nonsteroidal anti-inflammatory drug; OR, odds ratio.

hospitalization with dementia; the adjusted ORs were 0.53 (95% CI: 0.43-0.67), 0.78 (95% CI: 0.63-0.96), 0.57 (95% CI: 0.45-0.71), and 0.86 (95% CI: 0.69-1.06) for 1 to 4 prescriptions, 5 to 10 prescriptions, 11 to 19 prescriptions, and > 19 prescriptions, respectively.

The association between statin use and risk of hospitalization with dementia remained robust in all subanalyses when stratifying by age, sex, calendar period, and prior cardiovascular disease. The lowest risk estimates associated with statin use were found among patients younger than 60 years (adjusted OR: 0.48, 95% CI: 0.19-1.21) or 80 years or older (adjusted OR: 0.55, 95% CI: 0.44-0.68) and among patients with prior cardiovascular disease (adjusted OR: 0.46, 95% CI: 0.38-0.56). Results were also similar when restricting the analyses to cases with Alzheimer disease (adjusted OR: 0.72, 95% CI: 0.57-0.91).

In contrast, use of hormone replacement therapy among women was not associated with risk of hospitalization with dementia (adjusted OR: 0.96, 95% CI: 0.89-1.02).

Finally, we also looked at the risk estimates obtained for some of the known risk factors for dementia in our analyses to qualify the validity of our dataset. The adjusted ORs for a history of hypertension, stroke, or diabetes mellitus were 1.22 (95% CI: 1.15-1.30), 1.84 (95% CI: 1.72-1.96), and 1.58 (95% CI: 1.46-1.71), respectively.

DISCUSSION

In this population-based case-control study, users of statins had a clearly decreased risk of hospitalization with dementia. The association seemed not to be explained by a wide range of possible confounding factors, including socioeconomic factors, and the association remained robust in subanalyses stratified by age, sex, calendar time period, and prior cardiovascular disease, and when restricting the analyses to cases with Alzheimer disease. However, we found no clear association between numbers

of filled prescriptions for statin and the risk of hospitalization with dementia.

The main strengths of our study included the population-based design, and the ability to link different data sources with prospectively collected data. By use of these databases, the potential difficulties with recall bias are avoided.

Although the overall positive predictive value of a dementia diagnosis recorded in the National Patient Registry and the Danish Psychiatric Central Register is high (ie, approximately 85%),²⁸ an important limitation was the lack of details about the dementia diagnosis among the cases, including type of dementia and onset of symptoms. This information is relevant as the effects of statins on vascular dementia and Alzheimer disease may differ and as dementia is commonly first recognized or diagnosed several years after symptom onset. If an individual has early signs or symptoms of dementia, physicians might be less inclined to maintain preventive treatment, when the overall health deteriorates,²⁹ introducing indication bias in which an individual with prevalent or early dementia has less chance of receiving treatment than the control. The latter seemed not to be a major problem in our study as more cases than controls filled prescriptions for other drugs used for long-term primary and secondary prevention of cardiovascular disease, for example, antihypertensives and antiplatelets. Another limitation of our study includes lack of information on nonhospitalized dementia cases. However, this most likely leads to nondifferential misclassification and will thus bias the estimates toward unity. Similarly, any Berksons bias or surveillance bias arising owing to a potential increased probability of users of statins to be registered with a dementia diagnosis would also result in conservative risk estimates.

It has previously been discussed that statin users may be “healthy users”—younger, healthier, better educated, and socioeconomically privileged persons—who may be more likely to receive preventive treatments than the less privileged and frail.³⁰ In general, severe

confounding by socioeconomic differences is unlikely given Denmark's universal healthcare. However, we have recently reported a weak socioeconomic gradient in statin use among Danish men in the mid 90s,³¹ indicating that socioeconomic factors could also be a confounding factor in the association between use of statins and risk of dementia. However, such an explanation is unlikely as the gradient was not found among women and that it decreased in magnitude over time among men. Some of the existing observational studies that found a reduced risk of dementia did adjust for educational level as a crude measure of socioeconomic status.^{5-9,15} We added marital status, employment status, and gross income to get a better measure of socioeconomic status.^{32,33} After adjustment for these factors, we still found a decreased risk of dementia among statin users. Notably, use of hormone replacement therapy in women, which may also be subject to the "healthy user effect," had no effect on the risk of hospitalization with dementia in our study.

Although we adjusted for a number of confounding factors, we can, however, still not entirely exclude the possibility that our results remain influenced by residual, unmeasured, or unknown confounding.

Finally, it should be noted that despite the relatively large sample size, the low frequency of statin use among cases and controls in our study resulted in limited statistical precision for several of the presented risk estimates as indicated by the relatively wide confidence intervals. This low use of statins reflects a general low use of statins in Denmark.^{34,35}

Our finding that users of statins have an overall reduced risk of dementia is in agreement with other observational studies.^{4,5,7-9,11-13,19}

We found no association between the number of filled prescriptions for statins and the risk of hospitalization with dementia, which is consistent with the lack of a dose-response relationship between use of statins and dementia in other observational studies,^{10,12-14} and suggests that statins may not directly cause the reduced risk of dementia. However, an increased number of statin prescriptions could possibly also reflect more severe cardiovascular disease, which overwhelms the potential beneficial effects of the increased statin dosage.

Besides the beneficial effect on cholesterol levels, statins have other pharmacologic effects that may influence the pathogenesis of Alzheimer disease and dementia. The possible biologic mechanisms include (1) stimulation of α -secretase activity,³⁶ (2) modulation of amyloid precursor protein metabolism and A β production,³⁷ (3) immunomodulation leading to attenuation of inflammatory markers (eg, C-reactive protein and cytokines),^{38,39} (4) reduction of lipoprotein oxidation and free radical injury,⁴⁰ (5) increase in cerebral endothelial nitric oxide synthase and cerebral circulation,⁴¹ and (6) activation of signal transcription (Rho-GTPases) through depletion of downstream isoprenoids.⁴²

In conclusion, we found a reduced risk of hospitalization with dementia among users of statins, and our findings, therefore, provide some support to the hypo-

thesis that statins may protect against the development of dementia.

REFERENCES

- de la Torre JC. Alzheimer disease as a vascular disorder: nosological evidence. *Stroke*. 2002;33:1152-1162.
- Gorelick PB, William M. Feinberg lecture: cognitive vitality and the role of stroke and cardiovascular disease risk factors. *Stroke*. 2005; 36:875-879.
- Crisby M, Carlson LA, Winblad B. Statins in the prevention and treatment of Alzheimer disease. *Alzheimer Dis Assoc Disord*. 2002; 16:131-136.
- Wolozin B, Kellman W, Ruosseau P, et al. Decreased prevalence of Alzheimer disease associated with 3-hydroxy-3-methylglutaryl coenzyme A reductase inhibitors. *Arch Neurol*. 2000;57:1439-1443.
- Dufouil C, Richard F, Fievet N, et al. APOE genotype, cholesterol level, lipid-lowering treatment, and dementia: the Three-City Study. *Neurology*. 2005;64:1531-1538.
- Zandi PP, Sparks DL, Khachaturian AS, et al. Do statins reduce risk of incident dementia and Alzheimer disease? The Cache County Study. *Arch Gen Psychiatry*. 2005;62:217-224.
- Rockwood K, Kirkland S, Hogan DB, et al. Use of lipid-lowering agents, indication bias, and the risk of dementia in community-dwelling elderly people. *Arch Neurol*. 2002;59:223-227.
- Hajjar J, Schumpert J, Hirth V, et al. The impact of the use of statins on the prevalence of dementia and the progression of cognitive impairment. *J Gerontol A Biol Sci Med Sci*. 2002;57:M414-M418.
- Green RC, McNagny SE, Jayakumar P, et al. Statin use and the risk of Alzheimer's disease: the MIRAGE Study. *Alzheimer Dement*. 2006;2:96-103.
- Rea TD, Breitner JC, Psaty BM, et al. Statin use and the risk of incident dementia: the Cardiovascular Health Study. *Arch Neurol*. 2005;62:1047-1051.
- Jick H, Zornberg GL, Jick SS, et al. Statins and the risk of dementia. *Lancet*. 2000;356:1627-1631.
- Zamrini E, McGwin G, Roseman JM. Association between statin use and Alzheimer's disease. *Neuroepidemiology*. 2004;23:94-98.
- Qizilbash N, Boudiaf N, Feudjo-Tepie M, et al. Statins and dementia risk: more of the same from observational databases? *Age Ageing*. 2005;34:ii35.
- Li G, Higdon R, Kukull WA, et al. Statin therapy and risk of dementia in the elderly: a community-based prospective cohort study. *Neurology*. 2004;63:1624-1628.
- Yaffe K, Barrett-Connor E, Lin F, et al. Serum lipoprotein levels, statin use, and cognitive function in older women. *Arch Neurol*. 2002;59:378-384.
- Masse I, Bordet R, Deplanque D, et al. Lipid lowering agents are associated with a slower cognitive decline in Alzheimer's disease. *J Neurol Neurosurg Psychiatry*. 2005;76:1624-1629.
- Heart Protection Study Collaboration Group. MRC/BHF Heart Protection Study of cholesterol lowering with simvastatin in 20,536 high-risk individuals: a randomised placebo-controlled trial. *Lancet*. 2002;360:7-22.
- Shepherd J, Blauw GJ, Murphy MB, et al. Pravastatin in elderly individuals at risk of vascular disease (PROSPER): a randomised controlled trial. *Lancet*. 2002;360:1623-1630.
- Sparks DL, Sabbagh MN, Connor DJ, et al. Atorvastatin for the treatment of mild to moderate Alzheimer disease: preliminary results. *Arch Neurol*. 2005;62:753-757.
- Brauner DJ, Muir JC, Sachs GA. Treating nondementia illnesses in patients with dementia. *JAMA*. 2000;283:3230-3235.
- Moroney JT, Tseng CL, Paik MC, et al. Treatment for the secondary prevention of stroke in older patients: the influence of dementia status. *J Am Geriatr Soc*. 1999;47:824-829.
- Haley RW, Dietsch JM. Is there a connection between the concentration of cholesterol circulating in plasma and the rate of neuritic plaque formation in Alzheimer disease? *Arch Neurol*. 2000; 57:1410-1412.
- Andersen TF, Madsen M, Jørgensen J, et al. The Danish National Hospital Register. A valuable source of data for modern health sciences. *Dan Med Bull*. 1999;46:263-268.

24. Munk-Jørgensen P, Mortensen PB. The Danish Psychiatric Central Register. *Dan Med Bull.* 1997;44:82–84.
25. Wacholder S, McLaughlin JK, Silverman DT, et al. Selection of controls in case-control studies. I. Principles. *Am J Epidemiol.* 1992;135:1019–1028.
26. Gaist D, Sørensen HT, Hallas J. The Danish prescription registries. *Dan Med Bull.* 1997;44:445–448.
27. Roed AS, Juhl C, Kamper-Jørgensen F. The Danish Prevention Register. A comprehensive health and socio-economic, individual based register. *Dan Med Bull.* 1999;46:269–272.
28. Phung TK, Andersen BB, Høgh P, et al. Validity of dementia diagnoses in the Danish hospital registers. *Dement Geriatr Cogn Disord.* 2007;24:220–228.
29. Rodriguez EG, Dodge HH, Birzescu MA, et al. Use of lipid-lowering drugs in older adults with and without dementia: a community-based epidemiological study. *J Am Geriatr Soc.* 2002;50:1852–1856.
30. Glynn RJ, Schneeweiss S, Wang PS, et al. Selective prescribing led to overestimation of the benefits of lipid-lowering drugs. *J Clin Epidemiol.* 2006;59:819–828.
31. Thomsen RW, Johnsen SP, Olesen AV, et al. Socioeconomic gradient in use of statins among Danish patients: population-based cross-sectional study. *Br J Clin Pharmacol.* 2005;60:534–542.
32. Avlund K, Holstein BE, Osler M, et al. Social position and health in old age: the relevance of different indicators of social position. *Scand J Public Health.* 2003;31:126–136.
33. Huisman M, Kunst AE, Mackenbach JP. Socioeconomic inequalities in morbidity among the elderly; a European overview. *Soc Sci Med.* 2003;57:861–873.
34. Rasmussen JN, Gislason GH, Abildstrom SZ, et al. Statin use after acute myocardial infarction: a nationwide study in Denmark. *Br J Clin Pharmacol.* 2005;60:150–158.
35. Walley T, Folino-Gallo P, Schwabe U, et al. Variations and increase in use of statins across Europe: data from administrative databases. *BMJ.* 2004;328:385–386.
36. Kojro E, Gimpl G, Lammich S, et al. Low cholesterol stimulates the nonamyloidogenic pathway by its effect on the alpha -secretase ADAM 10. *Proc Natl Acad Sci USA.* 2001;98:5815–5820.
37. Fassbender K, Simons M, Bergmann C, et al. Simvastatin strongly reduces levels of Alzheimer's disease beta -amyloid peptides Abeta 42 and Abeta 40 in vitro and in vivo. *Proc Natl Acad Sci USA.* 2001;98:5856–5861.
38. Ridker PM, Rifai N, Pfeffer MA, et al. Long-term effects of pravastatin on plasma concentration of C-reactive protein. The Cholesterol and Recurrent Events (CARE) Investigators. *Circulation.* 1999;100:230–235.
39. Rosenson RS, Tangney CC, Casey LC. Inhibition of proinflammatory cytokine production by pravastatin. *Lancet.* 1999;353:983–984.
40. Sobal G, Sinzinger H. Effect of simvastatin on the oxidation of native and modified lipoproteins. *Biochem Pharmacol.* 2005;70:1185–1191.
41. Endres M, Laufs U, Huang Z, et al. Stroke protection by 3-hydroxy-3-methylglutaryl (HMG)-CoA reductase inhibitors mediated by endothelial nitric oxide synthase. *Proc Natl Acad Sci USA.* 1998;95:8880–8885.
42. Laufs U, Liao JK. Post-transcriptional regulation of endothelial nitric oxide synthase mRNA stability by Rho GTPase. *J Biol Chem.* 1998;273:24266–24271.