Cross-sectional study of the association between cataract surgery and age-related macular degeneration in the era of phacoemulsification in the national health and nutrition examination survey 2005–2008

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INTRODUCTION
Cataract and age-related macular degeneration (AMD) are leading causes of visual impairment and blindness in the USA1 and are often presenting concurrently in older persons.2 As the population ages, the burden of both age-related conditions is growing.

Cataract surgery is the only cost-effective intervention for improving visual function and quality of life in patients with cataract3 and is one of the most commonly performed surgeries in the USA.4 The presence of AMD is an established risk factor for poor visual outcome following cataract surgery.5 Although the introduction of anti-vascular endothelial growth factor (anti-VEGF) has recently revolutionised the management of exudative AMD,6,7 the high price of anti-VEGF imposes heavy economic and social burden. Furthermore, there is still no effective treatment for atrophic AMD.

The relationship between cataract surgery and AMD was first identified by a postmortem histopathologic study.8 Since then, concerns have been raised about whether cataract surgery increases the incidence and progression of AMD.9–28 Despite the extensive studies investigating this relationship, results remain contradictory.9–28 Furthermore, surgical-related inflammation and light toxicity due to aphakic cataract surgery (postulated mechanisms for the association between cataract surgery and AMD) have decreased...
A recent cross-sectional evidence from a large representative Korean population suggested no significant association between cataract surgery and the presence of AMD, except in left eyes, which was explained as a chance finding. On the contrary, a recent Taiwanese study using population-based claims data documented 1.5-fold increased risk of exudative AMD in those who underwent cataract surgery.

In the light of the rapidly ageing population, irreversible visual impairment and huge social economic burden of AMD, we aimed to elucidate the association between cataract surgery and AMD in a large, nationally representative sample of US middle-aged population.

METHODS
Study design and population
Two waves of National Health and Nutrition Examination Survey (NHANES, 2005–2006, 2007–2008) were combined. The NHANES collected health-related information in 2-year patterns and used a complex, multistage, probability sampling methodology to obtain nationally representative surveys of civilian, non-institutionalised US population. The NHANES intentionally oversampled the elderly and ethnicity minority groups. Standard protocols were used across different examination sites.

The research was performed in accordance to the tenets of the Declaration of Helsinki.

Patient and public involvement
Data used in this analysis were publicly available and deidentified NHANES data. No patient and/or public were involved in the design and conception of our study.

Cataract surgery and AMD status
Cataract surgery status was obtained from questionnaire. Prior to visual acuity examination, participants were asked the question ‘Have you ever had eye surgery to treat cataracts?’, and if so, they were further asked ‘Which eye(s) had cataract surgery?’, which included possible answers of right, left, both or do not know. Individuals who reported history of cataract surgery but did not know which eye(s) were excluded from the current analysis.

In 2005–2008 waves of NHANES survey, non-mydriatic fundus photographs were collected for participants older than 40 years using Canon CR6-45NM Ophthalmic Digital Imaging System and Canon EOS 10D digital camera (Canon USA, One Canon Park, Melville, New York, USA). Photographs were assessed by at least two graders at the University of Wisconsin, Madison according to the modification of the Wisconsin Age-Related Maculopathy Grading Classification Scheme. In brief, early AMD was defined as the presence of drusen and/or pigmentary abnormalities, and late AMD was based on the detection of exudative AMD signs or geographic atrophy.

Covariates
Demographic characteristics, including age, gender, race, education level, marital status and family income, were collected. Participants were categorised as non-Hispanic white, non-Hispanic black, Mexican American and other ethnicity. Educational level was analysed as a two-level categorical variable: less than high school and completion of or more than high school. Marital status (unmarried and other, married/with a partner) was analysed as a two-level categorical variable. The indicator for family income—the poverty income ratio—was categorised as below poverty (<1.00) and at or above poverty (≥1.00). Lifestyle factors included cigarette smoking and alcohol consumption. Former/current smoker was defined as participants who currently smoked and had smoked at least 100 cigarettes in their lifetime. Current drinker was defined as participants who currently drank.

General health was assessed by body mass index (BMI), physical activity, C reactive protein (CRP) and comorbidities. Height and weight were used to calculate BMI (kg/m²). Obesity was defined as BMI >30 kg/m². The 2008 Physical Activity Guideline for Americans suggests at least 2.5 hours of moderate-intensity or 75 min of vigorous-intensity aerobic physical activity per week or an equivalent combination of moderate-intensity and vigorous-intensity aerobic activity for health benefits. We categorised participants into two groups based on whether the 2008 Physical Activity Guideline was met or not. CRP was categorised as two groups (high CRP level: CRP ≥1 mg/dL or not).

Clinical comorbidities included diabetes mellitus, hypertension, hyperlipidaemia, chronic kidney disease (CKD) and self-reported history of cardiovascular disease (CVD). Diabetes mellitus was defined as serum glycosylated haemoglobin >6.5%,29 the use of insulin or diabetic tablets or self-reported history of diabetes mellitus. Hypertension was defined as systolic blood pressure of ≥140 mm Hg or diastolic blood pressure ≥90 mm Hg, the use of antihypertensive agents or self-reported history of hypertension. Hyperlipidaemia was based on total cholesterol ≥240 mg/dL, the use of lipid-lowering medications or self-reported history of high cholesterol. CKD was defined as estimated glomerular filtration rate <60 mL/min/1.73 m². Self-reported history of CVD was based on previous diagnosis of congestive heart failure, coronary heart disease, angina, heart attack or stroke.

Statistical analyses
All statistical analyses were performed using Stata (V.14.0; StataCorp., College Station, Texas, USA). The data set was set up with survey commands in Stata to account for the complex and multistage sampling design of NHANES to generate nationally representative estimates. Characteristics of eyes were reported by using means and SEs for continuous variables and numbers and weighted percentages for categorical variables. We used design-adjusted
RESULTS
From the 6797 eligible participants aged 40 years and older in 2005–2008 NHANES surveys, 5401 had complete data on cataract surgery status and gradable retinal photographs for right eyes were included in the present analysis. Compared with participants with complete data, participants with incomplete data were more likely to be older, non-Hispanic black, less educated, unmarried, in disadvantaged socioeconomic status, never smoker, abstainer/former drinker and unhealthier regarding physical activity and clinical comorbidities (all p<0.05). Other characteristics of participants excluded and included are illustrated in online supplemental table 1. Overall, 566 right eyes (8.0%) underwent cataract surgery. Of 338 right eyes with any AMD, 107 right eyes (28.9%) underwent cataract surgery. Table 1 provides the characteristics of right eyes. Univariable logistic regression models showed covariates, including age, race, marital status, smoking status, obesity, diabetes mellitus, hypertension, CKD and self-reported CVD, were significantly associated with any AMD and early AMD. There were statistically significant associations between late AMD and age, race, marital status, obesity, hypertension, hyperlipidaemia, CKD or self-reported CVD in the univariable logistic models. These results from logistic regression modelling are outlined in table 2. After adjusting for significant covariates identified in the univariable logistic regression models, significant associations were observed between cataract surgery and any AMD (OR 1.36; 95% CI 1.03 to 1.81) or late AMD (OR 2.48; 95% CI 1.01 to 6.09). However, no significant association was found between cataract surgery and early AMD after adjusting for multiple covariates for right eyes. Detailed results are provided in table 3. Similar results were observed when performing sensitivity analysis for one-way analysis of variance and Rao-Scott Pearson χ² for the comparison of continuous and categorical variables, respectively. Logistic regression modelling was used to determine the association of cataract surgery with AMD status. Separate analyses were conducted for any, early or late AMD. Unadjusted models were used to estimate ORs and 95% CIs for each status of AMD, to determine covariates significantly associated with outcomes. Significant covariates (p<0.05), along with cataract surgery status, were added to models investigating their effects on each status of AMD in multivariable-adjusted logistic regression models. In order to verify the robustness of our results, we also performed sensitivity analysis for left eyes and combing available data from both eyes. Multivariable-adjusted generalised estimating equation (GEE) models were used to account for correlations of paired eyes within the same participant and determine relationships between cataract surgery and AMD status. Two-sided p values <0.05 were considered significant for statistical inferences.

**DISCUSSION**
In this large nationally representative sample of the US population 40 years and older, our study provided evidence that previous cataract surgery may be associated with AMD, particularly late AMD. The relationship between cataract surgery and AMD has been intensively investigated through postmortem histopathologic study, cross-sectional study, cohort study, retrospective case–control study and randomised controlled trial, but results have been conflicting.9–28 The significant link between cataract surgery and AMD in our study is consistent with previous studies.8–10 12 13 17 Dated back to 1989, Liu et al.9 found that cataract surgery without intraocular implantation might place participants at increased risks of AMD using the nationally representative cross-sectional study (NHANES-I). After pooling results from three population-based studies, Freeman and colleagues claimed the strong association between cataract surgery and late AMD.11 Longitudinal studies, such as the Beaver Dam Eye Study and Blue Mountain Eye Study, documented higher risks of late AMD after cataract surgery.10 12 13 17 Similarly, cataract surgery was significantly associated with long-term incidence of late AMD in the Rotterdam Eye Study.8 15 Three recent clinical-based case–control studies consistently demonstrated that cataract surgical patients had higher risks of development and progression of AMD with follow-up length ranging from 1 to 5 years.14 18 19 Our results were contrary to findings from some previously published studies.20–28 Population-based

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**Table 1** Characteristics of participants by right eyes

| Characteristics          | Right eyes |
|--------------------------|------------|
|                         | (n=5401)   |
| Age (SE), years          | 56.1±0.39  |
| Female, number (%)       | 2713 (52.9)|
| Cataract surgery, number (%) | 566 (8.0) |
| Any AMD, number          | 338        |
| Age (SE), years          | 68.9±0.88  |
| Cataract surgery, number (%) | 107 (28.9) |
| Early AMD, number        | 296        |
| Age (SE), years          | 67.6±0.93  |
| Cataract surgery, number (%) | 81 (24.6) |
| Late AMD, number         | 42         |
| Age (SE), years          | 78.1±1.00  |
| Cataract surgery, number (%) | 26 (60.6) |

All proportions, means and standard errors are weighted estimates of the US population characteristics, taking into account the complex sampling design of the National Health and Nutrition Examination Survey. AMD, age-related macular degeneration.
## Table 2 Logistic regression models of risk factors for AMD

| Characteristics                        | Any AMD* OR (95% CI) | P value | Early AMD* OR (95% CI) | P value | Late AMD* OR (95% CI) | P value |
|----------------------------------------|----------------------|---------|------------------------|---------|-----------------------|---------|
| Age (SE), years                        | 1.10 (1.08 to 1.11)  | <0.001  | 1.09 (1.07 to 1.10)    | <0.001  | 1.24 (1.18 to 1.31)   | <0.001  |
| Female versus male                     | 0.94 (0.73 to 1.20)  | 0.59    | 0.85 (0.65 to 1.11)    | 0.22    | 2.07 (0.94 to 4.56)   | 0.07    |
| Race                                   |                      |         |                        |         |                       |         |
| Non-Hispanic white                     | Reference            | 0.007   | Reference              | 0.022   |                       | 0.045   |
| Non-Hispanic black                     | 0.31 (0.18 to 0.51)  |         | 0.31 (0.17 to 0.56)    |         | 0.31 (0.08 to 1.17)   |         |
| Mexican American                       | 0.53 (0.34 to 0.82)  |         | 0.60 (0.39 to 0.93)    |         |                       |         |
| Other                                  | 0.57 (0.30 to 1.06)  |         | 0.60 (0.31 to 1.18)    |         | 0.30 (0.06 to 1.59)   |         |
| High school and over versus less       | 0.75 (0.50 to 1.14)  | 0.17    | 0.74 (0.47 to 1.16)    | 0.19    | 0.82 (0.40 to 1.67)   | 0.57    |
| Married/with a partner versus unmarried or other | 0.62 (0.48 to 0.79)  | <0.001  | 0.69 (0.52 to 0.91)    | 0.010   | 0.29 (0.14 to 0.61)   | 0.002   |
| Income at or above poverty versus below| 0.78 (0.55 to 1.12)  | 0.17    | 0.85 (0.56 to 1.28)    | 0.42    | 0.49 (0.22 to 1.08)   | 0.08    |
| Former/current smoker versus non-smoker | 1.34 (1.04 to 1.72)  | 0.024   | 1.35 (1.02 to 1.79)    | 0.035   | 1.24 (0.58 to 2.66)   | 0.57    |
| Current drinker versus abstainer/former drinker | 0.74 (0.49 to 1.12)  | 0.15    | 0.79 (0.53 to 1.19)    | 0.26    | 0.45 (0.20 to 1.02)   | 0.05    |
| Obesity present versus absent          | 0.63 (0.47 to 0.85)  | 0.004   | 0.69 (0.51 to 0.94)    | 0.021   | 0.26 (0.10 to 0.71)   | 0.010   |
| Meet PA recommendation versus not meet | 0.79 (0.52 to 1.21)  | 0.27    | 0.79 (0.52 to 1.21)    | 0.27    | 0.79 (0.33 to 1.91)   | 0.59    |
| High CRP present versus absent         | 1.40 (0.92 to 2.13)  | 0.11    | 1.38 (0.89 to 2.13)    | 0.15    | 1.58 (0.60 to 4.19)   | 0.34    |
| Diabetes mellitus present versus absent | 1.44 (1.06 to 1.95)  | 0.020   | 1.54 (1.11 to 2.15)    | 0.012   | 0.73 (0.27 to 1.98)   | 0.53    |
| Hypertension present versus absent     | 2.20 (1.55 to 3.13)  | <0.001  | 2.19 (1.49 to 3.20)    | <0.001  | 2.31 (1.04 to 5.17)   | 0.041   |
| Hyperlipidaemia present versus absent  | 1.17 (0.82 to 1.66)  | 0.38    | 1.05 (0.73 to 1.52)    | 0.78    | 2.46 (1.18 to 5.14)   | 0.018   |
| CKD present versus absent              | 4.64 (3.32 to 6.48)  | <0.001  | 4.12 (2.95 to 5.75)    | <0.001  | 10.2 (4.70 to 22.3)   | <0.001  |
| Self-reported CVD present versus absent| 3.47 (2.48 to 4.85)  | <0.001  | 3.36 (2.42 to 4.65)    | <0.001  | 4.40 (2.14 to 9.07)   | <0.001  |

Boldface indicates statistical significance (P < 0.05).

*Comparisons were between each AMD group and the no AMD group.

AMD, age-related macular degeneration; CKD, chronic kidney disease; CRP, C reactive protein; CVD, cardiovascular disease; PA, physical activity.

Cross-sectional results from Visual Impairment Project, Beijing Eye Study and Korean NHANES found no significant association between cataract surgery and AMD. The only randomised controlled trial conducted on this topic suggested clear benefits of phacoemulsification for patients at high risk of AMD progression but without significant evidence for the progression to choroidal neovascularisation. However, the generalisability of this result might be subject to the small sample size (n=56). The lack of a significant association between cataract surgery and AMD was also documented in clinical-based studies, including a study of Armbrecht et al., a study of Baatz et al. and CSAMD. The AREDS concluded no significant effect of cataract surgery on the progression of AMD to advanced status. However, the analysis in the AREDS did find a statistically significant increase in the risk of advanced AMD in eyes with cataract surgery. Of note, many differences, including demographics, age range, sample size, study design, follow-up length, analytic paradigm and confounding factors, exist between our study and earlier studies, which might have led to the discrepancies. Previous studies investigating the association between cataract surgery and AMD are summarised in Table 4.

Several possible explanations for the association between cataract surgery and AMD have been proposed.
One possibility is the similar underlying mechanism behind cataract and AMD. Previous studies indicated that AMD appeared to be more prevalent in patients with cataract.\textsuperscript{9,11,31} Several physiologic or genetic mechanisms underlying lens opacity formation may also play important roles in the pathogenesis of macular degeneration.\textsuperscript{9,12} Due to the high level of cataract surgical coverage in the USA, the cataract surgery can be regarded as a surrogate of cataract. Alternatively, common risk factors shared by cataract and AMD potentially have effects on their relationship, including older age, smoking status and CVD.\textsuperscript{32,33} However, in our study, a comprehensive series of confounders were adjusted in final models and the significant association between cataract surgery and AMD still persisted. Of note, we could not exclude the possibility of incomplete adjustment for important unknown factors, such as genetic factors.

Another proposed explanation for the significant association between cataract surgery and AMD is surgical inflammation. Inflammatory response is common after cataract surgery and may exacerbate the short-term risk of occurrence and progression of AMD.\textsuperscript{34} A retrospective cohort study indicated higher susceptibility to cystoid macular oedema or deterioration of choroidal neovascularisation after cataract surgery among patients with wet AMD.\textsuperscript{38} Therefore, we speculated that cataract surgery might predispose eyes to the development or progression of AMD. Of note, techniques in cataract surgery have been improved greatly and phacoemulsification procedures have sharply reduced inflammatory responses. The significant association between cataract surgery and AMD in the current phacoemulsification era is, to less extent, due to the inflammation caused by cataract surgery. The third possible explanation for the increased risk of AMD after cataract surgery is the greater intensity of UV or short-wavelength light exposure after cataract surgery.\textsuperscript{35–37} Several studies indicated that lens might play a protective effect on retina health.\textsuperscript{14} Aphakic patients had been documented to show more severe macular degeneration than those phakic subjects.\textsuperscript{9} However, the greater likelihood of cataract surgery with implantation of intraocular lenses and advent of UV-blocking or blue light-blocking intraocular lenses had suggested to largely reduce the insult of UV or short-wavelength light on the macular. However, a very recent systemic review concluded that there was still unclear regarding the protective function of blue light-filtering intraocular lenses on macular health or the reduced risk associated with the development or progression of AMD.\textsuperscript{38} Other studies suggested that mechanisms underlying the significant relationship between cataract surgery and AMD could be the intraoperative photic damage caused by the operating microscope.\textsuperscript{39} Further research is needed to clarify possible mechanisms underlying the significant association between cataract surgery and AMD, if this significant association is confirmed in other studies.

Our study has several strengths, including the large nationally representative sample size, standardised protocol for grading AMD and availability of comprehensive confounders. Of note, our study was also limited by several points. First, as a cross-sectional study, our study could not provide the causal relationship between cataract surgery and AMD. More research is needed to investigate the longitudinal relationship between cataract surgery and AMD in the era of phacoemulsification. Second, excluded participants tended to be in more disadvantaged socioeconomic status and be unhealthier with respect to lifestyles and clinical comorbidities, which might bias the association between cataract surgery and AMD. Third, the self-reported status of cataract surgery might be subjected to the recall bias. Fourthly, we did not have detailed information on the presence of intraocular lens and/or the type of intraocular lens, therefore, we could not examine the effects of aphakia or different types of intraocular lenses on the status of AMD. However, the aphakic cataract surgery is currently scarce. Last but not the least, we could not exclude the possibilities of chance findings. Further studies are needed to corroborate our results.

In conclusion, in this nationally representative population-based study, we found evidence that cataract surgery was significantly associated with any AMD and late AMD, after adjusting for multiple covariates. This may emphasise the need to thoroughly evaluate the retina prior to cataract surgery and inform the potential risk of AMD to cataract patients when recommending cataract surgery. Furthermore, close retinal follow-up after cataract surgery is recommended, particularly in high-risk individuals. Longitudinal studies investigating the risk and progression of AMD after cataract surgery are needed to confirm this finding.

### Table 3 Multiple variable adjusted models for AMD and cataract surgery status

| AMD type          | Right eyes | OR (95% CI) | P value |
|-------------------|------------|-------------|---------|
| Any AMD*          | Reference  |             |         |
| Phakia            |            |             |         |
| Cataract surgery  | 1.36       | (1.03 to 1.81) | 0.033   |
| Early AMD*        | Reference  |             |         |
| Phakia            |            |             |         |
| Cataract surgery  | 1.20       | (0.91 to 1.59) | 0.18    |
| Late AMD†         | Reference  |             |         |
| Phakia            |            |             |         |
| Cataract surgery  | 2.48       | (1.01 to 6.09) | 0.048   |

Boldface indicates statistical significance.

*Adjusted for age, gender, race, marital status, smoking status, obesity, diabetes mellitus, hypertension, chronic kidney disease and self-reported cardiovascular disease.
†Adjusted for age, gender, race, marital status, obesity, hypertension, hyperlipidaemia, chronic kidney disease and self-reported cardiovascular disease.

AMD, age-related macular degeneration.
Table 4  Summary of previous studies investigating the association between cataract surgery and AMD

| Study                  | Country                  | Age range at baseline (y) | Sample size | Design                        | Follow-up | Results (OR/RR/HR, 95% CI)                                                                 | Adjustments                                                                 |
|------------------------|--------------------------|---------------------------|-------------|-------------------------------|-----------|--------------------------------------------------------------------------------------------|----------------------------------------------------------------------------|
| **Significant association**                                      |                          |                           |             |                               |           |                                                                                             |                                                                            |
| Liu et al              | America                  | ≥45                       | 3087        | Population-based cross-sectional | NA        | AMD: 4.6 (2.5 to 8.6)                                                                     | Age, sex and SBP.                                                          |
| Klein et al (BDES)10   | America                  | 43–86                     | 3684 at 5 years; 2764 at 10 years. | Population-based cohort | 10 y      | Late AMD: 3.81 (1.89 to 7.69); GA: 3.18 (1.33 to 7.60); Exudative AMD: 4.31 (1.71 to 10.9); Progression: 1.97 (1.29 to 3.02). | Age, sex, SBP; heavy drinking and smoking, vitamin use.                   |
| Freeman et al (SEE, PVER, BES)11 | America                 | ≥49                       | SEE: 2,520; PVER: 4,774; BES: 4396. | Population-based cross-sectional | NA        | Late AMD: 1.7 (1.1 to 2.6)                                                                 | Age, race, sex and smoking.                                                |
| Wang et al (BMES and BDES)12 | Australia and America | BMES:≥49; BDES:≥43      | 6019        | Population-based cohort | 5 y       | Late AMD: 5.7 (2.4 to 13.6)                                                               | Age, study site, gender, smoking, reticular drusen or pigmentary abnormalities. |
| Cugati et al (BMES)13  | Australia                | ≥49                       | 2335 at 5 years; 1952 at 10 years | Population-based cohort | 10 y      | Late AMD: 3.3 (1.1 to 9.9); NVAMD: 3.4 (1.1 to 10.9).                                      | Baseline age, gender, smoking, and early AMD.                              |
| Kaiserman et al14      | Israel                   | >50                       | 5913:29565  | Retrospective case–control    | 3 y       | PDT: 2.7 (2.4 to 5.7)                                                                     | Age, gender, country of birth, residency, socioeconomic status, hypertension, hyperlipidaemia, DM, congestive heart failure, ischaemic heart disease, and chronic renal failure. |
| Ho et al (Rotterdam Study)15 | Netherlands              | ≥55                       | 6032        | Population-based cohort | 5.7       | Dry late AMD: 3.44 (1.68 to 7.08).                                                    | Age, sex, follow-up time, smoking status and AMD stage at baseline.        |
| Fraser-Bell et al (LALES)16 | America                 | ≥40                       | 6357        | Population-based cross-sectional | NA        | Advanced AMD: 2.8 (1.0 to 7.8); Retinal pigment: 1.6 (1.0 to 1.5); Retinal pigment epithelial degeneration: 2.2 (1.1 to 4.4). | Age, gender and smoking.                                                 |
| Klein et al (BDES)17   | America                  | 43–86                     | 2626 early; 3329 late | Population-based cohort | 20 y      | Late AMD: 1.96 (1.28 to 3.02)                                                            | Age, age, sex, education, smoking, heavy drinking, history of CVD, DM, DBP |

Continued...
| Study          | Country           | Age range at baseline (y) | Sample size | Design                  | Follow-up | Results (OR/RR/HR, 95% CI) | Adjustments                                                                 |
|---------------|-------------------|---------------------------|-------------|-------------------------|-----------|----------------------------|----------------------------------------------------------------------------|
| Saraf et al   | America           | Case: 80.8±6.5 Control: 79.2±9.1 | 39:42       | Retrospective case-control | 1 y       | No more injections, but more susceptibility to cystoid macular oedema or exacerbation of choroidal neovascularisation. | NA                                                                         |
| Ho et al      | China, Taiwan     | ≥49                       | 3465:10 395 | Prospective case-control | 5 y       | NVAMD: 2.68 (1.55 to 4.66). | Geographical location, urbanisation level, monthly income, DM, hypertension, CVD and hyperlipidaemia. |
| McCarty et al (VIP) | Australia | ≥40                       | 4345        | Population-based cross-sectional | NA        | Not significant in the multivariate model. | Age, cigarette smoker, use of ACE inhibitor, use of cholesterol-lowering medication. |
| Armbrcht et al | Scotland         | ≥40                       | 40:43       | Prospective case control  | 1 y       | Only 2% of surgical eyes progressed to wet AMD. | NA                                                                         |
| Holger et al  | Germany           | –                         | 696:202     | Retrospective case-control | 1 y       | NVAMD: 1.30 (0.52 to 3.24). | Age, baseline visual acuity.                                               |
| Chew et al (AREDS) | America      | 55–80                     | 4577        | Cohort study             | 5 y       | NVAMD: 1.08 (0.65 to 1.72); Central GA: 0.98 (0.64 to 1.49). | Baseline age, gender, race, history of smoking.                            |
| Hooper et al  | Australia         | 67–92                     | 27:29       | Randomised controlled trial | 6 m       | 3.7% of CNV in surgical group compared with none in control group (p=1.0). | NA                                                                         |
| Xu et al (BES) | China            | ≥40                       | 3826        | Population-based cross-sectional | NA        | Unilateral surgery was not associated with intereye differences in presence of early (p=0.99) or late AMD (p=0.99). | NA                                                                         |
| Wang et al (CSAMD) | Australia | ≥65                       | 1244        | Clinic-based cohort      | 3 y       | Late AMD: 0.74 (0.23 to 2.36); Early AMD: 1.07 (0.74 to 1.65). | Age, sex, smoking or early AMD lesions at baseline.                        |
| Wang et al (CSAMD) | Australia | ≥65                       | 1057        | Clinic-based cohort      | 4–5 y     | Late AMD: 0.7 (0.4 to 1.2); Early AMD: 0.7 (0.5 to 1.1). | Age, sex, baseline AMD status.                                             |

Table 4 Continued
Table 4  Continued

| Study          | Country | Age range at baseline (y) | Sample size | Design                      | Follow-up | Results (OR/RR/HR, 95% CI) | Adjustments |
|----------------|---------|---------------------------|-------------|-----------------------------|-----------|-----------------------------|-------------|
| Park et al²⁸   | Korea   | ≥40                       | 17987       | Population-based cross-sectional | NA        | Any AMD: 1.02 (0.87 to 1.21); Early AMD: 0.98 (0.82 to 1.16); Late AMD: 1.42 (0.88 to 2.29). | Age, sex, smoking status, income, education level, occupation, DM, dyslipidaemia, overweight, hepatitis B surface antigens and anaemia. |

AMD, age-related macular degeneration; AREDS, age-related eye disease study; BDES, Beaver Dam Eye Study; BES, Beijing Eye Study; BMES, Blue Mountain Eye Study; CNV, choroidal neovascularisation; CSAMD, cataract surgery and age-related macular degeneration; CVD, cardiovascular disease; DBP, diastolic blood pressure; DM, diabetes mellitus; GA, geographic atrophy; LALES, Los Angeles Latino Eye Study; NA, not available; NVAMD, neurovascular age-related macular degeneration; PDT, photodynamic therapy; PVER, Proyecto VER; RR, risk ratio; SBP, systolic blood pressure; SEE, Salisbury eye evaluation; VIP, visual impairment project.

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