**Review Article**

Influence of Metformin on Age Related Macular Degeneration (AMD)

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**Abstract**

Metformin is the most commonly prescribed antihyperglycemic drug as first-line therapy in type II diabetic patients. In recent years, evidence is increasing that metformin has beneficial effects beyond its classical antihyperglycemic way of action. Those effects include anti-inflammation, anti-oxidation, anti-aging, anti-angiogenesis, anti-neoplasia, anti-apoptosis and neuroprotection.

The complex pathophysiology of age-related macular degeneration (AMD) includes age-related changes of the retinal pigment epithelium (RPE) and Bruch’s membrane. An inflammatory and oxidative damage component have also been described. The dry form of late AMD is especially characterized by degeneration of the RPE, Bruch’s membrane, the choriocapillaris and finally loss of the photoreceptors (geographic atrophy) and the wet form of late AMD is characterized by pathological neovascularization. An increasing number of reports about beneficial effects of metformin on AMD have been published in the last years. Several effects of metformin could be linked to the AMPK-pathway. A first prospective trial investigating the effect of metformin on dry AMD is ongoing with estimated results by the end of 2024. In this review, the current knowledge about the association of metformin and AMD is summarized.

**Keywords**: metformin, age related macular degeneration retina, insulin, diabetes, aging, drug therapy, AMPK-pathway

# **INTRODUCTION**

Metformin is one of the most commonly used oral antidiabetic drug. Classically, it is used in non-insulin-dependent type 2 diabetic patients and in most of the cases as the first oral antidiabetic medication. Metformin inhibits the formation of glucose in the liver and improves glucose turnover in the periphery (the muscles) of the body, thereby lowering the blood glucose level (1,2).

There is increasing evidence that metformin may exert several beneficial effects beyond its original antidiabetic function (3–5). In summary, in vitro and in vivo investigations report anti-angiogenic, anti-inflammatory, anti-oxidative, anti-apoptotic, anti-aging and neuroprotective effects of metformin (6,7). Most of these effects also play a crucial role in many retinal diseases such as diabetic retinopathy (DR), age-related macular degeneration (AMD), glaucoma, uveitis or inherited retinal dystrophies as retinitis pigmentosa.

AMD is a vision-threatening disease of the elderly population worldwide with increasing prevalence. Wong et al. calculated an increase from 196 million affected people in 2020 to 288 million affected people in 2040 (8). Together with diabetic retinopathy and glaucoma, AMD accounts for the majority of legal blindness cases in developed countries. In Germany, for example, it is estimated that AMD is responsible for up to 50% of legally blind people. (9)

AMD is a progressive, multi-factorial disease with a complex pathophysiology which is still not fully understood in all its details. The main risk factor is age. It is also known that a history of smoking, hyperlipidemia, ethnicity and a certain genetic disposition as well as inflammatory processes play a role (10,11). different stages: slate stage AMD(12)The early and intermediate stages are characterized by the size of the drusen deposits and by the presence or absence of pigmentary changes. The early and intermediate stage(10)progressThe symptoms include d in the macular regionIf left untreated, fibrosis and permanent vision loss are the consequences (10). (10,11)

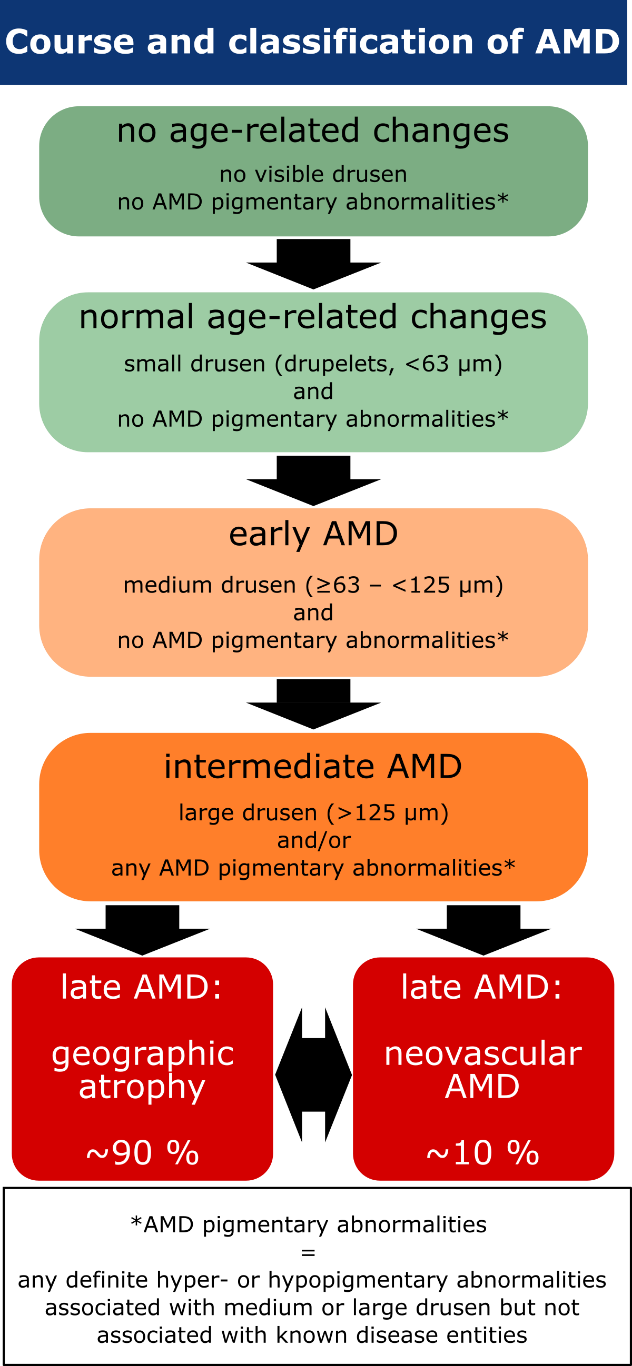


Figure 1: Classification and course of age-relate macular degeneration (AMD). The earliest precursor signs of AMD are small drusen that are classified as normal age-related changes. Early AMD is characterized by the presence of medium drusen but the absence of AMD pigmentary changes which are defined as any hyper- or hypopigmentary abnormality. Intermediate AMD shows large drusen and/or the presence of any AMD pigmentary abnormalities. The late stages of AMD are its two distinct forms: neovascular AMD (wet AMD) and geographic atrophy (dry AMD) with the latter being the more common form. Both forms may merge into one another or be present simultaneously. (modified from Ferris et al. (12))

(10)(10)Approved treatment options are currently only available for late, neovascular AMD. block angiogenic factors that induce. Pathologic neovascularisation leads to(13)(14) The late, dry stage of AMD remains untreatable to date thus efforts are made to find a way to modify the disease. In the last years, an increasing number of scientific publications report on several potential associations of metformin with the course of the disease. This is true for both the development and the treatment of AMD. In this review, we summarize the current knowledge about these associations and the potential underlying (patho)physiological mechanisms.

(15)(15)(16)(17)

# **METHOD**

Systematic literature search was performed using the PubMed library. The search term “metformin age-related macular degeneration” revealed a total of 35 publications (search was performed on July 20th 2022). After screening of titles and abstracts, 22 publications qualified as being suited for the topic of this review. Further database searches with adjusted search terms (metformin AMD, metformin macular degeneration pathways, etc.) did not reveal any further relevant articles.

Additional publications have been included for the introductory part as well as for the background part on the pathophysiology of AMD and on the mode of action of metformin. These publications were identified by direct database search as well as by backward citation searching.

(11)

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(13)(14)

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# **MAIN TEXT**

## **Pathophysiology of AMD**

As mentioned above, the pathophysiology of AMD is complex and several risk factors are associated with this disease. As a neurosensory tissue, the retina, especially the photoreceptor cells, are metabolically highly active. This requires a constant balance between breakdown of metabolic waste products and supply of necessary nutrients, including oxygen. In the healthy retina, the RPE with its tight contact to the photoreceptor cells, Bruch’s membrane and the choroidal vasculature executes this important task (18). The RPE cells form a single cell layer with neighboring cells being connected via tight junctions. This single cell layer is supported by Bruch’s membrane lying underneath. This complex forms the selective diffusion barrier known as the blood-retina-barrier (BRB) which precisely regulates the passage of ions, water, nutrients, proteins and oxygen (19). Any change to this sensitive interface, irrespective of its cause (e.g. age, disease, environmental factors), will affect the precise metabolic balance (11,19,20).

The three main risk factors for AMD are age, environmental risk factors and genetic predisposition. Age itself has an influence on the viability of both the RPE cells and Bruch’s membrane (11). Age has several negative effects on many intracellular structures of the RPE cells, finally leading to changes of RPE metabolism (11). Similarly, Bruch’s membrane suffers from age-related changes, such as thickening and other structural changes that change its permeability. Altogether, these age-related changes have the potential to negatively influence the integrity of the RPE/Bruch’s membrane interface leading to accumulation of debris that ultimately leads to the formation of drusen (11). However, this assumption still has to be confirmed. According to Anderson et al. local inflammation as a response to debris accumulation plays a critical role in the formation of drusen (21). Analyses of the composition of drusen have shown, that they are composed of lipids, polysaccharides, glycosaminoglycans and proteins (21–23). Additionally, many proteins showed oxidative modifications, supporting the hypothesis that oxidative stress is a further contributor to the pathophysiology of AMD (23).

The most important environmental risk factors are smoking and diet. Smoking increases the risk to develop AMD by two- to three-fold. Moreover, there is evidence that there is a dose-dependent association as well as a reversibility in case of quitting smoking (24). Regarding an individual’s diet, healthy forms, e. g. the mediterranean diet, are associated with a reduced risk due to high content of antioxidants and vitamins (25,26). In contrast, high fat or high glucose/fructose diets represent a significant risk factor for AMD. Both direct influences of the nutritional components as well as more indirect influences like dysbiosis of the gut microbiota are thought to be associated with AMD formation. The latter is thought to result in a systemic low-grade inflammation (27).

To date, the largest study of the genome-wide association of AMD revealed 52 gene variants across 34 loci (28). 45 out of 52 were classified as common variants, the remaining 7 as rare variants. Furthermore, the analyses showed that the genetic risk is shared between the neovascular and the non-neovascular form of AMD except for one genetic variant that seems to be exclusive for neovascular AMD (28). Further enrichment analyses narrowed down the following molecular mechanisms that could be affected by the identified gene variants: lipid metabolism, extracellular matrix organization and assembly as well as the complement pathway (28). The possible role of the complement system in the pathophysiology of AMD was recently reviewed by Armento et al. (11). In summary, increasing evidence supports the involvement of the activation of the alternative pathway of the complement system, both in a local fashion as well as on a systemic basis. (11,29)(11)

It is hypothesized that the crucial anatomic site where AMD pathophysiology begins is the complex of RPE cells, Bruch's membrane, and the choroid. In the healthy retina, this complex does not only mediate the precisely regulated exchange of nutrients and metabolic waste products, but it also inhibits the activation of the alternative pathway of the complement system. As soon as the AMD pathophysiology has been triggered through one or more of its risk factors, normal function of the complex is unbalanced. Consequently, both the integrity of RPE cells and Bruch’s membrane becomes more and more impaired. This leads to a cascade of events that disturb retinal homeostasis: accumulation of metabolic end products, oxidative stress, activation of the complement system thereby inducing local inflammation and cell senescence(11,29,30). The exact temporal relationships between degeneration of RPE and photoreceptor cells and collapse of BRB are not yet clear. It is possible that first the BRB collapses and then RPE degenerates, or the other way around. The latter is probably more likely to be the case. Likewise, it is not clear at what point the photoreceptors first become damaged. Possibly already before the RPE degenerates.

## **Metformin**

Metformin is a synthetic derivative of the naturally occurring galegine from the plant *Galega officinalis* (1). Chemically, metformin is a biguanide consisting of two coupled guanidine molecules with some additional substitutes. As a derivative of a naturally occurring molecule, metformin has not been designed to target specific pathways, nor did it go through the regulatory process of preclinical and clinical trials which are mandatory today. After its safety and efficacy had been established, metformin has been used as glucose-lowering agent since the 1950s (1). FDA approval followed in 1994 and since the UK Prospective Diabetes Study in 1997 (UKPDS) had clearly demonstrated the beneficial effects of metformin, it has been recommended as first-line treatment for type 2 diabetic patients (2).

### **Metformin mechanisms of action**

The classical antihyperglycemic function of metformin takes place at multiple sites of action in the body and through multiple molecular mechanisms that have been described in detail elsewhere (1,2). Briefly, its blood glucose-lowering ability is a combination of effects that metformin exerts in the liver, the gastrointestinal tract and in the muscles.

In the liver, gluconeogenesis is downregulated through both AMPK-dependent and -independent signaling pathways. The AMPK-pathway is the cellular energy sensor and regulator of the cell’s energy homeostasis. If the ratio of AMP:ATP increases, the AMPK-pathway induces a switch from ATP-consuming pathways to ATP-generating pathways. This includes downregulation of gluconeogenesis and hence, a reduction of glucose levels (1,2).

In the gastrointestinal tract, metformin is thought to increase glucose uptake and metabolism by colonic enterocytes (1). Moreover, increased glucagon-like peptide-1 receptor (GLP-1 receptor) secretion has been reported in response to metformin. Activation of the GLP-1 receptor results in increased insulin release (2). Finally, metformin seems to be related to shifts in the composition of the gut microbiome, but it remains unclear if and how changes of the gut microbiome lead to glucose-lowering effects (2). It is postulated that a healthier gut microbiome suppresses postprandial hyperglycemia and that levels of inflammatory cytokines are reduced (1).

In skeletal muscles, metformin has been reported to increase insulin-stimulated uptake of glucose. Newer investigations, however, indicate that this effect is of a more secondary nature by the metformin-induced overall improvement of glycemic control and reversal of glucose toxicity (2).

### **Metformin and AMD**

As described above, the pathophysiology of AMD primarily affects the interface of photoreceptors, RPE cells, the choroid and choriocapillaris. The association of metformin with AMD has been investigated in some preclinical trials, some retrospective trials and some systematic reviews and meta-analyses based on the reported mechanisms of action of metformin. Before reporting the results of these trials, we will summarize the proposed mechanisms of action of metformin, that could play a role in its influence on the AMD pathophysiology.

(Figure 2)(31)(31)(32)(2,7)(33)

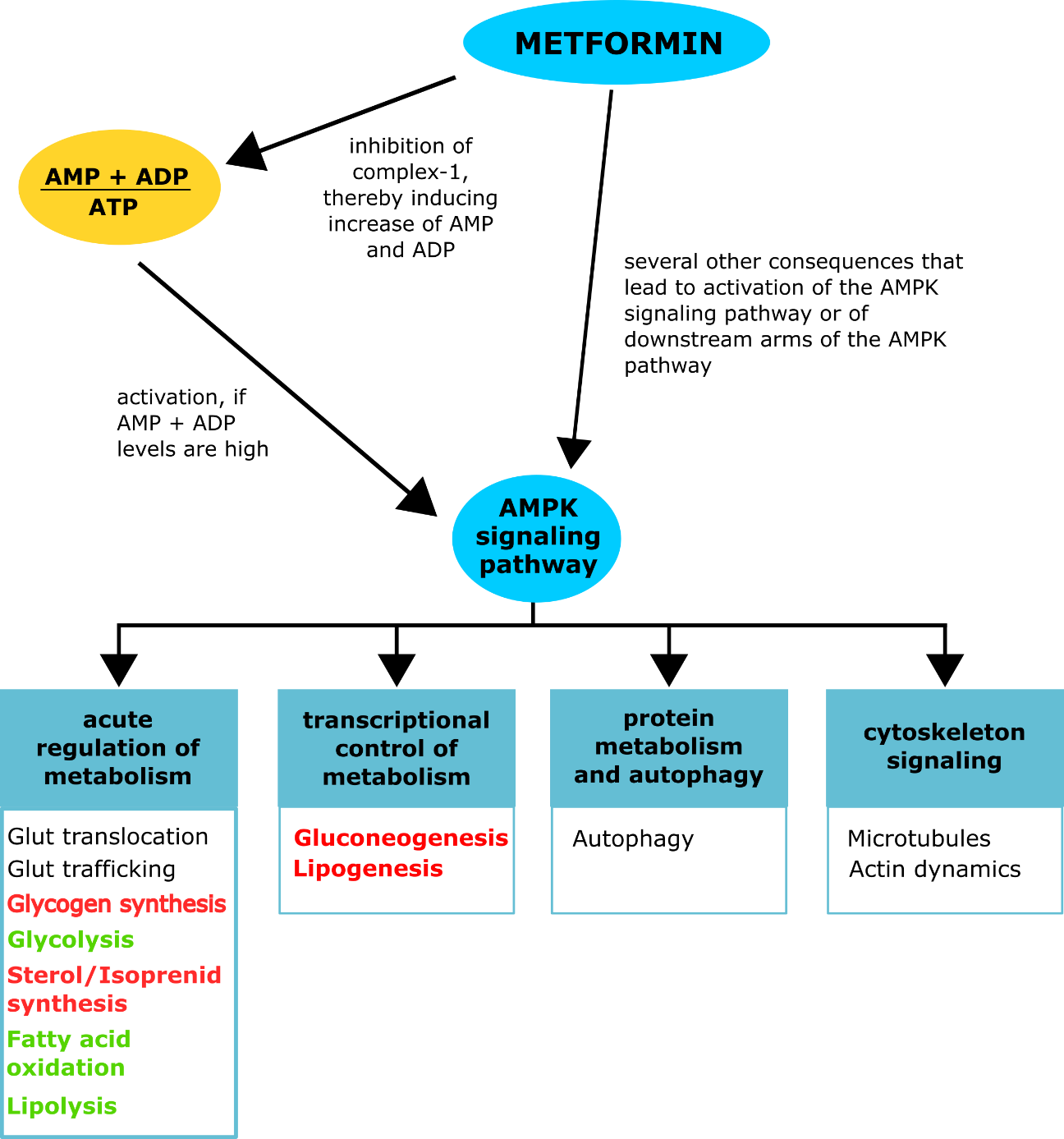


Figure 2: Influence of metformin on the AMPK signaling pathway and consequences of AMPK activation. Without metformin, the AMPK signaling pathway is activated when the cellular levels of AMP and ADP increase. Activation of the pathway lead to a switch from energy consuming metabolism to energy providing metabolism. Metformin has been shown to exert parts of its function through activation of the AMPK-pathway. A confirmed mechanism is that metformin is able to inhibit complex1 of the respiratory chain, thereby inducing accumulation of AMP and ADP. Furthermore, several other, more direct influences of metformin on downstream components of the AMPK-pathway have been reported. AMP = adenosine-monophosphate, ATP = adenosine-triphosphate, ADP = adenosine-diphosphate, AMPK = adenosine-monophosphate dependent kinase, Glut = glucose transporter. Bold red font: inhibition, bold green font: promotion.

### (34)(34)(35) is in line with the finding that fluoxetine, a direct inhibitor of NLRP3, with alikewise (36)**Effects of metformin in preclinical trials**

The group of Ying et al. investigated the effects of metformin in a mouse model of laser-induced CNV as well as in the human umbilical vein endothelial cell (HUVEC) line (37). Mice treated with metformin had significantly smaller CNV lesions with reduced vascular density than the control group. Their experiments with HUVEC cells showed that activin receptor-like kinase 1 (ALK1), a receptor which is essential for vascular development, remodeling and pathological angiogenesis, is inhibited by AMP-activated protein kinase (AMPK) and that metformin is a potent activator of AMPK (37).

(38)(38) The authors did not specify which AMPK-independent pathways are involved in the mode of action of metformin. However, their experiments showed that suppression of NFkB and interleukin-8 by metformin were independent from the AMPK-pathway.

Qu et al. examined the effect of metformin on the human retinal pigment epithelium cell line ARPE-19. Cells were put under oxidative stress via glyoxal-induced cytotoxicity (39). Metformin was able to protect ARPE-19 cells by inhibiting cell death, by reducing intracellular reactive oxygen species (ROS) production, by decreasing the apoptosis rate and by increasing intracellular nitric oxide (NO) levels, an important molecule for maintaining retinal homeostasis (39). A subset of experiments confirmed that metformin influences antioxidant and autophagy pathways to exert its function (39). Similar experiments have been performed by Zhao et al. using two different human pigment epithelium cell lines (40). Their experiments showed that H2O2-induced oxidative damage was attenuated by metformin. Metformin stimulated autophagy via the AMPK-pathway (40).

The *in-vivo* experiments performed by Xu et al. using different mouse models for retinal and photoreceptor degeneration corroborate the results of the above described *in-vitro* experiments (41). Xu et al. used the albino BALB/cJ mouse strain to analyze whether metformin is able to protect light-induced photoreceptor loss. If mice were pretreated with metformin at least 4 days before light damage was induced via 4 h exposure to 4.000 lx bright white fluorescent light, photoreceptor loss was prevented. In a subset of experiments, the group used knockout-mice for the AMPKa1- and AMPKa2-subunit, and showed, that presence of the a2-subunit was crucial for the protective effect of metformin. As the protection by metformin was the same between systemic and local (intravitreal) injection, the authors followed that metformin's protection is based on local influences. Xu et al. used a second mouse model, the Rd10 model for inherited retinal degeneration to analyze the protective effect of metformin. Starting on postnatal day 16, Rd10 mice aggressively lose their rod photoreceptors followed by cone photoreceptor loss. Treatment with metformin starting on postnatal day 13 delayed the loss of both photoreceptor types. Via mitochondrial protein expression experiments, Xu et al. could associate metformin’s protection with an increased metabolic activity. In a third set of experiments, the group injected sodium-iodate into BALB7cJ mice to induce acute oxidative stress to the RPE and to the Photoreceptors. This oxidative stress mimics the early oxidative stress factor of early AMD. If mice were pretreated with metformin, either 30 or 35 mg/kg the RPE and photoreceptor were resistant to the damage in a dose-dependent manner: ~50% and ~90% of cells were protected (41). None of the used mouse models is a perfect model for AMD and such a model does not exist. But each experiment gives insights into relevant parts of the AMD pathophysiology and how metformin possibly intervenes.

(38)(38)

### **Effects of metformin use on AMD in retrospective clinical trials**

Eight retrospective studies have analyzed the association of metformin use with AMD, of which one was a cross-sectional study (42), four were cohort studies (43–46), two were case-control studies (47,48), and one was a nested case-control study (49). Five studies exclusively determined the association of metformin use and the risk of developing AMD in diabetic patients (42–46), whereas three studies included broader patient groups according to their cohort definition (47–49). One study examined the association of metformin use with dry AMD only (46), while the remaining studies considered all forms of AMD or did not further specify. All studies took into account possible confounders like age, sex, ethnicity, smoking status, insurance status, other oral (antidiabetic) medications, insulin use, cardiovascular disease, hypertension, hyperlipidemia, obesity, BMI, HbA1c, kidney disease, or charlson comorbidity index as far as data were available.

Stewart et al. performed a cross-sectional study using the electronical medical record database of the University of California, San Francisco (42). They included 3.120 diabetic patients who had documented ophthalmologic examinations and a documented metformin use prior to or at their first documented ophthalmologic exam. The outcome of interest was a diagnosis of either non-neovascular or neovascular AMD at the first ophthalmologic exam. Using propensity score-weighted logistic regression models, Stewart et al. found that metformin use was significantly associated with a reduced odd ratio (OR) to develop AMD (OR 0.70, 95% confidence interval, p-value 0.003). The association was even stronger, when analyzing non-neovascular AMD alone (OR 0.59, 95% CI, p-value < 0.001). All other antidiabetic drugs studied showed no association. Limitations of this study are the retrospective nature, the relatively small sample size, the exclusion of drusen as an early stage of AMD, because authors found the diagnosis of drusen to be unreliable, and missing information about the duration of metformin use.

Chen et al. investigated the association of metformin use and the risk of AMD in a cohort study with type 2 diabetic patients (43). They included 68.205 patients who had a diagnosis of type 2 diabetes mellitus during the study period. Patients were followed up to identify the onset of AMD (unspecified, non-exudative or exudative). The main independent variable was use of metformin, which was true for 66.7% of the identified patients. Adjusted hazard ratios (HRs) were obtained via multivariate Cox regression analyses. Patients taking metformin had a significantly lower HR to develop AMD than metformin non-users (HR 0.54, 95% Ci, p-value <0.0001). Chen et al. also calculated HRs for the duration and cumulative dose of metformin and their association with the development of AMD and found that both significantly lowered AMD risk. Limitations of this study are the retrospective nature, some missing details in the database like smoking status, diet, and laboratory values.

Another cohort study was performed by Jiang et al. (44). The group reviewed medical records of the ophthalmology department of the China-Japan Friendship Hospital in Bejing, China. 324 patients with a diagnosis of diabetes mellitus type 2, lasting for at least 10 years, were identified and followed-up over 5 years. Patients were excluded if they had a diagnosis of AMD before diagnosis of diabetes. AMD was graded into early and late-stage AMD. Metformin users and non-users were compared using X2 test and multivariate logistic regression models were used to characterize the influence of confounders. AMD occurrence in the metformin group was 15.8% and 45.2% in the metformin non-users (p<0.0001), thus patients taking metformin had significantly less risk to develop any AMD. Subgroup analysis revealed, that metformin use only influenced development of early AMD and not late AMD. Further analysis showed that both duration of metformin use and cumulative metformin dose were associated with significantly lower risks to develop any and early but not late AMD. The retrospective design, the small sample size, and missing data on some important confounders are the limitations of this study.

Gokhale et al. performed a further cohort study investigating the influence of metformin on the risk of AMD in patients with type 2 diabetes (45). The group identified 173.689 patients with newly diagnosed type 2 diabetes from the United Kingdom IQVIA Medical Research Data. Patients were excluded if they had AMD diagnosis before diabetes diagnosis and if they had no prescription for antidiabetic medication. 89% of the identified patient had a prescription for metformin alone or in combination with other antidiabetic drugs. The control group had any medication except of metformin. The outcome of interest was a diagnosis of AMD during the study period. HRs were defined in a time-dependent manner using extended Cox proportional hazard models. For the time-dependent analysis the follow-up intervals were set to 3 months. AMD occurred in 3111 (1,8%) of the patients. Gokhale et al. did not find an association between metformin and the development of AMD. This finding was independent from the use of other antidiabetic drugs as well as from the duration of diabetes and the duration of metformin use. Limitations of this study include the retrospective design, the missing differentiation between AMD stages (early, late) that could for example mask findings if metformin was only protective for certain AMD stages.

The group of Eton et al. investigated the association of metformin and dry AMD only (46). In their cohort study, they included patients with a diagnosis of diabetes mellitus and sufficient follow-up visits. Furthermore, Eton et al. distinguished between current and historical use of metformin. Current metformin use was defined as metformin use during the study period, historical metformin use was based on any metformin use before the patients’ enrollment date (defined as aged 55 years or more, a diagnosis of diabetes mellitus and at least two years of follow up data). Current metformin use was associated with a small, but significant increased HR to develop dry AMD (HR 1.08; 95% CI, p<0.0001). Historical metformin use, however, showed a protective effect (HR 0.95; 95% CI, p=0.002). The analysis of the cumulative dose of metformin revealed slightly decreased HRs for cumulative doses below 720,000 mg, but slightly increased HRs for cumulative doses above 720,000 mg. Overall, the study by Eton et al. showed conflicting results for the effect of metformin on the development of dry AMD. Study limitations include the retrospective design, potentially the restriction to dry AMD only and a probably observation bias.

Lee et al. used a different study design, a nested case-control study, and they also had a broader definition for the study eligibility as they not only included patients with a diagnosis of diabetes mellitus type 1 and 2 but also patients with a diagnosis of cardiovascular disease (49). Above that, they were not only interested in the effect of metformin, but also in the effects of statins, angiotensin-converting enzyme (ACE)-inhibitors and angiotensin II receptor blockers on AMD. 2330 patients developed AMD during the study period. For each case 10 controls were matched by sex, age and cohort entry date, leading to a control group of 23,278 patients. Study outcomes were, that none of the investigated drugs had a protective effect on the development of AMD. These findings were independent of the duration of drug use. The nested case-control design overcomes some of the disadvantages of traditional case-control studies, as for example the reduction of selection bias. The retrospective design and its disadvantages remain, and sample sizes were relatively small.

Two case-control studies investigated the effect of metformin on AMD independent from a diagnosis of diabetes (47,48). However, both studies examined diabetic patients separately as subgroups of the initial total study cohort. Cases were defined as patients who had a diagnosis of AMD during the study period. Brown et al. included patients with all types of AMD (non-exudative, exudative, or unspecified), controls had no AMD and were propensity score matched using age, charlson comorbidity index (CCI), hypertension and anemia as matching variables (48). They found that metformin was associated with statistically significant decreased odds of developing AMD (OR 0.58; 95% CI, p=0.0005). Other diabetic and non-diabetic medications showed no association with AMD. The subgroup analysis of diabetic patients taking metformin versus non-metformin users showed that metformin was significantly associated with decreased odds of developing AMD in univariate and multivariate logistic regression (OR 0.68; 95% Ci, p=0.002 and OR 0.7; 95% CI, p=0,043). Blitzer et al. defined their study cohort as patients with newly diagnosed AMD during the study period and powered their study to detect ORs of 0.95 with 90% power in a subgroup of diabetic patients (47). Controls were selected 1:1 and matched based on age, anemia, hypertension, region and CCI score. The effects of diabetes were tested after control matching. Metformin use was similar in case and control group (12,8% and 13,0%). Use of any metformin was significantly associated with decreased odds of developing AMD (OR 0.94; 95% CI, p<0.001). In addition, it was found that low to moderate total metformin doses had a dose-dependent effect, while there was no association between high metformin (> 1080 g cumulative dose) doses and AMD. The subgroup analyses of diabetic patients showed similar results. Metformin use significantly decreased the odds for developing AMD (OR 0.95; 95% CI, p>0.001) and again a dose-dependent effect for low to medium cumulative metformin doses was found.

In summary, five out of eight retrospective studies found associations of metformin with decreased odds of developing AMD (42–44,47,48), one study found conflicting associations (46) and two studies report no association of metformin use with the development of AMD (45,49). Three studies found positive associations with either the duration of metformin use or dose-dependent effects (43,44,47), while one study did not detect an association with longer metformin use (49). A meta-analysis of Romdhoniyyah et al. over five retrospective trials did not find a significant association between metformin use and the risk to develop AMD (3). (3)

The main limitation of all retrospective studies is that they can only detect associations but cannot determine causal relationships. The latter is only possible in the context of prospective trials. In addition, retrospective trials are prone to other limitations such as selection bias, recall bias, loss to follow-up, and confounding factors (50). Nevertheless, the majority of the described retrospective analyses found that metformin was associated with decreased odds to develop AMD. Selection bias is especially small for cohort studies like those of Chen et al., Jiang et al., Gokhale et al., and Eton et al. (43–46,50). All eight studies considered confounding factors and comorbidities in their analyses. The limitation of loss to follow-up was reduced by adjusting the eligibility criteria and only patients for whom sufficient follow-up visits were available were allowed to enter the study cohorts. As metformin is predominantly described to type 2 diabetic patients, five studies exclusively investigated the effect of metformin on AMD in diabetic patients. Three studies included broader patient groups. Two of them found that metformin decreased the odds of developing AMD independently from a diagnosis of diabetes. This suggests that diabetes itself probably has little influence on the development of AMD.

**Prospective clinical trials**

(GA)(51)A planned population of 186 subjects will be stratified 1:1 into a treatment and an observation group. The treatment group will be assigned to oral metformin for 18 months. At an additional follow-up visit at month 24, the progression of geographic atrophy will be measured and compared between groups. The primary outcome measures are the change in area of GA or drusen growth. Secondary outcome measures include best corrected and low-luminance visual acuity, ocular and systemic safety of metformin use and score changes of the National Eye Institute Visual Function Questionnaire. Subjects with type 1 or 2 diabetes are excluded from the study as well as subjects that are already taking metformin for other reasons. Results of this study may for the first time confirm the trending results of the previously only retrospective trials.

**Limitations of metformin use**

Beyond all the reported beneficial properties of metformin, there are also some disadvantages associated with the use of metformin that should be taken into account before using metformin as a "cure it all medication". Reported disadvantages include vitamin B12 deficiency, increased risk of lactic acidosis, and alteration of 745 proteins with uncertain consequences (7). In addition, metformin is known to have various gastrointestinal side effects.

Furthermore, a study of Ebeling et al. that analyzed the influence of metformin on individual patient-derived RPE cell-lines indicated that the effect of metformin was not uniform across all patients. The group suggests that patient specific responsiveness to metformin should be taken into account before prescription and that approaches towards personalized medicine are necessary (52).

(53)(53)

### (54)(54)

(31)(31)(32)(2,7)(33)

(34)(34)(35)(36)

# **CONCLUSION**

Evidence is increasing that metformin, the most commonly prescribed oral antihyperglycemic drug, influences a variety of physiological functions besides its classical glucose-lowering effect. Essentially, this includes anti-inflammatory, anti-angiogenic, anti-oxidative, anti-apoptotic, neuroprotective and anti-ageing effects. In addition, it has been shown that metformin is able to enhance the effect of anti-VEGF agents in the treatment of diabetic macular edema via regulation of the AMPK/mTOR signaling pathway (53). As for DME, anti-VEGF agents are the standard of care treatment for late stage neovascular AMD. that are treated with anti-VEGFs If a similar effect is true for nAMD treatment, metformin use could probably be an option for anti-VEGF non-responders to increase their benefit from anti-VEGF therapy. (53)(53)

(7)(52)

The ongoing prospective trial about the effect of metformin on the progression of geographic atrophy could deliver first results for this subgroup of late-stage AMD patients. In the future, more prospective trials are needed to confirm in more detail how the beneficial effects of metformin influence the pathophysiology of AMD and if metformin qualifies as a treatment option in patients with a diagnosis of AMD or even as a protective therapy before any AMD diagnosis. Additionally, prospective trials should not only concentrate on late-stage dry AMD but consider all AMD stages. Jiang et al. found for example, that especially the early stage of AMD was associated with a beneficial effect of metformin. Finally, prospective trials should consider patients with and without a diagnosis of diabetes to rule out possible confounding effects of the diabetic disease. From the patients' point of view it would be highly desirable if the progression of AMD could be stopped or substantially slowed down, irrespective of a diagnosis of diabetes or not. (51)

# Author’s contributions

The authors contributed equally to the article.

# Conflicts of interest

I declare that I have no conflicts of interest.

# Ethical approval and consent to participate

Not applicable.

# Consent for publication

Not applicable.

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