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**Keeping an eye on the diabetic foot: The connection between diabetic eye disease and wound healing in the lower extremity**

Ramsey DJ *et al*. The diabetic foot and eye connection

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**Abstract**

Diabetic eye disease is strongly associated with the development of diabetic foot ulcers (DFUs). DFUs are a common and significant complication of diabetes mellitus (DM) that arise from a combination of micro- and macrovascular compromise. Hyperglycemia and associated metabolic dysfunction in DM lead to impaired wound healing, immune dysregulation, peripheral vascular disease, and diabetic neuropathy that predisposes the lower extremities to repetitive injury and progressive tissue damage that may ultimately necessitate amputation. Diabetic retinopathy (DR) is caused by cumulative damage to the retinal microvasculature from hyperglycemia and other diabetes-associated factors. The severity of DR is closely associated with the development of DFUs and the need for lower extremity revascularization procedures and/or amputation. Like the lower extremity, the eye may also suffer end-organ damage from macrovascular compromise in the form of cranial neuropathies that impair its motility, cause optic neuropathy, or result in partial or complete blindness. Additionally, poor perfusion of the eye can cause ischemic retinopathy leading to the development of proliferative diabetic retinopathy or neovascular glaucoma, both serious, vision-threatening conditions. Finally, diabetic corneal ulcers and DFUs share many aspects of impaired wound healing resulting from neurovascular, sensory, and immunologic compromise. Notably, alterations in serum biomarkers, such as hemoglobin A1c, ceruloplasmin, creatinine, low-density lipoprotein, and high-density lipoprotein, are associated with both DR and DFUs. Monitoring these parameters can aid in prognosticating long-term outcomes and shed light on shared pathogenic mechanisms that lead to end-organ damage. The frequent co-occurrence of diabetic eye and foot problems mandate that patients affected by either condition undergo reciprocal comprehensive eye and foot evaluations in addition to optimizing diabetes management.

**Key Words:** Foot ulcer; Diabetic; Wound healing; Diabetes complications; Amputation; Diabetic retinopathy; Corneal ulcer

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**Core Tip:** This review explores the epidemiological and pathophysiological interconnections between diabetic foot and eye disease, especially the shared mechanisms that impact wound healing. Since diabetic foot and eye problems are often concurrent, it is imperative that patients affected by one or the other condition promptly undergo reciprocal examinations to reduce the risk of further complications. The best outcomes for patients with diabetic foot and eye disease are achieved by a team-based strategy that incorporates regular examinations, often performed by specialists, provides preventative health education, and delivers effective long-term management of the underlying diabetes and its associated metabolic consequences.

**INTRODUCTION**

An estimated 131 million people worldwide have lower extremity complications related to diabetes mellitus (DM), such as diabetic foot ulcers (DFUs), peripheral vascular disease (PVD), neuropathy, and amputations[1,2]. Similarly, an estimated 103 million people worldwide have diabetic eye disease, including nearly one million people aged 50 and older who are blind from diabetic retinopathy (DR)[3,4]. The frequent co-occurrence of diabetic eye and foot problems makes it imperative for patients affected by either condition to promptly undergo reciprocal examinations to reduce the risk of further complications (Figure 1). It is essential that individuals who have DM and the clinicians who care for them understand the likelihood of this association. With the worldwide prevalence of DM increasing because of changes in diet and lifestyle, aging of the population, and the ability of individuals to live longer with the disease, the need for well-informed clinicians has never been greater[3].

The connection between diabetic eye and foot problems is related, in part, to shared risk factors. In particular, the duration of DM and level of glycemic control as reflected by hemoglobin A1c (HbA1c) level strongly govern both the rate of onset and severity of diabetic foot disease[5,6] and DR[7]. Molecular biomarkers, particularly ceruloplasmin, have been demonstrated to be elevated in people with DM[8].Other risk factors, such as age, male gender, race and ethnicity, smoking, insulin use, type of diabetes, and individual comorbid factors such as hypertension, elevated low-density lipoprotein (LDL), decreased high-density lipoprotein (HDL), coronary artery disease, cerebral vascular disease, PVD, neuropathy, and nephropathy, have been assessed, but not all studies agree on which of these risk factors affect the incidence or progression of these diabetic complications[5]. Some of this variation may possibly be ascribed to differences in DM care, improvements in treatment over time, and other less well-defined differences between individual populations studied. This paper reviews the shared pathogenic mechanisms underlying these conditions and the importance of comprehensive diabetes care to reduce morbidity and prevent disability.

**Diabetes-Associated Lower Extremity Complications and their Ocular Parallels**

***DFUs***

Individuals with DM are at a significantly increased risk of developing DFUs.DFUs are full-thickness wounds that penetrate the dermis (the deep vascular and collagenous inner layer of the skin) and are located below the ankle in patients with both type 1 and type 2 DM (Figure 2)[1,2]. DFUs arise from a combination of micro- and macrovascular compromise related to hyperglycemia and associated metabolic dysfunction that causes impaired growth and wound healing, immune dysregulation, and PVD[9]. In addition, the loss of protective sensation and proprioception caused by diabetic neuropathy and vision loss from diabetic eye disease predisposes patients to repetitive lower extremity trauma, with DFUs a common complication, especially among older adults[10,11]. Risk factors for DFUs include age, deformity or prior ulceration, repetitive trauma, sensory and autonomic neuropathy, peripheral arterial disease (PAD), and infection[12].Up to one third of patients with DM will be affected by a DFU in their lifetime[13]. DFUs are associated with a 10- to 20-fold increased risk of amputation[1] and have a one-year mortality rate as high as 5%[14]. As many as 20% of DFUs remain unhealed after one year of treatment,with unhealed ulcers posing a risk for infections, gangrene, amputation, and even death[15,16].

There is a strong association between the development of DFUs and DR (Figure 2)[5]. Depending on the population studied, most individuals affected by DFUs also have DR[6,17-19], and those with DR are two to four times more likely to have DFUs or more serious forms of diabetic foot disease (Table 1)[20-22].Even more concerning is the strong association between DFUs and proliferative diabetic retinopathy (PDR), with 31% to 55% of individuals having this more severe stage of DR (see below)[23]. Furthermore, patients with nonproliferative diabetic retinopathy (NPDR) who develop comorbid non-healing DFUs have a greater than 50% increased risk of progressing to PDR relative to those without this condition[24]. Finally, diabetic keratopathy is an important ocular parallel of DFUs. It is a disruption of normal corneal wound healing and loss of protective mechanisms of corneal sensation and aqueous tear production. These aberrations create an ideal environment for persistent corneal epithelial defects, microbial infection, and ulceration. Around half of patients with DM are affected by this condition[25]. The pathophysiology accounting for diabetic structural and functional alterations in the cornea is discussed in depth below.

***Microvascular* *complications***

Microvascular dysfunction in the lower extremity contributes to impaired function and impedes wound healing, which promotes the development of DFUs. Damage to endothelial cells from chronic hyperglycemia, oxidative stress-induced injury, generation of advanced glycation end-products, increased polyol flux regulated by aldose reductase, activation of protein kinase C (PKC), and other pro-inflammatory processes from immune dysregulation cumulatively disrupt normal blood flow and affect vascular permeability[26]. In its most severe form, this compromise of the microvasculature leads to ischemia and a relative hypoxic state in the involved tissue. As a result, there is increased expression of hypoxia-inducible factor-1 (HIF-1) leading to the production of vascular endothelial growth factor (VEGF), a protein principally responsible for restorative angiogenesis[27]. However, in DM there are disturbances in cytokine growth factor expression and locally decreased concentrations of VEGF, which render the lower extremity vulnerable to poor wound healing[9]. Failure of the microvasculature also contributes to peripheral neuropathy and local immune dysfunction, including impaired cellular response, cytokine expression, and vascular tone[28].

In the eye, these same pathways driven by hyperglycemia and other diabetes-associated factors lead to progressive damage to the retinal microvasculature and cause the development of DR[29]. DR develops in roughly one quarter of patients with DM, with the prevalence being highest in Africa (36%) and lowest in South and Central America (13%)[3,30]. Initially, the disease manifests as clinically detectable changes in the retinal vasculature, including the development of microaneurysms and loss of capillaries which are the hallmarks of early NPDR[31]. As the disease progresses, the production of VEGF and other diabetes-associated factors promotes further dysfunction, vascular leakage, and bleeding (dot-blot hemorrhages)[32]. At this stage, visual acuity is increasingly likely to be affected and is often further limited by swelling in the center of the retina, known as diabetic macular edema (DME)[33]. The development of neovascularization on the optic nerve or at locations in the peripheral retina signifies the progression to PDR. This is the most vision-threatening complications of the disease also primarily driven by the abnormal expression of VEGF[29,30,34]. These fragile new vessels, which grow into the vitreous cavity and along the inner retinal surface, often bleed, causing vitreous hemorrhages, traction, or retinal detachment and thereby impair vision[3,32]. Finally, excessive expression of VEGF may also affect the anterior segment of the eye by causing neovascularization on the iris and ciliary body. When the growth of this fibrovascular tissue extends to the anterior chamber angle it may block outflow of aqueous humor through the trabecular meshwork causing eye pressure to rise to levels capable of damaging the optic nerve in a disease process known as neovascular glaucoma[4,32]. When left unaddressed, irreversible blindness results.

Clinical examination supplemented by diagnostic color fundus photography and fluorescein angiography are the mainstays for staging DR; however, emerging modalities including ultrawide-angle imaging and optical coherence tomography and angiography increasingly allow clinicians to directly and noninvasively visualize the diseased retina and its microvasculature[31]. Advances in therapeutic modalities, such as intraocular injection of agents that target VEGF, steroids that target inflammation, and panretinal laser photocoagulation, have improved clinical outcomes for patients with DR[33-36]. However, effective long-term management is largely dependent upon regular follow-up care. Patients who fail to return for care are more likely to suffer vision loss[37,38].

The intraocular administration of agents that target VEGF are now the most common treatments for DR and DME[29,36]. Thankfully these agents are very unlikely to negatively impact wound healing in the lower extremity, especially at the doses employed to treat eye disease[39].Similarly, intraocular corticosteroids, such as dexamethasone, and intravitreal steroid implants utilized to treat DME have been found to have no detectable influence on HbA1c or renal function[40,41]. However, when larger doses are administered as subconjunctival or peribulbar injections, some patients can experience elevations of blood glucose, similar to that observed with oral and intravenous administration of corticosteroids[42]. Finally, topical steroid drops have only very rarely been associated with endocrinological side effects in case reports[43].

***Wound healing***

Wound healing in the lower extremity requires coordinated cellular responses that cause an organized release of growth factors and cytokines. Under normal conditions, when an injury occurs, multiple cell types, including macrophages, fibroblasts, and epithelial cells, release VEGF and other cytokines in response to local ischemia caused by the wound[44]. However, in patients with DM, disturbances in cytokine and growth factor expression, including fibroblast growth factor, insulin-like growth factor, platelet derived growth factor, and VEGF, among others, lead to a condition that subsequently permits prolonged hypoxia[9,45]. Additionally, keratinocytes and fibroblasts in DFUs have demonstrated attenuated cellular migration, proliferation, and protein synthesis, resulting in impaired re-epithelialization which further exacerbates the oxygen-restricted wound[46]. Moreover, hyperglycemic states reduce the stability and function of HIF-1, which further impairs the wound healing response as a downstream consequence of sustained hypoxia[9]. Increased free radical damage is also a known causative factor in impaired wound healing in patients with DM. Inappropriately elevated concentrations of reactive oxygen species (ROS) and impaired antioxidant enzyme activity can cause nerve damage and directly contribute to the progression of peripheral neuropathy[47].

Individuals with DM also have abnormal wound healing pathways in the eye. Notably, corneal thinning is thought to be the earliest detectable pathological manifestation of DM in the eye[48].Diabetic keratopathy leads to persistent corneal epithelial defects and neurotrophic corneal ulcers that respond poorly to treatments applied in the hyperglycemic environment[49]. Abnormalities in corneal cell morphology, varied number and disorganization of epithelial cell layers, impaired cellular migration, reduced endothelial cell number, and accumulation of acellular debris all contribute to poor wound healing[50]. Sectorial thinning, bullae, and persistent corneal epithelial defects from diabetic keratopathy often lead to corneal ulcers, scarring, and reactive neovascularization, which cause decreased visual acuity or permanent vision loss[50,51]. Although the cornea itself is avascular and ischemia does not play a significant role in diabetic keratopathy, wound healing in the cornea, like in the lower extremity, requires highly structured cellular processes which are impacted by hyperglycemia. These involve proliferation and migration of epithelial cells, fibroblasts, and the expression of numerous growth factors, including transforming growth factor beta, epidermal growth factor, insulin-like growth factor, and platelet derived growth factor[51]. Finally, diabetes-associated hyperglycemia may also impair vision by accelerating the progression of diabetic cataract and impact the health of the lens epithelium[52].

Most treatments for diabetic foot and eye problems are applied locally, but some treatments to aid the lower extremity may have theoretical consequences on the eye, and vice versa. Several adjuvant therapies have been found to reduce DFU healing times and amputation rates, including non-surgical debridement agents, topical dressings and agents, negative pressure wound therapy, oxygen therapies, acellular bioproducts, and human growth factors[53]. Oxygen is required for almost every step of the wound healing, affecting cell proliferation, collagen synthesis, and re-epithelialization, as well as immunologic defense against bacteria and other pathogens[54]. Oxygen may be delivered in the form of local, hyperbaric, or supplemental inspired oxygen therapy. Hyperbaric oxygen therapy has proven to be particularly useful in managing chronic, non-healing DFUs, especially in the relatively ischemic diabetic foot, albeit at a high financial cost[55]. Patients have been observed to have increased tissue concentrations of VEGF after completing hyperbaric therapy sessions; this has been attributed to the sharp decline of relative oxygen concentration once a session is completed[56]. As previously mentioned, the presence of VEGF is the primary driver of DR, so there is a theoretical risk that systemic or local oxygen therapy could exacerbate this condition. However, empiric evidence does not suggest that oxygen therapy is harmful to the diabetic eye[23]. It has even been reported that patients with concurrent DR have benefitted from the administration of hyperbaric oxygen therapy through supranormal levels of oxygen delivered to the retina[57]. Nevertheless, it remains essential that the status of DR is assessed and regularly monitored in any patient undergoing oxygen therapy for DFUs.

Many growth factors that have been identified as integral to wound healing are also potential therapeutic targets. In the diabetic foot, among the most promising are hydrogels which contain recombinant PDGF, approved by the United States Food and Drug Administration for topical administration having demonstrated improved rates of DFU healing in randomized clinical trials[58]. In the diabetic cornea, recombinant human nerve growth factor (NGF), epidermal growth factor, and metalloprotease inhibitors have demonstrated some success in trials for the treatment of diabetic keratopathy[59]. Of note, the opioid antagonist naltrexone has been demonstrated to improve wound healing, corneal surface sensitivity, and tear secretion in diabetic animal models[60,61]. The future will also likely include gene- and cell-based therapies to accelerate wound healing, including in DFUs and diabetic cornea[62,63].

***Diabetic neuropathy***

Approximately half of adults with DM will be affected by peripheral neuropathy in their lifetime[64]. Peripheral neuropathy typically begins with diminution or loss of protective sensation. In addition, loss of proprioception contributes to injuries and falls[11].Moreover, autonomic dysregulation in the foot may contribute to impaired cutaneous blood flow, sweating dysfunction, and loss of vascular tone that compromise integument integrity and wound healing[65]. Lower extremity deformities may also occur, such as hammer toes or claw toes, which are associated with loss of function[11]. Finally, delays in the identification of accidental and iatrogenic injuries because of reduced sensation may cause patients to fail to seek care in a timely fashion and increase the risk for infections[64].

As mentioned above, chronic hyperglycemia from DM causes microangiopathic changes. In the case of diabetic neuropathy, hyperglycemia may affect the endoneurial microvasculature by directly reducing perfusion and impairing nerve function[66]. Many of the same cellular and biochemical mechanisms linked to chronic hyperglycemia injure the peripheral nerves, including increased glycolytic processes producing oxidative stress, generation of advanced glycation end-products, increased polyol flux regulated by aldose reductase, PKC activation, and other pro-inflammatory processes from immune dysregulation[67]. Damage to mitochondria also plays an important role in the pathogenesis of diabetic neuropathy and contributes to nerve dysfunction, cell death, and loss of neurotrophic support provided by neurotrophin-3 and NGF[68]. Patients who develop PAD also have more severe diabetic neuropathy (see below)[64].

The cornea is the most densely innervated tissue of the human body and is 100 times more sensitive than skin[69], but this declines with age[70] and is further reduced by DM[71]. The loss of protective sensation of the diabetic cornea impacts various homeostatic functions, such as blinking, aqueous tear production, and the release of growth factors[51,52]. As a result, the incidence of dry eye disease and the need for artificial tears is increased among patients with DM, particularly among those with worse diabetes-related outcomes[72]. A recent meta-analysis estimated that DM conferred 30% increased odds for dry eye syndrome[58]. Dry eye disease and DR are also associated with each other[73]. These changes result in neurotrophic keratopathy marked by persistent epithelial defects and chronic erosions that may develop into corneal ulcers, corneal scarring, and neovascularization, all of which contribute to visual dysfunction[74] and predispose patients to infectious keratitis[75]. The recent application of *in vivo* confocal microscopy has allowed for visualization of diabetes-associated structural changes in the nerves of the corneal epithelium, including nerve thickening[76] and decreased nerve length and density[77]. Anterior segment optical coherence tomography is another emerging diagnostic modality used to evaluate and manage diabetic keratopathy by enabling the direct visualization of the cornea structure and nerves[50,71].

Diabetes-associated hyperglycemia has also been shown to cause direct injury to the neuronal retina, leading to thinning of the nerve fiber layer from the loss of ganglion cells and the death of other retinal neurons, including photoreceptors[78]. This may lead to decreased visual function, impaired contrast sensitivity, and diminished night vision[29,32]. Finally, the eye may also be suddenly and directly affected by diabetic cranial neuropathies, manifesting as double vision from ophthalmoplegia, which is the paralysis of the muscles that move the eye (see below).

Current recommendations for the management of painful diabetic neuropathy include gabapentinoids, serotonin and norepinephrine reuptake inhibitors (SNRIs), and tricyclic antidepressants (TCAs)[79].Gabapentin is a well-tolerated medication from an ophthalmic standpoint; its most common adverse effect is a reversible nystagmus[80]. SNRIs, such as venlafaxine, have been associated with acute angle closure glaucoma in some case reports[81], along with increased cataract development[82]. Finally, TCAs are associated with blurred vision in up to one third of patients, likely due to the anticholinergic action of these drugs[83]. These side effects further emphasize the importance of communication and collaboration with ophthalmologists when treating diabetes-associated complications of the lower extremity.

***Macrovascular complications***

Common macrovascular complications of DM that affect the lower extremity include PAD and chronic venous insufficiency (CVI), which may lead to lower extremity amputation (Figure 1)[84].DM induces and accelerates the development of atherosclerosis *via* multiple mechanisms that include metabolic derangements, smooth muscle dysfunction, oxidative stress, potentiated platelet function, increased coagulability, and chronic inflammation[85]. The availability of the potent vasodilator nitrous oxide, which is produced in the endothelium and is a primary mediator in local vascular endothelial tone, is reduced in hyperglycemic states; DM also promotes the production of endothelin-1, which indirectly increases vasoconstriction and vascular smooth muscle hypertrophy[86]. The result may be an overt occlusion, sometimes acutely when a thrombus forms, or when increasingly stenotic vessels result in reduced perfusion[86]. Patients with co-existing severe PAD are also more likely to have CVI[84], which contributes to poor wound healing by increasing hydrostatic pressure in the lower extremity, thereby promoting wound exudation[87].

While not directly a macrovascular complication, it is important to recognize that DR is strongly associated with lower-extremity PAD. Patients with DR have an approximately two-fold increase in the need for lower-limb revascularization and a five-fold increase in lower-limb amputation[88]. Patients with PAD benefit from additional medical management and risk factor modification for atherosclerotic disease. In addition to optimizing diabetes control, this includes counseling about smoking cessation, antiplatelet and statin therapies, as well as blood pressure control[89]. Exercise also plays a fundamental role in the treatment of PAD, leading to reductions in pain and improvement in functional capacity[89]. The clinical benefit of newer medications on amputation prevention remains uncertain.

In the eye, DM-associated macrovascular disease can manifest as an ocular ischemic syndrome (OIS), a rare, but vision-threatening condition associated with severe carotid artery occlusive disease that leads to ocular hypoperfusion[90]. Like PAD, atherosclerosis affecting the vessels supplying the eye is the main cause of the disease, and most patients with OIS have a diagnosis of DM[91]. Patients typically report dull eye or periorbital pain associated with gradual vision loss as the retina experiences progressive ischemia. Consequently, VEGF levels rise, which may cause neovascular glaucoma in the anterior segment and reduce the final visual potential; neovascularization can also develop in the retina, but it is less prominent than in DR because of reduced retinal perfusion[92]. OIS entails an overall poor visual prognosis, which means that the ophthalmologist’s diagnosis is crucial for the systemic health of those patients because OIS may be the presenting sign of impending serious cerebrovascular and ischemic heart disease. Finally, DM can sometimes cause an ischemic optic neuropathy, which is a direct infarct of the optic nerve[52].

Another condition involving a main function of the eye where macrovascular disease in DM manifests is ophthalmoplegia. Ophthalmoplegia is the paralysis of one or more of the extraocular muscles (EOM). It can arise from traumatic, autoimmune, infectious, and vascular etiologies. Usually involving the third (oculomotor), fourth (trochlear), or sixth (abducens) cranial nerves, double vision is the characteristic symptom of ophthalmoplegia[93]. The vascular supply for the EOMs comes from branches of the ophthalmic artery, which is itself a branch of the internal carotid artery. Additionally, the cranial nerves responsible for the EOMs themselves have a complex vascular supply. Focal cranial nerve ischemia due to atherosclerosis within the microvasculature is thought to contribute to the development of ophthalmoplegia in patients with DM[94].

**Prevention and Management**

Preventing diabetic foot and eye problems is best achieved through regular examinations, diabetes education, and optimal management of underlying DM and its associated metabolic consequences. Tight control of blood glucose, as reflected by HbA1c level, is the most important element for prevention of these two interrelated diabetes-associated complications, closely followed by optimization of blood pressure and lipid levels[1,7,37,38]. This is accomplished through a combination of regulation of diet, lifestyle modification, body mass reduction, and medications, such as insulin and/or oral antidiabetic therapies, as appropriate. Monitoring alterations in serum biomarkers, such as HbA1c, ceruloplasmin, creatinine, uric acid, LDL, and HDL, is also important because these biomarkers are associated with both the onset and severity of DR and DFUs[8,20].

The standard practices in DFU management include cleansing, surgical debridement, application of clean dressings to maintain a moist environment and control exudates, wound off-loading, vascular optimization (including revascularization procedures), treatment and prevention of infection, and glycemic control[88,95]. Proper instruction is also required to prevent accidental or iatrogenic injuries which can result from ordinary hygiene and grooming of the feet and lower extremity[29]. Infection prevention is best achieved through protective footwear, proper hygiene, and offloading interventions[11]. Similarly, preventing complications from diabetic keratopathy focuses on limiting repetitive trauma, neurosensory deformities, exposure, and infections. Injuries may be caused by eye droppers, abnormal eye lashes, cosmetic applicators, fingers, facial towels, and bedding[96]. Infection can occur from overgrowth of the ocular flora or opportunistic infection enhanced by hyperglycemia, or it can take the form of chronic and recurrent herpes simplex and zoster[97].

Many parallels exist between management of ulcers in the cornea and those in the lower extremity. Both are treated with clean dressings, antimicrobial ointments, and salves. Wound infections may be polymicrobial, but the bacterial species most associated with DFUs include gram-positive species, *e.g.*, Staphylococcus aureus and Streptococcus species, but gram-negative infections with Pseudomonas aeruginosa and Enterobacteriaceae species also occur and are notably more common in ischemic or deep wounds[98]. Infection of corneal ulcers involve many of these same organisms, including Staphylococcus, Pseudomonas aeruginosa, and Streptococcus pneumoniae[70,99]. Special dressing and vacuum-assisted wound closure have been used with good result in the management of DFUs[100]. Non-healing diabetic corneal ulcers are often treated in conjunction with bandage contact lenses, which can lengthen the time antibiotic treatments are in contact with the ocular surface and serve as a reservoir for pharmacologically active compounds to aid wound healing[69]. In contrast to DFU management, patching should generally be avoided in patients with DM and corneal disease because of an increased risk of infection[101]. Amniotic membrane grafts have been studied for their potential of facilitating epithelial migration and healing of corneal ulcers and in very severe cases, corneal transplantation may be necessary[102].

Emerging research has placed an emphasis on developing therapeutic options that offer additional ways of preventing diabetic complications, treating them at earlier stages, or in more effective ways. As discussed earlier, inflammation has been implicated in the pathogenesis of diabetes-associated complications. Cytokines, such as interferon-γ, are being investigated as potential therapeutic targets in attenuating inflammatory cascades given that many of these cytokines contribute to altered vascular permeability and angiogenesis[103]. Given the high metabolic rate of the retina in conjunction with the metabolic stress induced by chronic hyperglycemia, reducing free radical stress may be an effective strategy[104]. Polyphenols, such as epigallocatechin-3-gallate found in green tea, are known for their antioxidant and anti-inflammatory properties and in diabetic animal models, have been shown to attenuate ROS concentrations in the retina[105]. Other polyphenol compounds, carotenoids, thiols, and vitamin supplementation are being investigated to address the several pathways involved in ROS generation and inflammation[104].

Finally, a multidisciplinary care team is essential to care optimally for the diabetic foot and eye, preserving function and quality of life for those with DM (Figure 3). Primary care providers and endocrinologists play a crucial role in coordinating care, including providing a formal assessment of the degree of diabetic control, screening for symptoms related to diabetic complications affecting other organ systems such as diabetic nephropathy, prescribing DM treatment, and involving specialists who manage diabetic complications such as foot or eye problems[1]. A diabetes care team should also include pharmacists who provide medication therapy management, dieticians, psychologists, diabetes care managers, and nurse educators. By working together, a coordinated care team can effectively reduce the healthcare burden associated with DM and its complications through prevention, screening, and management. In the future, the integration of smartphone technology and telehealth may not only streamline care coordination, but also allow for remote diagnosis and long-term monitoring of disease[106].

**CONCLUSION**

The identification of any ophthalmic or lower extremity complication in a patient with type 1 or type 2 DM should immediately prompt a review of DM management and coordination of diabetes care, including referral for reciprocal comprehensive foot or eye evaluations in patients with either complication[1,37,38].Although diabetic foot disease is slightly more common among patients with type 1 DM and those who use insulin[6],optimizing diabetes management remains the most important step in preventing diabetes-associated complications no matter what the type of DM[37,38]. While many patients may report symptoms related to diabetic foot disease or observe vision loss in the setting of diabetic eye problems, many others may be asymptomatic or have such mild signs and symptoms that they are easily overlooked, dismissed, or fail to receive clinical attention unless specifically assessed[64,107]. Primary care providers and endocrinologists should perform regular diabetic foot examinations because they provide insight into the presence and degree of PVD, neuropathy, skin breakdown, and other pre-ulcerative changes. Providers must also screen for signs and symptoms of eye disease, in part because their identification may help triage the urgency of any necessary referrals[37,38]. Because diabetic eye and foot diseases so commonly occur in conjunction, it is essential that clinicians take the necessary steps to reduce the impact of these diseases through regular screening, prompt referral to specialists, and providing a coordinated, team-based approach to management.

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Grade A (Excellent): 0

Grade B (Very good): B

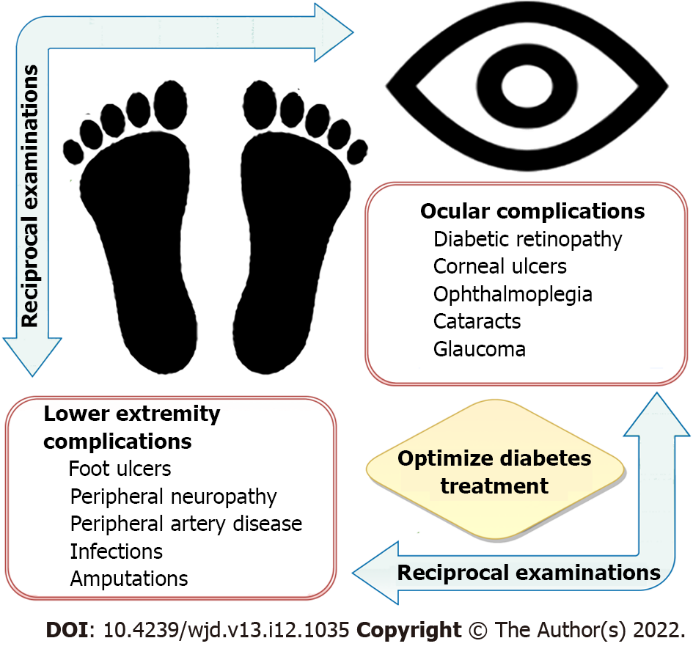
Grade C (Good): C, C

Grade D (Fair): 0

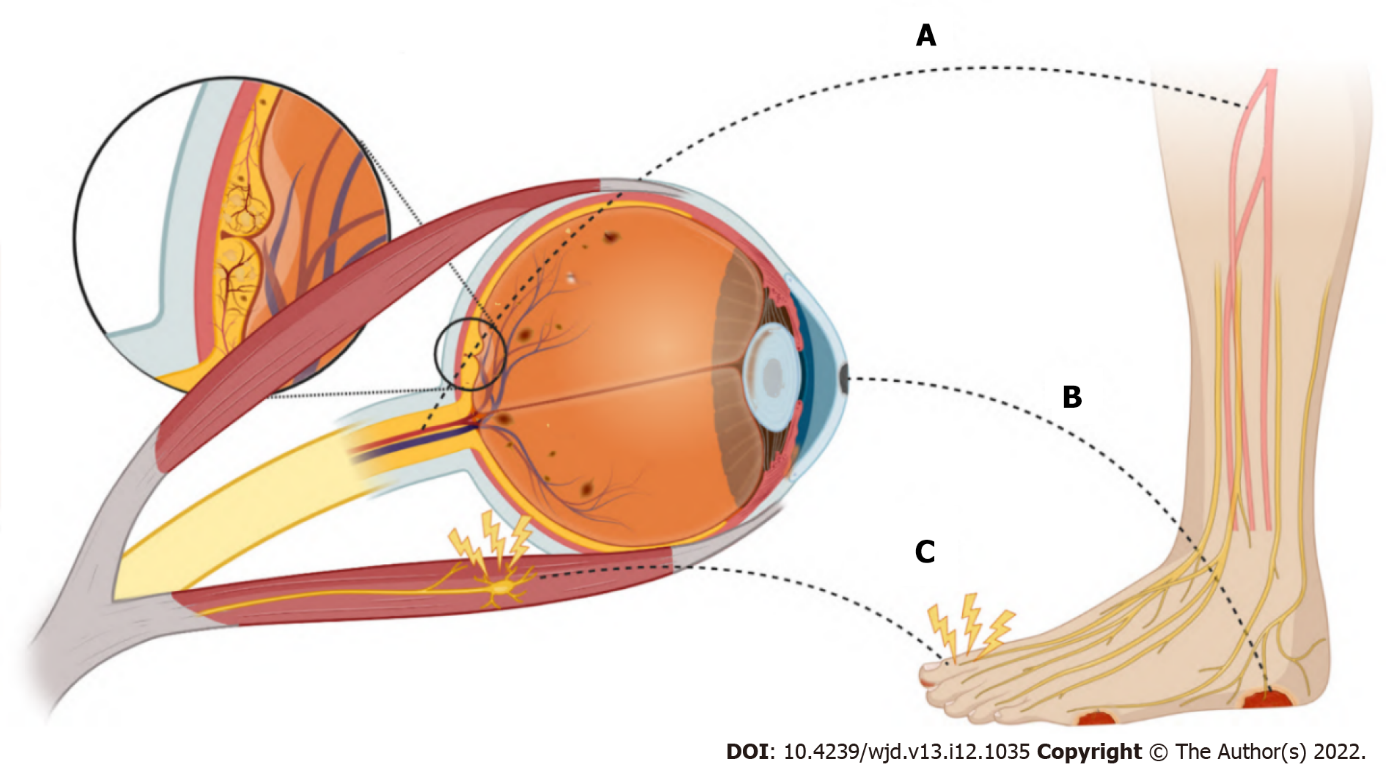
Grade E (Poor): 0

**P-Reviewer:** Morya AK, India; Wu QN, China **S-Editor:** Fan JR **L-Editor:** A **P-Editor:** Fan JR

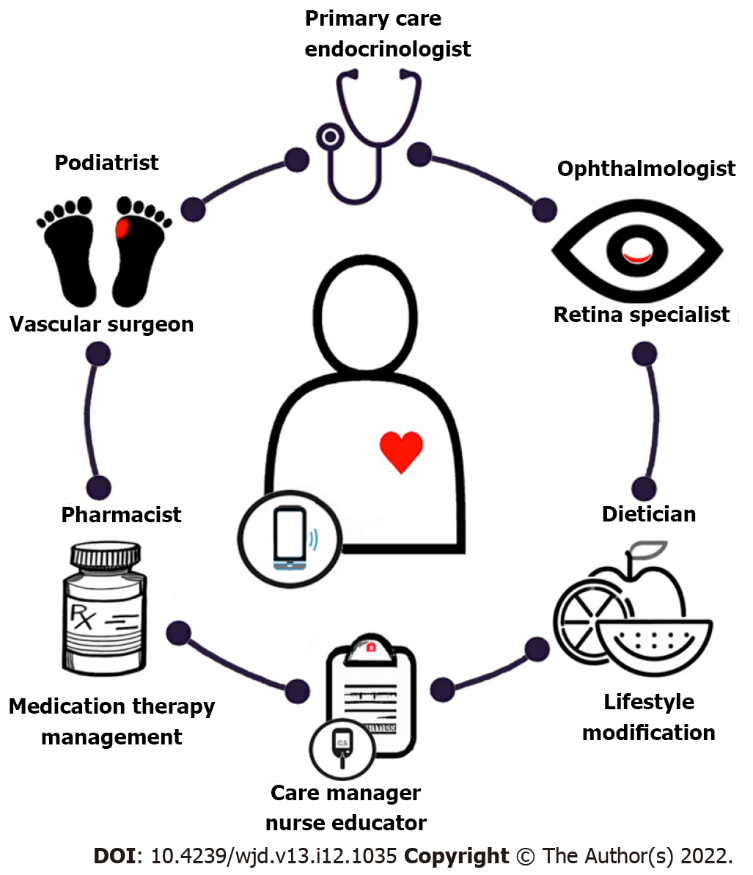
**Figure Legends**



**Figure 1 Upon identification of one or more of problems involving the lower extremity or the eye, reciprocal examinations are recommended to reduce the risk of further complications.** Preventing diabetic foot and eye problems is best achieved through regular examinations, diabetes education, and optimal management of underlying diabetes mellitus and its associated metabolic consequences.



**Figure 2 The connection between diabetic eye disease and diabetic foot and wound healing.** A: Diabetic micro- and macrovascular complications; B: Diabetic ulcers; C: Diabetic neuropathy. Each of these diabetic complications has multifactorial etiologies. Figure 2 was made in ©BioRender-biorender.com.



**Figure 3 Diabetic care team process.** Primary care and endocrinology physicians are central to comprehensive diabetes mellitus (DM) evaluation including assessment of the level of glycemic control, prescription of medications, determining level of treatment adherence, and identification of gaps in care and risk of complications. The frequent concurrence of diabetic eye and foot problems mandate that patients affected by either condition should undergo reciprocal comprehensive eye and foot evaluations, in addition to optimizing diabetic control. Specialists are often required to manage diabetic foot problems, including referral to podiatry, lower extremity wound care specialists, or vascular surgery, each engaging treatment algorithms according to their expertise. Eye care is typically provided by ophthalmologists or optometrists, but often requires the expertise of a retinal specialist capable of providing the medical and surgical management of diabetic eye disease. Pharmacists provide medication therapy management and are an important sources of diabetes education. Dietitians, lifestyle coaches, and psychologists offer counseling that works toward improving or maintaining glycemic targets through nutrition, achieving weight management and physical activity goals, and implementing behavior changes. Diabetic care managers and nurse educators help individuals with DM establish long-term commitments. They provide instruction on foot and skin care; the use of medications, including the administration of insulin; the monitoring of blood glucose levels; and maintenance of proper diet and exercise. They develop an overall management strategy aimed at reducing risk factors linked to diabetes-associated complications. The integration of smartphone technology and telehealth may streamline the care coordination and communication between the patient and each component of the diabetic care team[106]. Figure 3 was made in ©BioRender-biorender.com. The authors generated parts of the digital images used in Figure 3 by using the Generative Pre-trained Transformer 3 (GPT-3) autoregressive language model that employs deep learning to generate digital images from natural language descriptions (DALL·E, OpenAI, San Francisco, CA, labs.openai.com). The authors reviewed, edited, and revised these images and take ultimate responsibility for the content included in this publication.

**Table 1 Studies examining association of diabetic foot disease and diabetic retinopathy**

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Ref.** | **Year** | **Type of study** | **Sample size** **(DFU; no DFU)** | **Source of population** | **Main findings** |
| Jayaprakash *et al*[17] | 2009 | Prospective Case Study | 94 | India | 73.4% prevalence of DR in patients with DFUs |
| Hwang *et al*[22] | 2017 | Retrospective Cohort | 100 | South Korea | 90% prevalence of DR in patients with type 2 DM and DFUs; 55% had PDR |
| Karam *et al*[18] | 2018 | Cross-sectional | 182 | India | 67.6% prevalence of DR in patients diabetic foot disease (including neuropathy, deformation, DFUs, or amputation) |
| Zafar *et al*[19] | 2019 | Cross-sectional | 530 (225; 305) | Pakistan | 96% of patients with DFUs had DR |
| Sellman *et al*[23] | 2020 | Case Control | 270 (90; 180) | Sweden | 31% prevalence of PDR in patients with DFUs |
| Banik *et al*[21] | 2020 | Cross-sectional | 680 (8; 672) | Bangladesh | 65.9% prevalence of DR in patients with DFUs |
| Ye *et al*[20] | 2014 | Retrospective Cohort | 829 (61; 768) | China | OR 2.026 for DFUs in patients with DR |
| Al-Rubeaan *et al*[6] | 2015 | Retrospective Cohort | 62681 (2071; 60610) | Saudi Arabia | OR 4.45 for diabetic foot disease (including DFUs, gangrene, and amputation) in patients with DR |
| Harris Nwanyanwu *et al*[24] | 2013 | Retrospective Cohort | 4617 | United States | 1.54 HR for those with comorbid non-healing DFUs to progress from NPDR to PDR in three to five years |

DR: Diabetic retinopathy; NPDR: Nonproliferative diabetic retinopathy; PDR: Proliferative diabetic retinopathy; DFU: Diabetic foot ulcer; OR: Odds ratio; HR: Hazard ratios.



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