Directed Therapy for Exfoliation Syndrome

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Abstract: Exfoliation syndrome (XFS) is an age-related disorder of the extracellular matrix that leads the production of abnormal fibrillar material that leads to elevated intraocular pressure and a relatively severe glaucoma. Exfoliation material is deposited in numerous ocular tissues and extraocular organs. XFS is associated with ocular ischemia, cerebrovascular disease, neurodegenerative disease and cardiovascular disease. Current modalities of treatment include intraocular pressure lowering with topical antihypertensives, laser trabeculoplasty and filtration surgery. The disease paradigm for XFS should be expanded to include directed therapy designed specifically to target the underlying disease process. Potential targets include preventing the formation or promoting the depolymerization of exfoliation material. Novel therapies targeting trabecular meshwork may prove particularly useful in the care of exfoliative glaucoma. The systemic and ocular associations of XFS underscore the need for a comprehensive search for neuroprotective agents in its treatment.

1. PATHOGENESIS AND PRESENTATION OF EXFOLIATION SYNDROME

Epidemiology and Clinical Presentation

Exfoliation syndrome (XFS) is an age-related disease of the extracellular matrix characterized by the production and deposition of a fibrillar extracellular material in numerous ocular tissues. It is now recognized as the most common identifiable cause of open-angle glaucoma, affecting an estimated 60 to 70 million people worldwide [1, 2]. Its reported prevalence rates vary extensively due to true population variations as well as the clinical criteria used and the ability of the examiner to detect the clinical signs of the disease.

The diagnosis of XFS is confirmed by the presence of white exfoliation material (XFM) at the pupillary border or on the anterior lens capsule in a characteristic distribution. A central disc is separated from a granular peripheral zone by an intermediate clear zone demarcated by XFM by friction between the iris and anterior lens capsule during pupillary movement. The material on the anterior lens capsule causes rupture of iris pigment epithelial cells leading to pigment dispersion, trabecular meshwork hyperpigmentation, iris transillumination defects at the pupillary margin and loss of the pupillary ruff. Pigment dispersion after dilation can lead to marked IOP rise and may be one of the earliest manifestations of the disease.

Patients with XFS are twice as likely to convert from ocular hypertension to detectable glaucoma [3]. Exfoliative glaucoma (XFG) is more severe than primary open angle glaucoma (POAG). There is greater diurnal intraocular pressure (IOP) fluctuation, greater visual field loss and optic nerve head damage at presentation, poorer response to medications, faster visual field progression and increased need for surgical intervention. At any given IOP, eyes with XFS are at significantly greater risk for glaucomatous damage, suggesting that pressure-independent factors play a significant role in visual field loss in these patients.

The prevalence of XFS increases with age in all affected populations. Approximately two-thirds of patients have unilateral disease on clinical examination; however, XFS is often detectable in the contralateral eye with conjunctival biopsy [4]. This finding supports the hypothesis that the development of XFS is similar to uveitis in that an underlying predisposition plus an environmental or immunologic trigger leads to disease manifestation.

Pathogenesis of XFS

The pathogenesis of XFS is defined by the accumulation of an abnormal fibrillar material biochemically related to the composition of the extracellular matrix that results either from excessive production and/or insufficient breakdown. The product has a unique structure as seen by electron microscopy [5]. Streeter et al. first hypothesized that XFS is an elastosis based on similarities of XFM and zonular fibers [6]. These characteristic fibrils, composed of microfibrillar subunits surrounded by an amorphous matrix, contain epitopes of elastic fibers such as fibrillin-1, elastin, tropoelastin, amyloid P and microfibril-associated glycoprotein, vitronectin and the cross-linking enzyme lysyl-oxidase [7, 8].

Two common single nucleotide polymorphisms (SNPs) in the coding region of the lysyl-oxidase-like 1 (LOXL1) gene on chromosome 15 have been identified and account for virtually all cases of XFS in several populations [9-12]. While the SNPs confer susceptibility to XFS, not all people with the SNPs will develop the disease.
LOXL1 is a member of the lysyl oxidase enzyme family responsible for formation, stabilization, and remodeling of elastic fibers. They are important for cross-linking of elastin, for providing a scaffold that ensures spatially defined deposition of elastin and for regulating the production of elastin [13, 14]. The currently proposed pathogenesis of XFS is a stress-related, excessive production by elastogenic cells of elastic microfibrils that aggregate into a typical configuration via an enzymatic cross-linking process [15]. There is evidence that derangement of the balance of matrix metalloproteinases and their tissue inhibitors, increased oxidative stress and an impaired stress response contribute to impaired degradation and accumulation of fibers [16-19]. These fibers impair trabecular meshwork outflow resulting in an increased rise of IOP.

**Ocular and Systemic Associations**

XFS is associated with both chronic open-angle glaucoma and angle closure. The zonules and ciliary body are often diffusely coated with XFM and zonules are often frayed and broken. Zonular fragility and poor pupillary dilation contribute to an increased incidence of serious complications during cataract extraction, including zonular dehiscence and vitreous loss. Phacodonesis and even complications during cataract extraction, including zonular dehiscence and vitreous loss. Phacodonesis and even long-term worsening of trabecular function that may be particularly detrimental in XFS [42].

Unfortunately, pilocarpine is underutilized in the treatment of glucoma due to the possibility of decreased vision or the misconception that it must be used q.i.d. It is our experience that pilocarpine 2% q.h.s. can provide significant limitation of pupillary movement without the inconvenience of q.i.d. dosing and dimming of vision. Pilocarpine can also blunt an early morning IOP spike [43].

Argon laser trabeculoplasty is particularly effective in XFS [44]. Primary ALT can delay the need for topical therapy in some patients, although there is a general decrease in efficacy over time, with 35-55% remaining medication free at 3-5 years [45]. A minority of patients will experience a sudden and dramatic rise in IOP within two years following treatment, presumably due to continued pigment liberation overwhelming the meshwork. Pilocarpine 2% q.h.s. may provide protection from this. Limited data on selective laser trabeculoplasty (SLT) suggests that success may be comparable, although prospective studies are necessary to confirm this and also to compare results with argon laser trabeculoplasty [46]. Marked IOP elevations associated with pigment release in patients with heavily pigmented trabecular meshworks necessitating filtration surgery may occur after SLT [47]. Trabeculectomy with antifibrotic therapy leads to comparable IOP reduction in XFS and OAG, and offers greater 24-hour IOP control than medical therapy alone [48].

The investigation of surgical therapy directed at trabecular dysfunction is particularly sensible for XFS. Trabeculectomy has been reported as successful in XFS although additional trials are required [49-51]. Goniotomy is also successful (Kim JH, Sbeity Z, Ritch R: Long-term results of goniotomy and cataract surgery for exfoliative glaucoma. Southeast Asian Glaucoma Interest Group, Seoul, Republic of Korea, Sept. 25, 2008.) Jacobi and Krieglstein [52] described a procedure in which trabecular aspiration of elastic microfibrils is performed. Goniotomy is also successful (Kim JH, Sbeity Z, Ritch R: Long-term results of goniotomy and cataract surgery for exfoliative glaucoma. Southeast Asian Glaucoma Interest Group, Seoul, Republic of Korea, Sept. 25, 2008.)

**Directed Ocular Anti-hypertensive Therapy**

The management of XFS currently hinges on ocular antihypertensive therapy and initially can be achieved with topical therapy in many cases. Prostaglandin analogues such as latanoprost and travoprost reduce IOP at each time point in the 24-hour diurnal curve as compared to untreated IOP, with travoprost conferring a slightly higher level of reduction [40]. Latanoprost reduced IOP and narrowed the range of diurnal fluctuation in comparison to timolol in XFS [41].

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bind to free actin in trabecular cells leading, to temporary disintegration of the existing actin cytoskeleton thereby increasing aqueous outflow [53, 54]. Unlike previous compounds with similar mechanisms of action such as sodium-EDTA and cytochalasin B [55, 56], Latrunculin has no known adverse effects on retinal and corneal function [57-59]. Ideally, latrunculin would be effective as a depot medication with once a week or once a month dosing.

**Treatment Directed at XFM**

Future therapy of XFS should be directed at preventing formation of XFM or depolymerizing existing XFM. An intriguing line of research is an investigation of the factors that protect the contralateral eye from developing XFS. If the development of the disease in the second eye is actively suppressed, as by an immune mechanism, then elucidation of this could lead to a novel treatment. The systemic ramifications of XFS and the morbidity associated with these diseases highlight the need for a therapy that prevents aggregation of XFM microfibrils disaggregating the microfibrils.

Homocysteine is an amino acid, elevations of which result from a disturbance of methionine metabolism. Hyperhomocysteinemia is a widely recognized cardiovascular risk factor and is reversible with dietary supplementation with folic acid, pyridoxine (vitamin B6) and cyanocobalamine (vitamin B12) [60, 61]. Hyperhomocysteinemia is associated with disruption of the elastic fibers of the extracellular matrix leading to vascular disease [62]. Elevated homocysteine levels are present in blood, aqueous and tears of patients with XFS, and there are decreased serum levels of cyanocobalamin, pyridoxine and folic acid [63-66]. The concept of a relationship between hyperhomocysteinemia is supported by the propensity for cardiovascular and cerebrovascular disease in patients with XFS. By the same virtue, it is possible that treatment of hyperhomocysteinemia could be beneficial in XFS as it is in cardiovascular disease.

Folic acid deficiency leads to altered expression of genes involved in cell signaling, the cytoskeleton and the extracellular matrix [67]. Actin disrupting agents, such as latrunculin B, reversibly increase the proportion of receptors to extracellular matrix [67]. Latrunculin has no known adverse effects on retinal and corneal function [57-59]. Ideally, latrunculin would be effective as a depot medication with once a week or once a month dosing.

**Treatment Targeting Non-pressure Dependent Mechanisms**

Patients with XFS have an increased likelihood to have coexisting cerebrovascular and cardiovascular disease, and systemic agents that provide neuroprotection and vasoprotection may prove to be an important therapeutic tactic in the treatment of XFS. Pharmacologic treatment of non-IOP dependent mechanisms in glaucoma has traditionally been limited to calcium channel blockers that are widely used in treatment of hypertension, coronary artery disease, migraines and Raynaud’s disease. The effect of calcium channel blockers on visual field progression and ocular blood flow is disputed [74-76].

Gingko biloba extract (GBE) has been used in Chinese medicine since 3000 BC for the treatment of aging, dementia, tinnitus, bronchoconstriction and sexual dysfunction with minimal side effects. It improves peripheral, cerebral and ocular blood flow, prevents platelet aggregation and protects against free radical damage [77, 78]. Neurons in tissue culture are protected from toxicity-induced apoptosis with GBE, and there is mixed evidence that GBE improves function in patients with Alzheimer’s disease and vascular dementia [79, 80]. Further elucidation by prospective trials of GBE in glaucoma is warranted.

Curcumin is an anti-oxidant extracted from the plant *Curcuma longa* and is a component of the commonly used Indian spice turmeric. Curcumin has antioxidant, anti-inflammatory, anti-angiogenic and anti-neoplastic activity via the inhibition of numerous mediators of inflammation [81-85]. Clinical trials for the treatment of gastrointestinal disease and other neoplasms as well as Alzheimer’s disease are under way. Common therapeutic doses cause only mild gastrointestinal discomfort in a minority of patients.

Resveratrol is a compound found in the skin of red grapes. It is an effective antioxidant and protects against degeneration of neurons in ischemia. It is hypothesized that resveratrol is responsible for the “French paradox”, the observation that the French have a lower incidence of heart disease despite a high fat diet due to the consumption of red wine. It is an effective antioxidant and prevents neuron degeneration [84, 85].

**3. CONCLUSION**

XFS is a systemic disorder with components of low-grade inflammation and oxidative damage leading to accumulation of XFM. The ocular morbidity associated with XFS far exceeds that of open-angle glaucoma, and a host of systemic diseases with considerable morbidity and mortality are associated with XFS. Further elucidation of the genetics and pathophysiology of XFS is crucial to our ability to treat this disease effectively. There is a role for non-IOP lowering treatment modalities in XFS as there is significant evidence of generalized ischemia and neurodegeneration. With this
directed investigation, there is hope that XFS will be both a preventable and curable disease.

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