Retinal arteriolar occlusions due to cytomegalovirus retinitis in elderly patients without HIV

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Abstract

Background: Five of 7 (71%) elderly immunocompetent patients with cytomegalovirus retinitis had retinal arteriolar occlusions versus 2 of 8 (25%) elderly immunocompromised patients and 1 of 19 (5%) younger HIV-infected patients. Compared to HIV-infected patients, elderly patients were more likely to have occlusive events, neovascularization or hemorrhage, and underlying vasculopathy. The purpose of this study is to report the novel finding of extensive retinal arteriolar occlusions and neovascularization in immunocompetent patients with cytomegalovirus retinitis. This is a retrospective observational cohort study of cytomegalovirus retinitis (CMVR) in a university setting. Seven patients were elderly but not immunocompromised, 8 were elderly and iatrogenically immunocompromised, and 16 were HIV-infected. All patients underwent polymerase chain reaction testing of intraocular fluid. Primary outcome measure was visual acuity. Secondary outcome measures were vascular occlusions, ischemic complications, and response to treatment.

Results: Mean age was 73, 70, and 41 years for immunocompetent, immunocompromised, and HIV-infected patients, respectively. Diabetes and vascular disease were common in the elderly. Vision loss to less than 5/200 occurred in 50% of the immunocompetent elderly patients, and 17% of CMV eyes in immunocompromised and HIV patients. Occlusion of the entire retinal vasculature occurred in 4/7 (57%) of immunocompetent patients despite lack of Zone I involvement, and rubeosis occurred in three, disc neovascularization in one, and vitreous hemorrhage in two patients. Vascular occlusive events were less common in immunocompromised patients and rare in the HIV-infected.

Conclusions: CMVR in non-HIV-infected elderly patients is associated with retinal arteriolar occlusions. An intact host immune response may increase damage to retinal vessels. Prompt diagnosis may avert catastrophic vision loss.

Keywords: Cytomegalovirus retinitis, Immunocompetent, Immunocompromise, Aged, HIV, Retinal arteriolar occlusion
Results

Fifteen patients age 60 and older were identified who were not HIV-infected and had a diagnosis of CMVR after diagnostic PCR testing. Seven, with an average age of 73 ± 7.4 years, were judged to be immunocompetent. Of these, one was taking low doses of oral corticosteroids for healed giant cell arteritis, and four had controlled diabetes mellitus; one had chronic lymphocytic leukemia (CLL) that was controlled with no medication (Table 1). Eight patients, with an average of 72.2 ± 12.2 years, were judged to be immunocompromised. Of these, two were organ transplant recipients, and two had lymphoma being treated with chemotherapy; the remaining patients were receiving immunosuppression for a variety of autoimmune disorders (Table 2). Sixteen HIV-infected patients with CMVR were selected for comparison because the diagnosis had been confirmed with PCR, and they were contemporaneous (2000–2005) with the non-HIV-infected patients. Average age was 40.3 ± 11.6 years.

Males accounted for approximately 70% of the patients in each group (P = 0.56). None of the immunocompetent patients had bilateral disease (0%), but 4 of the 8 (50%) immunocompromised non-HIV-patients had bilateral disease, as well as 4 of the 16 (25%) HIV-infected patients (P = 0.6). Profound, permanent vision loss worse than 20/400 occurred in 8 of the 19 eyes in the elderly, and 10 of the 20 eyes in the HIV-infected (P = 0.14). In the eight cases of profound vision loss in the elderly, five were due to arteriolar occlusion, two to CMVR; one other case had both prior vein occlusion and extensive retinitis. Among HIV-infected patients, vision loss was attributable to arteriolar occlusion in only one case (P < 0.001). The remaining vision loss was due to retinal detachment or to CMVR.

More arteriolar occlusions occurred in the eyes of the elderly, immunocompetent patients than in those of elderly, immunocompromised patients (5 of 7 (71%) vs. 1 of 12 eyes (8%)), P = 0.07. When arteriolar occlusions in non-HIV-infected patients were compared to HIV-infected, the results were highly statistically significant (6 of 19 eyes (32%) vs. 1 of 20 eyes (5%)), P < .001. Findings associated with ocular ischemia, such as neovascularization, rubeosis, and vitreous hemorrhage, were also more common in non-HIV-infected patients (6 of 19 (32%) vs. 4 of 20 eyes (20%)), P < 0.003.

Vasculopathic diagnoses such as cardiovascular disease and diabetes were more common in the elderly, especially those who were not considered to be otherwise immunocompromised (Tables 1 and 2). Compared to HIV-infected patients, vasculopathy was present in 10 of the 15 patients (67%) vs. 1 of 20 (5%), P < 0.001. Retinal detachment was more common in eyes in HIV-infected patients than in non-HIV-infected patients (7 vs. 2, P = 0.03). Table 3 lists the medical and surgical interventions, and visual and anatomic outcomes.

Case reports

Case 1

A 71-year-old Indian man with hypertension and diabetes had blurred vision in the right eye for 3 months. Vision was 20/40+ in the right eye; iritis and vitreous...
inflammation were present. Peripapillary and macular cotton-wool spots were noted and there was peripheral nonperfusion with scattered intraretinal hemorrhages temporally in the right eye. Diabetic retinopathy was not present. The left eye was normal.

Fluorescein angiography at that time showed intact choroidal filling and delayed filling of the arteriolar vasculature in the right eye with incomplete filling in the midperiphery. Subtenon triamcinolone injection was given elsewhere to treat the inflammation. Toxoplasmosis. Intravenous ganciclovir was prescribed for HSV 1 and 2, varicella zoster virus (VZV), and Epstein-Barr virus. 

Lab testing including complete blood count (CBC), anti-neutrophil cytoplasmic antibody (ANCA), rapid plasma reagin (RPR), fluorescent treponemal antibody, human T-cell lymphotropic virus-1 antibody, leptospira antibody, and human immunodeficiency virus (HIV) antibody were negative or normal. PCR analysis of the aqueous humor was positive for CMV DNA and negative for HSV 1 and 2, varicella zoster virus (VZV), and toxoplasmosis. Intravenous ganciclovir was prescribed but stopped after 3 weeks by an infectious disease specialist, who found no other signs of CMV disease.

Three months later, the retinitis remained active. Vision was bare light perception, and the retinal circulation appeared more occluded. Diagnostic pars plana vitrectomy with retinal biopsy, membrane peel, and endolaser, and intravitreal triamcinolone acetonide and C3F8 intraocular gas tamponade was performed. PCR of vitreous fluid was again positive for CMV DNA and negative for HSV, VZV, toxoplasmosis, and Epstein-Barr virus. Retinal biopsy showed cytomegalic inclusion bodies and CMV by immunohistochemistry (Figure 3). At surgery, the nummular lesions appeared as multiple, intraretinal, white snowflake opacities with adhesion to the underlying RPE. No bleeding occurred during retinal incision. All retinal arteries were sclerotic, with a crystalline appearance, and the optic nerve head was completely atrophic. On final follow-up, the patient had a visual acuity of light perception in his right eye with neovascularization of the iris, ectropion uveae, and dense vitreous hemorrhage despite endolaser at surgery.
Figure 1 Case 1, posterior fundus, right eye. Left frame: there are sclerotic central vessels and cotton-wool spots located inferotemporal to the optic disc and fovea. Hemorrhages are located along the superior and inferior vascular arcades. Right frame: late-phase fluorescein angiography shows minimal retrograde filling from the optic nerve circulation of the retinal vasculature with late intraretinal staining around the optic nerve. There is extensive capillary nonperfusion peripherally.

Table 3 Medical and surgical interventions in elderly, immunocompetent, and elderly immunocompromised non-HIV-infected patients with cytomegalovirus retinitis

| Case | Medical treatment                                                                 | Surgical interventions                                      | Retinovascular disease                                                                 |
|------|------------------------------------------------------------------------------------|-------------------------------------------------------------|---------------------------------------------------------------------------------------|
| 1    | Intravenous ganciclovir                                                            | Diagnostic paracentesis, diagnostic vitrectomy, retinal biopsy | Disc neovascularization, rubeosis iridis, ectropion uvea, vitreous hemorrhage          |
| 2    | Oral valganciclovir, intravitreal foscarnet, intracameral bevacizumab               | Diagnostic paracentesis, PRP                                 | Rubeosis iridis                                                                      |
| 3    | Clarithromycin, atovacuoquine, valacyclovir, intravenous and oral ganciclovir       | Diagnostic vitrectomy, trabeculectomy, diagnostic paracentesis, glaucoma drainage device | Rubeosis iridis, neovascular glaucoma                                                  |
| 4    | Prednisone, subtenon triamcinolone, valacyclovir, ganciclovir drug delivery device | Diagnostic vitrectomy, CE/IOL                                 | Vitreous hemorrhage                                                                  |
| 5    | Intravitreal ganciclovir and foscarnet, valganclovir                               | Diagnostic paracentesis                                     | None                                                                                  |
| 6    | Intravitreal ganciclovir and foscarnet, intravenous ganciclovir                    | Diagnostic paracentesis                                     | Sclerotic arterioles in Zone I                                                        |
| 7    | Intravitreal ganciclovir and foscarnet, valganclovir                               | Diagnostic paracentesis, RD repair                           | None                                                                                  |
| 8    | Lost to follow-up                                                                  | Prior diagnostic vitrectomy and PRP, diagnostic paracentesis | Vitreous hemorrhage                                                                  |
| 9    | Intravitreal ganciclovir and foscarnet, valganclovir                               | Diagnostic paracentesis                                     | None                                                                                  |
| 10   | Valganciclovir                                                                     | Diagnostic paracentesis                                     | None                                                                                  |
| 11   | Valganciclovir, intravitreal ganciclovir and foscarnet                            | Diagnostic paracentesis                                     | None                                                                                  |
| 12   | Valganciclovir, intravitreal ganciclovir and foscarnet                            | Diagnostic paracentesis                                     | Vitreous hemorrhage                                                                  |
| 13   | Intravenous ganciclovir                                                            | Diagnostic paracentesis                                     | None                                                                                  |
| 14   | Intravenous ganciclovir, intravitreal ganciclovir and foscarnet                    | Diagnostic paracentesis, RD repair                           | None                                                                                  |
| 15   | Intravenous ganciclovir, intravitreal ganciclovir and foscarnet                    | Diagnostic paracentesis                                     | None                                                                                  |

CE/IOL, cataract extraction with intraocular lens implantation; RD, retinal detachment; PRP, panretinal photocoagulation.
Case 2
An 81-year-old white man experienced decreased vision and redness in his right eye that worsened over 3 weeks. Past medical history was significant for multiple sclerosis in 1950 with bilateral optic neuritis in remission, cerebrovascular accident in 2000, hyperlipidemia, and diabetes. Granulomatous anterior segment inflammation and a central retinal artery occlusion of the right eye were noted by an outside ophthalmologist. Serum CMV IgG antibody and varicella zoster antibody were positive. CMV IgM antibody, HSV I and II IgG antibodies, and syphilis antibody were negative. Carotid artery stenosis was 70% to 80% on the right and 50% on the left on carotid Doppler ultrasound. Varicella-related necrotizing retinitis was suspected, and famciclovir 500 mg three times daily was initiated.

On referral 2 months after onset, vision was counting fingers at 6 feet in the right eye and 20/20 in the left eye with a 2+ relative afferent pupillary defect in the right eye. There were large white keratic precipitates on the right corneal endothelium, 2+ anterior chamber cell, and 1+ flare. On dilated examination of the right eye, the optic nerve was pale, with sclerotic, attenuated arterioles, segmentation of the venous blood columns, and no evidence of recanalization (Figure 4). There was a patchy, granular retinitis in the inferotemporal quadrant. The diagnosis of necrotizing retinitis with central retinal artery occlusion was made. The left eye was free of inflammation.

Diagnostic AC paracentesis for PCR for CMV, varicella zoster virus, and herpes simplex virus was positive for CMV DNA only. HIV serology was negative. An intravitreal injection of foscarnet was given and oral valganciclovir was started. Three months later, there was florid neovascularization of the iris. The retinitis was inactive. Bevacizumab was injected intracameral, and PRP was performed with regression of the NVI. Vision remained stable at hand motions. Valganciclovir was continued at 450 mg daily adjusted for renal function to complete 3 months of treatment.

Case 3
A 66-year-old white man presented with 3 weeks of blurred vision and floaters in his left eye. Past medical history was significant for chronic open angle glaucoma and CLL status post chemotherapy, with last treatment 1.5 years prior. The patient denied pain, redness, or photophobia. Vision was 20/20- in the right eye and 20/30 in the left eye. Slit lamp exam showed 2+ flare with rare cells and fine keratic precipitates in the left eye. Fundus examination on the left revealed 2+ vitreous cells. There was a small patch of chorioretinitis present in the superotemporal periphery. A presumptive diagnosis of toxoplasmosis chorioretinitis was made. Clarithromycin and atovaquone were started; laboratory studies included complete blood count with differential, complete metabolic profile,
rheumatoid factor, ANCA, RPR, lyme antibodies, HLA-B27, and angiotensin converting enzyme. CBC was remarkable with 63,300 WBC with 92.4% lymphocytes. All other laboratory studies were within normal limits; HLA-B27 antigen was present. Subsequent toxoplasmosis IgG was positive, with a negative IgM.

After 1 week, the retinitis appeared to resolve, but one month later, vision dropped to 20/200. Intraocular pressure was 32 mmHg with 1+ anterior chamber cell and flare and 2+ vitreous cells. Fundus examination showed extensive arteriolar closure with segmentation of the venous circulation confirmed by fluorescein angiography (Figure 5). The focus of chorioretinitis superotemporally had enlarged slightly (Figure 6, left). Diagnostic vitrectomy was performed 2 days later; vision prior to surgery had declined to 6/200 ‘E’. The vitrectomy was combined with trabeculectomy with mitomycin C. Vitreous fluid was sent for PCR analysis for CMV, HSV 1 and 2, and VZV DNA, as well as toxoplasma IgG, flow cytometry and cytology, and viral culture. All of these tests were negative.

Empiric valacyclovir was started. The retinal vessels appeared attenuated with macular ischemic whitening consistent with central retinal artery occlusion. Two weeks later, vision declined to 3/200 ‘E’. Anterior chamber paracentesis was performed, and CMV DNA was detected by PCR. Intravenous ganciclovir was started 300 mg every 12 h for 14 days followed by oral ganciclovir 1 gram three times daily. Within 2 months, the retinitis appeared healed; however, neovascular glaucoma developed requiring Baerveldt glaucoma drainage implant. The intraocular inflammation and retinitis reactivated, with focal nummular white lesions encroaching on the superior arcade. No further therapeutic intervention was recommended due to poor visual prognosis, and the retinitis eventually progressed to involve most of the periphery (Figure 6, right).

Figure 4 Case 2, posterior fundus, right eye. Left frame: 5 weeks after vision loss from central retinal artery occlusion, there is no recanalization of the artery. The venous blood columns are segmented with a few intraretinal hemorrhages. Retinitis is barely visible inferotemporally to the macula. Right frame: the inferotemporal periphery shows nummular lesions of a granular retinitis similar to Case 1. The vascular closure is generalized rather than specific to the area of retinitis.

Figure 5 Case 3, posterior fundus, left eye. Left frame: the vasculature is perfused centrally but there are sheathed arterioles and intraretinal hemorrhages. Right frame: arteries and veins are truncated with extensive capillary nonperfusion in the macula.
Discussion
The pathogenesis of CMVR likely involves hematogenous dissemination with initial infection of the vascular endothelium and subsequent spread into adjacent tissues [5]. In HIV infection, breakdown of the blood-retinal barrier by HIV retinopathy [6] may facilitate viral spread into the retinal tissue. Pro-inflammatory cytokines may contribute to the vasculopathic effects of the virus [7] with expression of endothelial adhesion molecules, leading to increased platelet and leukocyte adhesion, induction of anti-phospholipid antibodies, and enhanced factor VIII synthesis and secretion [7].

CMVR and HIV-related branch retinal artery occlusions have been rarely reported [8,9]. There have also been a few case reports of retinal vascular occlusion in HIV/AIDS patients without CMVR [10-15]. In the five cases of ARN-like CMVR in immunocompetent patients reported in the literature [2,3,16,17], one had branch retinal artery occlusion [14]. Designation of patients as immunocompetent or immunocompromised is imprecise even when special testing of the immune system is performed. The elderly patients we deemed immunocompetent were not receiving immunosuppression for organ transplantation or chemotherapy for cancer, which are the usual causes for CMVR in non-HIV patients. They were elderly with associated co-morbidities and, therefore, in a gray zone of immunocompetency. They represent a large population at risk for CMVR with few apparent risk factors other than advanced age.

Well-established risk factors for vascular occlusion were present in our patients including hypertension, increased white blood cell counts, uncontrolled glaucoma, giant cell arteritis, cardiovascular disease, and carotid stenosis [18]. CMV infection has been associated with atherosclerosis and with impaired circulation [19,20]. Vascular occlusion contributed to poor visual acuity outcomes independent of macular involvement or retinal detachment, which are the usual means by which patients with CMVR lose vision but were not present in these patients. A weaker than normal immune system in these elderly patients that allowed for CMVR combined with a sufficient immune response to damage and close vessels moderately impaired by other vasculopathies may have produced this unique presentation. In cases 1 and 4, the early use of corticosteroids without antiviral cover before the diagnosis of CMVR was made may have allowed sufficient worsening to permit this unusual constellation of findings.

Extensive vascular occlusion occurred when the retinal surface area involved with retinitis was still quite small. Although risk factors for vascular closure were present, the artery occlusions were not typical for embolic disease, for example, typical cherry-red spots and recanalization did not occur. Despite extensive occlusions as in cases 1 and 3, at least some portion of the central retinal circulation remained patent until late in the course. The process therefore was more likely progressive nonperfusion that finally affected the central vasculature rather than an acute event such as a central retinal artery occlusion. The incremental closure of the total retinal circulation may have led to other atypical features such as the high frequency of secondary neovascularization, which is unusual in central retinal artery occlusion and also unusual in eyes of patients with CMVR involving the entire retina.

Conclusion
Retinal artery occlusion can occur in conjunction with CMVR in elderly, immunocompetent patients. Recognition of CMV as a rare cause of necrotizing retinitis and institution of early antiviral treatment may reduce the
risk of this devastating complication. Since generalized vascular occlusion is an uncommon manifestation of CMVR, older age, concurrent illnesses, and relatively intact immune defenses with intravascular inflammation may have contributed to this devastating outcome.

Methods

Patient lists were searched to identify non-HIV-infected patients with CMVR from 2000–2007. Medical records were retrospectively reviewed for age and sex; medical status including iatrogenic immunosuppression; ocular characteristics including extent of retinitis, vascular occlusions, neovascularization, and hemorrhage; clinical estimate of grade of inflammation; clinical course including timing of vascular occlusion relative to onset of symptoms; results of diagnostic studies; medical treatment and response to medication; surgical interventions; and visual outcomes. Narrative accounts of three cases considered to be immunocompetent who developed retinal arteriolar occlusion were created. Control patients were selected from a group of HIV-infected patients with CMVR confirmed by polymerase chain reaction (PCR) testing and previously reported from this institution [21].

Paired McNemar statistics were used to compare the presence and absence of bilateral disease, profound vision loss, arteriolar occlusion, neovascularization or vitreous hemorrhage, and retinal detachment between the non-HIV-infected and HIV-infected patients. Sample size hampered comparisons between the elderly, immunocompromised and elderly, immunocompetent groups, and the elderly, immunocompetent and HIV-infected groups. Expedited approval for retrospective review of medical records was granted by the Human Subjects Review Board by the University of Miami Miller School of Medicine with waiver of HIPAA authorization.

Competing interest

The authors declare that they have no competing interests.

Authors’ contributions

JLD conceived of the study, provided patients, performed the statistical analysis, and wrote the final manuscript. PH and KH provided patients and drafted the manuscript. All authors read and approved the final manuscript.

Authors’ information

The authors presented this study at the Joint Meeting Retina Society, Macula Society, American Society of Retina Specialists, New York, NY on October 2, 2009. KH and PH participated in the study while in residency training at Bascom Palmer Eye Institute and are now practicing ophthalmologists.

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