Introducation

Chorioretinal folds (CFs) are most often fluctuating or streaking, which is also found in the posterior pole, which can also include internal choroid, Bruch’s membrane, retinal pigment epithelium (RPE) and occasionally neurosensory retinas. A series of thin lines or streaks, usually arranged in light-dark color that extends parallel to each other but can be more or less vertical or oblique and very rarely extends beyond the equator. CFs were first described by Edward Nettleship in a patient with atrophic papilledema due to an intracranial mass in 1884 [1]. The yellow, hypopigmented overhangs are the areas where the RPE is stretched and tapered, while the darker and deeper bands are the areas where the RPE is under compression. This appearance of RPE is also the cause of hyper and hypolorescence in funds fluoresce in angiography (FFA). In 1972, Newell stated that the CFs was formed due to the close connections between the Bruch membrane and the underlying chori-capillaries [2].

When the choroid swells and expands, the upper Bruch membrane becomes forced to fold and the clinical appearance of the CFs is formed. Idiopathic chorioretinal folds associated with benign conditions such as hypermetropia are the most common, but the orbital mass, inflammation, sclerotic, sciera buckling, choroid mass, chorioidal neovascular membrane (CNV), chronic papilloma, central serous chorioretinopathy (SSCR), thyroid eye disease, uveal effusion syndrome, age-related macular degeneration, trauma, drug use, and hypotonic have been reported [3-12]. Idiopathic CFs is diagnosed when no etiologic factor is detected. Systemic research is essential for the identification of a specific etiology if folds are seen on the fundus examination. Patients may be asymptomatic, as well as with hyperopic or metamorphosis. In this article, we aimed to present a series of five cases of CFs with different etiologies.

Case 1

A 21-year-old male patient was referred due to low vision and intraocular pressure (IOP) in the right eye. The patient stated that he had been wounded with a wire in the right eye about thirty days ago and cataract developed because of the trauma. He stated that IOP was low during the postoperative examinations and that he was referred for this reason. In the right eye; the visual acuity (VA) was 6/10 with correction of +3.00 diopter (D), the IOP was 4 mm Hg, biomicroscopic examination revealed a penetrating line (Seidel negative), which was sutured to the temporal limbus and pseudo phobia, funds examination revealed a tortuosity increase in all vascular structures starting from optic disc, wrinkles and mauls in the per papillary region were observed. The left eye was completely normal. Axial length measurements performed with A-mode ultrasound showed 22.19 mm in the right and 22.70 mm in the left eye. Optical coherence tomography (OCT), ultrasonic biomicroscopy (UBM) and funds auto fluorescence (FAF) images were obtained.

As a result of the examinations, traumatic cyclodialysis induced hypotonia maculopathy was diagnosed for right eye. Topical
corticosteroid and cyclopentolate therapy was initiated, and 3 days later the right eye axis was measured as 22,80 and the refraction rate was measured as +0,75. The IOP returned to 18 mmHg and the VA was 10/10. Comparing the OCT images taken at the end of treatment with the first received OCT, the choroid, RPE and wavy appearance of the retina (Figure 1a). Were seen to be healed at the end of treatment (Figure 1b). Cyclodialysis was detected in the temporal quadrant (Figure 2a). By UBM and normal culinary anatomy was obtained at the end of the treatment (Figure 2b). There was a normal appearance at the end of the treatment (Figure 3a), while significant increase in tortuosity was observed in vascular structures at the first examination of the FOF (Figure 3b).

Figure 1: Right eye view (a) at first examination and (b) post-treatment OCT images.

Figure 2: a. Temporal quadrant UBM images of the right eye at first and b. At the end of treatment.

Figure 3: a. Right-eye at first and treatment outcome, b. FOF images taken at 30° magnification.

Figure 4: OCT images (a,b).
Case 2

A 53-year-old male presented with complaints of blurred vision. Patient who did not use glasses and had uncorrected VA were 0.7 in both eyes, with +3.00 D correction in both eyes the Varies up 10/10, and the IOP was 15 mmHg in both eyes. Anterior segment biomicroscopic examination was normal in both eyes. Light-dark horizontal lines extending from the optic disc to the macula were observed in both eyes in the funds examination. In OCT, CFs was observed in choroid, RPE and neurosensory layers in both eyes (Figures 4a & 4b). The patient’s FFA did not reveal any features other than CFs. Radiologic imaging studies for orbital, intracranial, and systemic pathologies were performed and the patient was diagnosed as CFs associated with an acute acquired hyperopic.

Figure 5: Fundus appearance in both eyes (a,b).

Case 3

A 50 year old male presented with complaints of blurred vision. The VA was 9/10 in both eyes with no refractive correction need. Anterior segment biomicroscopic examinations were of course normal in both eyes. IOPs were 14 mmHg in both eyes. In the fundus examination, several angioid streaks and light-dark colored bands were seen in the per papillary region in both eyes (Figure 5a & 5b). Wrinkles were observed in choroid, RPE and neurosensory layers in OCT (Figure 6a & 6b). In FFA, more prominent angioid streaks were observed around the optic disc in the left eye, and hyper fluorescence due to RPE cracks was observed (Figure 7a & 7b). Microperimetry (MP) revealed that sensitivity was normal in folding areas while sensitivity was decreased in areas with RPE defect (Figure 8a & 8b). Radiologic imaging studies and laboratory examinations for orbital, intracranial and systemic pathologies were within normal limits. CFs associated with angioid streaks was diagnosed.

Figure 6: OKT images (a,b).

Figure 7: FFA images (a,b).
Case 4

During the follow-up of an 81-year-old diabetic female, oblique choroid folds were detected outside the vascular arquades in both eyes. The VA was 8/10 in the right eye and 9/10 in the left eye. In biomicroscopic examination, posterior chamber intraocular lens was present in both eyes and IOPs were 13 mmHg bilaterally. In the fundus examination, moderate diabetic retinopathy findings and CFs were observed in both eyes. In OCT, RPE was observed to be fluctuating in the cross sections of the superior vasculature (Figure 9a & 9b). In FFA, CFs were observed in both eyes as light-dark lines with more obvious oblique outcrops than the upper main vascular arquade, and there were micro aneurysms occasionally due to diabetic retinopathy (Figure 10a & 10b).

Case 5

An 81-year-old male who was followed up for age-related macular degeneration, horizontal-extending CFs were noted in the posterior pole of the left eye. VA was in the right eye at the level of hand movement perception and at the level of 7/10 in the left eye. Biomicroscopic examination revealed pseudophakia in both eyes. IOPs were 15 mmHg in both eyes. In the fundus examination, peripapillaryatrophy, macular scar and RPE detachment were seen in the right eye (Figure 11a), with light-dark banding in the upper vascular quadrant, especially in the left eye (Figure 11b). In the right eye, wide RPE detachment and subretinal fluid were observed in the right eye (Figure 12a), while choroid and RPE wrinkles were observed in the left eye (Figure 12b). FFA revealed age-related neo vascular macular degeneration (Figure 13a). Horizontal extension of the CFs to the upper half of the macula and vein ligation around the optic disc were observed in the left eye (Figure 13b). Intravitreal ranibizumab therapy for the right eye of the patient was continued while left eye was taken in the direction of the CFs.
Discussion

CFs is not common, but they are easy to diagnose when they are seen on fundus examination because of their characteristic appearance. Clinically, the posterior pole is usually seen as diffuse light-dark banding. CFs may be parallel, radial or randomly distributed. Patients may be asymptomatic or may be presenting with metamorphosis or hypermetropia. The cause of visual impairment is distortion of the overlying photoreceptor layer, but in chronic cases the neurosensory retina and RPE may develop permanent damage. If it can be detected, treatment is directed to etiology. According to the currently accepted view, any intraocular or extrinsic event that disrupts the anatomical relationship of sclera with choroid and retina may cause CFs by affecting choroid, Bruch’s membrane, RPE and neurosensory layer. The CF is a kind of compensatory response given to the disruption of the normal spherical shape of the globe. Clinical examination is sufficient for the diagnosis of CFs, but FFA is the gold standard imaging technique.

Lately it has become easier to diagnose, thanks to OCT, which is increasingly used and allows visualization of the cross-sectional anatomy of the choroid and retinas. Another non-invasive imaging modality, B-mode USG, augments the etiologic investigation by showing thickening of the sclera and choroids, especially during inflammation and mass-induced tumors [13]. We also found that the UBM which we used in our first case and demonstrated cyclodialysis is invaluable in that it provides us with the opportunity to assess the anatomy of the anterior segment, is a non-invasive imaging method. As in our case, idiopathic CFs are usually bilateral in healthy hypermetropic individuals and best corrected visual acuity (BCVA) is preserved. Choroid tumours (melanocytes, choroid hemangiomas and stomas), mainly melanomas, and metastases
from organs such as the lungs and breast may be the cause of CFs. Regardless of whether it is malignant or benign, they may cause IOP rise and may result in CFs.

Cavernous hemangioma, lymphangioma, hemangiopericytoma, gloom, meningioma, rhadomyo sarcoma, lymphoma, fibrous histiocytoma, epithelial tumors of lacrimal gland, neuroblastoma and metastatic tumors are the examples. Intracranal masses of these types can cause hypermetropia and the extracranial ones induce astigmatism and cause refraction error. The collapsing materials used in the treatment of retinal detachment, radiotherapy implants, and implants used for orbital fractures may also cause CFs through globe compression [13]. Autoimmune connective tissue diseases such as posterior sclerotic, orbital inflammatory pseudo tumor, thyroid eye disease, rheumatoid arthritis, vacuities such as Wegener and Poliarteritis nodosa (PAN) are causes of inflammation-induced CFs. These patients may also present with pain, tenderness, proptosis, and lack of vision. It is noted in the literature that self-limiting CFs may develop after local supracroidal hematoma due to mass effect when it is not even solid [14].

CFs can be seen in the hypotoniamaculopathy caused by wound infiltration, excessive filtration after glaucoma surgery, and cyclodialysis. As seen in the first case, edema-like appearance in the optic disc and engorgement in vascular structures are noteworthy findings. With surgery for etiology or medical treatment as in our case, the fundus appearance usually returns to normal and visual acuity increases, but after extended hypo tony, the folding may persist. As an example of postoperative CFs in the literature, Liu et al. with 25 gauge Trans conjunctives alutureless vitrectomy for recurrent epiretinal membrane showed choroid detachment and CFs in a patient with hypo tony. After 3 months of treatment, the folds persisted and the VA increased to 20/40 from the hand movements [15]. Choroidal neovascular membrane due to age-related macular degeneration, spontaneously or through postlaser contraction, may cause CFs. Intracranial masses, pseudo tumor cerebri, benign intracranial hypertension and papilledema due to duralarteriovenous malformation may also be the cause of CFs. CFs may persist after papilledema treatment, but VA is rarely affected.

Talban et al. showed an increase in BCVA despite the persistence of CFs after tumor excision in a patient with parasellar meningioma [16]. Choroidal fold progression due to globe compression was reported due to medial wall injury of the orbital wall during nasal polypectomy [17]. It has been reported that in a patient with persistent choriorretinal adhesion due to trauma, the systemic steroid response was not available and that after 12 months the intravitreal injection of triamcinolone acetonide resulted in a level of VA of 0.2 [18]. CFs has been reported with pseudo vitally for Mmauculopathy, nanoftalmus, and micro ptalmus [19-21]. It has been reported in Vogt Koyanagai Harada’s syndrome, and even in the literature that this exacerbation period of the syndrome may be a leading indicator [22]. In some patients using topiramate, it has been reported that reversible macular folds can be the observed [23]. In diabetic retinopathy cases, some CF cases have

been reported. While CFs can be seen in diverse stages of diabetic retinopathy, they are usually asymptomatic, with symmetrical, macular, varying lengths.

The relationship between diabetic retinopathy and CFs is not well defined, and it is thought that about half of the cases can be seen in diabetic patients with increased arterial hypertension and increased choroidal thickness [24,25]. As a result, one of the five CFs was evaluated as traumatic hypertonic, one was associated with acute acquired hyperopic, one was associated with algioid streaks, one was associated with diabetic retinopathy, and one was associated with age-related macular degeneration. As seen in the literature and in our cases, the CFs which will keep the idiopathic cases out is not a primary pathology but rather a compensatory response given to many pathologic conditions which may or may not be related to the eye but distort the spherical shape of the eye. In such cases, detection of etiologic factor is important because it will determine the treatment. Every ophthalmologist should be aware of orbital or systemic pathologies that may cause chorioretinal folding. A careful history should be taken and medical history, inflammatory, eulastic, infectious and infiltrative diseases should be questioned diligently.

References


