INTERNATIONAL CONGRESS OF THE EGYPTIAN OPHTHALMOLOGICAL SOCIETY

In collaboration with:

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Ocular masquerade in adults: Challenges in diagnosis and management

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In our Practice

We encounter a myriad of unexplained signs in unusual situations

Opaque ocular media pause a diagnostic dilemma

Think Ocular Tumors

Case 1: 44 YOM

Spontaneous hyphema OD







► History:

- No bleeding tendency
- No trauma

Investigations:

Normal hematological profile



Management



Management





Hemorrhagic Lesions Simulating Melanoma

Case 2:

62 YOM
POAG on Rx
Rapid diminution VA OS x 2 weeks
HM /GP
PMH:
DM/HT

► IHD

OD: WNL
OS:
NS
IOP: WNL
Fundus: Vitreous hemorrhage







Suggested Management



- Clues for the decision:
 - Blood thinners
 - Uncontrolled HT
 - ► US features



Old standing choroidal lesion in opaque media

Case 3:

- ► 44 YOM
- OD: amblyopia
- Corneal opacity OD
- ► US:
 - Axial length +/- 24.5 mm
 - Pseudophakia
 - Elevated Mass temporal to ON involving the macular area with high initial reflectivity and internal irregula low to medium reflectivity with no RE
 - Size: 8.2 mm x 2.9 mm







Hypocellular proliferating astrocytes, H&E, x20



Tumor, H&E x40



GFAP, Immunohistochemistry x20

S100, positive, Immunohistochemistry x40.



SOX10 showing negative tumor with positive retinal covering, Immunohistochemistry x40

Reactive retinal astrocytic tumor

Vasoproliferative tumor of the ocular fundus

- Definition:
 - Elevated red-pink mass with no feeding artery and a draining vein
 - A mixture of glial cells and a network of fine capillaries with some larger dilated blood vessels
- Aetiology:
 - Primary: unilateral, solitary, IT (74%)
 - Secondary (26%):
 - Intermediate uveitis
 - Retinitis pigmentosa
 - Ocular toxocara
 - Coats' disease
 - Chronic retinal detachment
 - Ocular trauma
 - Inflammation

CP

- Third or fourth decade
- Both sexes are equally affected
 - Globular yellowish-pink vascular mass in the inferior peripheral retina
 - Lacking the feeder vessels typically seen in retinal capillary hemangioma
 - Sub-retinal exudation (80%)
 - Macular fibrosis and edema may lead to visual loss.



Peripheral Subretinal mass: Is it truly a choroidal lesion ??

Case 4

► 43 YOM

- Complicated cataract surgery
- Intraocular mass OD x 3 years
- ► VA: 6/60 (+14 D)





10 x 6 mm



MRI coronal: T1W: mildly hyperintense dome shaped nasal intraocular lesion

MRI: T1W with contrast: Enhancement





MRI axial: T1W: mildly hyperintense dome shaped nasal intraocular lesion



MRI: T1W with contrast: Enhancement



MRI: T2W: Hypointense lesion



Systemic workup

- CT chest: Normal
- Abdominal US: Normal

DD: Atypical choroidal lesion

Metastatic ?

- Hemangioma ?
- Amelanotic Melanoma ?
- **Other rare entities:**
 - Hemangiopericytoma
 - Schwannoma
 - Neurofibroma

Therapeutic options

- **Tumor thickness < 6mm:**
 - **Prachytherapy:**
 - Possible non-regression specially if benign
 - ▶ If non-regression \rightarrow enucleation
 - **Enucleation:** ?? Why
 - Surgical excision:
 - Nasal mass
 - Peripheral
 - Accessible
 - Young healthy individual (hypotensive anesthesia)
 - Adjust/replapce the IOL
 - No need for FNAB (20% error)





A well circumscribed mass hypercellular area shows bland spindle cells with whorling patterns. Antoni A (×200).

Schwannoma.



Postoperative: 1 year VA: 1.0





Unilateral Intraocular and Extraocular Lesions: Limit your DD

- ▶ 57 YOF
- DVA OS for 3 months









Lymphoma: 60 YOF



FNAB

Plasmacytoma in a 65 YOF



Choroidal Metastases

Commonest uveal malignancy (8-10% of pts with metastatic disease)

Uveal sites:

- Choroid: 88%
- ► Iris: 9%
- ► CB: 2%
- Primary site:
 - Breast 47%
 - ► Lung 21%
 - ► GI: 14%
- Unknown primary: 34%



Guidelines of FNAB

Rarely indicated for diagnosis
 Effective when indicated and properly managed
 Indications:

- Confirm uveal metastasis
- Resolve uveal melanoma vs metastasis
- Confirm uveal lymphoma
- Confirm uveal granuloma
- Confirm vitreoretinal lymphoma
- Limitations
- Uveal melanoma prognostication

Conclusion:

- Obscure Intraocular masses in adults pause a diagnostic dilemma
- Diagnsotic tools include US, MRI, PET CT when indicated
- Different approaches for diangosis and management
- Well equipped ophthalmic oncology centers
- Cooperation between ophthalmic oncologist, pathologist and general oncologist

