

POOR ANTI VEGF RESPONSE IN PATIENTS WITH EXUDATIVE AGE RELATED MACULAR DEGENERATION AND CANCER

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► Purpose

Exudative age related macular degeneration (EAMD) and cancer are common disease of the elderly and there are patients with cancer on treatment for EAMD.

Both these diseases are dependent on vascular endothelial growth factor (VEGF) for progression. Systemic VEGF production is required and augmented in cancer. It is possible that this increased production could augment EAMD and also lead to poor response to anti VEGF agents.

There is paucity in the literature documenting this association

► Methods

This was a descriptive study of consecutive patients presenting with EAMD and cancer or patients while on treatment for EAMD who developed cancer between 2012 and 2015, at the Canberra Retina Clinic, Canberra, Australia.

EAMD was confirmed with standard imaging studies that included fluorescein angiography and spectral domain ocular coherence tomography. The cancer type, stage, treatment and temporal relation to wet AMD presentation were obtained from case record.

Poor response to anti VEGF agents in EAMD was defined as persistence of sub-retinal and intra-retinal fluid on a monthly treatment schedule.

► Results

Six patients with wet AMD and co existent cancer was identified. Two patients had bilateral wet AMD. The male female distribution was 50%. The patient demographic data, imaging data, cancer type and treatment and EAMD subtype and anti VEGF agent used are detailed in table 1 and figure 1.

► Conclusion

Active cancer is associated with a poor response to anti VEGF agents in this group of patients with exudative age related macular degeneration. These results require further investigation.

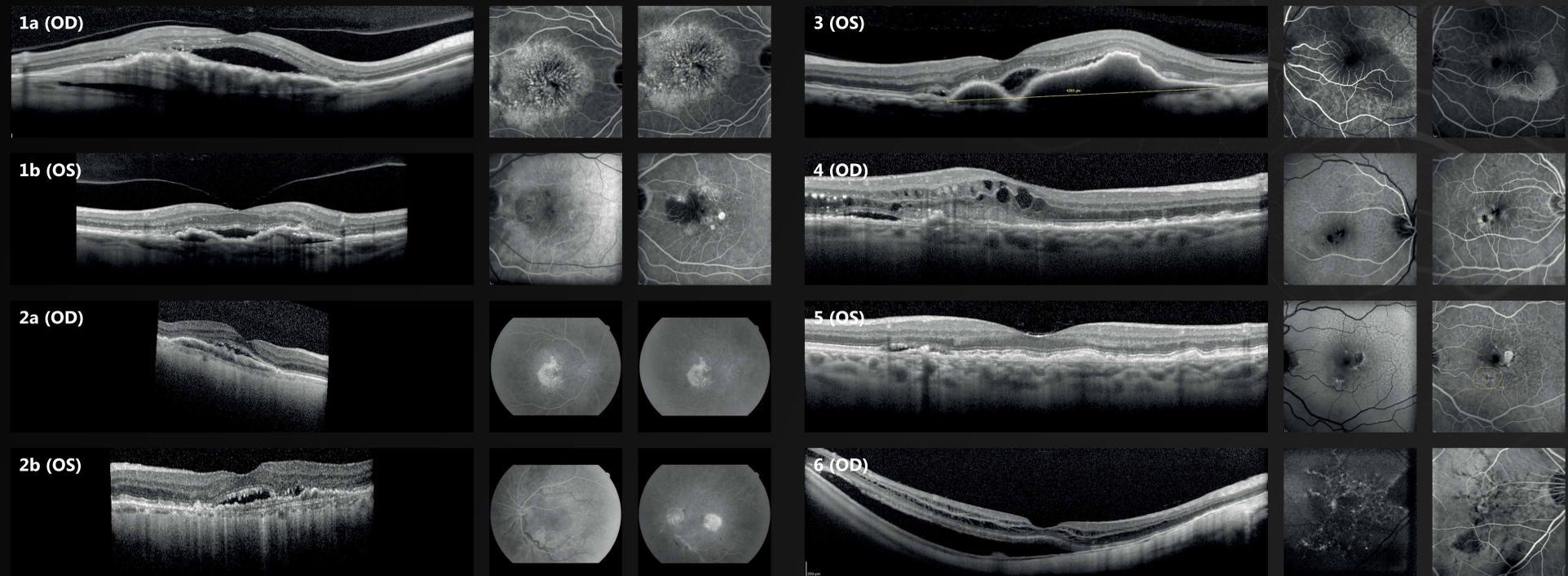


Table 1: Summary of cases

| Pt | Age | Sex | BCVA on presentation | | Latest BCVA | | Eye affected | Diagnosis | Type of cancer | Time interval between cancer diagnosis and AMD diagnosis or deterioration | Anti-VEGF treatment ¹ | No. of injections | Cancer treatment | Cancer outcome | AMD outcome |
|----|-----|-----|----------------------|-------|-------------|-------|--------------|------------------------------------------|-----------------------------------------------------------------|-----------------------------------------------------------------------------|----------------------------------|-------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------|------------------------------------------------------------------------------------------------------------------------------------|
| | | | OD | OS | OD | OS | | | | | | | | | |
| 1 | 80 | M | CF | 6/9-1 | 6/12-2 | 6/9 | OD | Exudative AMD with large subfoveal CNV | Grade 2 distal pancreatic non-functioning neuroendocrine tumour | 6 months | R | 15 | Distal pancreatectomy and splenectomy | In remission | Left eye developed exudative AMD 4 months later but with anti-VEGF treatment both eyes now have good control of exudative features |
| 2 | 71 | M | 6/15 | 6/6 | 6/15-2 | 6/12 | OD | Large central PED with CNVM progression* | Right lung T3N2M0 adenocarcinoma | 2 months | R | 20 | Planned for pneumonectomy but inoperable. Chemotherapy (gemcitabine, carboplatin, vinorelbine), radiotherapy and immunotherapy with nivolumab but ceased after a month | Metastasis of cancer | Left eye developed PED with serous component 13 months later Latest review – OD persistent SRF and PED but left eye nil SRF/RF |
| 3 | 60 | F | 6/5 | 6/12 | 6/6-2 | 6/9-1 | OS | PED + SRF + occult CNV | Left breast high grade ductal carcinoma in situ | 6 weeks | R | 4 | Wide local excision, sentinel lymph node biopsy and adjuvant radiotherapy. | In remission | Left eye trace subretinal fluid |
| 4 | 88 | F | 6/12 | 6/6-2 | 6/9 | 6/9 | OD | Subfoveal occult CNV | Right breast T2 N0 M0 grade 3 invasive ductal carcinoma | 6 months | R | 15 | Wide local excision, sentinel lymph node biopsy & adjuvant radiotherapy. Recurrence – mastectomy, adjuvant radiotherapy and endocrine therapy with letrozole | Possible liver metastasis | PED, nil SRF/RF |
| 5 | 74 | M | 6/9-1 | 6/6-2 | 6/6-3 | 6/9 | OS | Exudative AMD* | Stage yp T3 N3a M0 gastric adenocarcinoma | 3 months | R | 10 | Total radical gastrectomy, neoadjuvant chemotherapy (MAGIC chemoprotocol using eripubicin, cisplatin and fluorouracil) | In remission | PED and SRF |
| 6 | 83 | F | 6/12-2 | 6/9 | 6/12+2 | 6/6-1 | OD | Subfoveal CNV AMD* | Stage 3c melanoma | 2 months (between discontinuation of chemotherapy and vision deterioration) | R | 3 | Resection, axillary lymph node clearance. Local recurrence and metastatic disease a BRAF inhibitor vemurafenib but ceased after a month | Recurrence and metastatic disease | Reduction in SRF |

R, Ranibizumab; A, Aflibercept; TNM, tumour-node-metastasis system classifies cancer by the size and extent of the primary tumour (T), involvement of the regional lymph nodes (N), and the presence or absence of distant metastases (M); yp, pathological data following systemic therapy prior to surgery (American Joint Committee on Cancer, 2016).

*Patient had known stable AMD before sudden deterioration; ¹All patients received Ranibizumab and was then switched to Aflibercept except for patient 2 who received Ranibizumab, Aflibercept and then switched back to Ranibizumab and patient 4 who have received only Ranibizumab.