Pituitary apoplexy in setting of Dengue Hemorrhagic Fever with thrombocytopenia: case report and review of literature

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Abstract: Pituitary apoplexy is an acute clinical syndrome. It may occur spontaneously or as a result of several precipitating factors; one such factor being thrombocytopenia. Acute febrile illness accompanying with bleeding tendency is the main clinical feature of dengue. If the diagnosis is made in time, urgent treatment in the form of decompression of optic nerves may help to save vision. According to literature, only seven cases have been reported with pituitary apoplexy in setting of Dengue hemorrhagic fever. We report eighth case of Pituitary apoplexy in patient having Dengue hemorrhagic fever with its management and review of literature.

Key words: Dengue Hemorrhagic Fever, Pituitary apoplexy, Thrombocytopenia

Introduction

Pituitary apoplexy is an acute clinical syndrome characterized by sudden headache, vomiting, visual disturbances, ophthalmoplegia, altered consciousness, secondary to infarction or hemorrhage within a pituitary tumor or nontumorous pituitary gland. [1,2] The diagnosis is often missed in the early stages. CT and MRI of the brain are helpful in making the diagnosis of pituitary apoplexy. Pituitary apoplexy may occur spontaneously or as a result of several

precipitating factors; one such factor being thrombocytopenia.

Dengue is an infectious disease which can be seen worldwide. Acute febrile illness accompanying with bleeding tendency is the main clinical feature of dengue. [3] Dengue hemorrhagic fever (DHF) is characterized by fever, hemorrhagic tendencies, thrombocytopenia, and increased vascular permeability. [4] According to literature, only seven cases have been reported with pituitary apoplexy in setting of Dengue hemorrhagic

fever. We report eighth case of Pituitary apoplexy in the setting of Dengue hemorrhagic fever with its management and review of literature.

Case report

A 48 year old female presented with history of high grade fever 3 days. She was evaluated at local hospital where serology for Dengue infection was positive and she had low platelet count. Patient was referred to our hospital for further management. There was no history of petechial rash and spontaneous bleeding from mucosal surfaces. On third day of admission she complaint of sudden diminution of vision and severe headache. The platelet count decreased to 17,000. (Table 1)

Table 1 - Daily Platelet count

Platelets /mm ³)	On Admission	Day 1	Day 2	Day 3	Day 4	Day 5
Platelet count	72,000	50,000	22,000	17,000	33,000	131,000

NCCT Head was done which was suggestive of presence of sellar/suprasellar well defined heterogeneously hyperdense mass (25mm x 20mm x 18mm) with hyperdense area of bleed. (Figure 1)

Figure 1- NCCT Head Axial sections:
Sellar/suprasellar well defined heterogeneously hyperdense mass with hyperdense area of bleed

MRI brain showed well defined in homogeneously enhancing sellar- suprasellar mass with internal area of bleed. (Figure 2, 3 &4)

Figure 2 - MRI T2 WI Coronal Sections:
Heterogeneously hyperintense signal with presence of areas of bleed

Figure 3- MRI FLAIR Axial sections: Well defined extraaxial mass in sellar / suprasellar region measuring 2.6 x 2.3 x 2 cm in size

Figure 4 - MRI contrast enhanced Coronal sections:
Mild to moderate inhomogenous post contrast enhancement with hemorrhagic area

The optic chiasm was draped on its superior surface with no visualized area of in

between planes. On examination patient had Bitemporal hemianopia by confrontation method. Patient's hormone profile revealed decreased serum thyroid and prolactin hormone levels [Growth Hormone (GH) = 2.30 ng/ mL; Luteinizing Hormone (LH) = 2.64 mIU/mL; Prolactin (PRL) = 2.23 ng/mL; Cortisol = 6.53 µg/dL; Follicle Stimulating Hormone = 5.85 mIU/mL; T3 = 47.87 ng/dL; Thyroxine (T4) = 4.82 µg/dL; Thyroid Stimulating Hormone (TSH) = 1.220 µIU/mL]

Patient underwent several platelet units transfusion and was operated after platelet count reached above 100,000. Urgent Transnasal transsphenoidal decompression of the pituitary macroadenoma was performed. Intraoperatively, there was evidence of bleed inside the tumor Post operative period was uneventful and patient was discharged subsequently. Histopathology was consistent with Pituitary apoplexy with bleed. On three month follow up, patient had hypocortisolemia with no visual defect.

Discussion

Pituitary apoplexy may occur spontaneously or as a result of several precipitating factors such as head trauma, dynamic tests to evaluate pituitary function, surgery (mainly cardiac surgery), coagulation disorders, medications including aspirin, estrogens, heparin, and dopamine agonists and some conditions such as chronic systemic hypertension, diabetes mellitus, and radiotherapy. [1, 2, 5, 6]

The precipitants of pituitary apoplexy can be classified into 4 categories: (1) reduced

blood flow into the pituitary adenoma, resulting in infarction; (2) acute increase in blood flow to the pituitary gland from the hypothalamus portal system because it may increase the intra sellar pressure; (3) stimulation of the pituitary gland, as occurs in any stress inducing states; and (4) thrombocytopenia because of increased risk of bleeding. [5, 7]

Dengue fever is endemic in tropical countries and can range from a nonspecific febrile illness to DHF, which is characterized by high grade fever, marked thrombocytopenia ($<100,000/\mu\text{L}$), and increased vascular permeability. [4] In DHF, patients may have episodes of hypotension and hypovolemia, leading to oscillations in blood pressure. [4] Because it is an acute systemic illness, there is an increased demand on the pituitary gland and pituitary stimulation. In addition, DHF is characterized by severe thrombocytopenia and increased hemorrhagic tendencies. All these factors combined may have a compounding effect on the risk of pituitary apoplexy in a pre existing pituitary adenoma. However, the association of DHF and PA is exceedingly rare and only few case reports document their co occurrence. [6, 8, 9, 10, 11, 12] Other causes of visual deterioration in patients with dengue fever are optic neuropathy, maculopathy, retinal capillary occlusion, foveolitis, and retinal hemorrhage. [8, 13] The review of literature regarding the incidence of pituitary apoplexy in the setting of DHF revealed seven previously reported cases. The summary of these cases along with their comparative features has been described in Table 2. (Table 2)

Table 2 - Summary of the reported cases of pituitary apoplexy in setting of Dengue Hemorrhagic Fever

Reference	Age/ Sex	Clinical Feature	Platelet counts	Hormonal Profile	MRI findings	Management	Follow up
Kumar <i>et al</i> (2011)	31/F	Vision loss, H/A, vomiting	Hypothyroid, rest of the hormonal profile normal	Low (45,000/m ³)	Pituitary macroadenoma (16×22 mm) with acute bleed	Endoscopic endonasal decompression of the tumor	Residual bitemporal field defects at 3-month F/U
Wildenberg <i>et al</i> – Case 1 (2012)	40/M	H/A, vomiting, acromegalic features	Acromegaly; Hypogonadotropic hypogonadism	Mildly decreased (98,000/m ³)	Intrasellar lesion s/o – pituitary apoplexy	Transsphenoidal surgery 20 days after apoplexy	Acromegaly persists; other hormones normal
Wildenberg <i>et al</i> – Case 2 (2012)	38/M	Prolactinoma on medical Mx, presents with visual field deficits	Hypogonadism; Prolactin decreased on medical management	Mildly decreased (79,000/m ³)	Pituitary mass with areas of bleed	Urgent transsphenoidal surgery	Visual field defects improved. Hormonal deficits persist
Tan <i>et al.</i> (2014)	53/M	Acute-onset Lt third, sixth nerve palsy, Rt temporal hemianopia	Hyperprolactinoma; thyrotropin and gonadotropin deficiency	Low (16,000/m ³)	37×24×31 mm hemorrhagic pituitary adenoma with B/L cavernous sinus extension	Underwent transsphenoidal surgery for decompression (two surgeries)	Prolactin normal, anterior pituitary deficits and right eye visual defects persist
Mishra <i>et al</i> (2015)	43/M	Decreased vision, H/A, vomiting	Normal hormonal profile	Low (47,000/m ³)	23×21×20 mm sellar and suprasellar mass with bleed	Urgent transsphenoidal decompression of tumor	Visual acuity improved. Residual visual field deficits persist
Balaparameswara Rao <i>et al</i> (2016)	45/M	Severe H/A, vomiting and LOC	Normal hormonal profile	Low (27,000/m ³)	Pituitary apoplexy (sellar and suprasellar lesion) and obstructive hydrocephalus	EVDs to manage HCP; followed by endoscopic endonasal decompression	Vision normal. Anterior pituitary deficits present

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Varma et al (2016)	39 /F	Fever, melena, vision loss, altered sensorium	Hypothyroid, Hypogonadotropic hypogonadism	Low (11,000/m ³)	Pituitary adenoma with apoplexy	Urgent Decompression of Tumor	Symptoms Subsided
Our Case	48/F	Fever, H/A, Vision Loss	Hypothyroidism & Hypoprolactinoma	Low (17,000)	Pituitary apoplexy (sellar and suprasellar lesion)	Urgent transsphenoidal decompression of tumor	Visual acuity improved, hypocortisolemia

In our case we had a smooth outcome but as reported by Balaparameswara Rao et al, it can be stormy i.e. Fluid & electrolyte imbalance which can occur in pituitary surgery. In all cases Decompression of the tumour by transphenoidal technique was done and was successful. Vision improved in half of the cases.

Conclusions

Pituitary apoplexy can occur in setting of Dengue hemorrhagic fever. Dengue hemorrhagic fever should be considered as a cause of pituitary apoplexy. Transsphenoidal decompression of the tumor has good results in these cases. If the diagnosis is made in time, urgent treatment in the form of decompression of optic nerves may help to save vision.

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