Diabetes Insipides in a child disclosing localized Langerhans’ Cell Histiocytosis

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ABSTRACT

Langerhans cell histiocytosis (LCH) is monoclonal neoplastic condition of aberrant bone marrow histiocytes. The latter are part of the innate immune system and certain exogenous/endogenous stimuli may trigger its expansion. Hence LCH can present with limited or multiple organ involvement that may include; bones, lung, endocrine, skin, lymph nodes, spleen and bone marrow. In this case report, we describe a 3-year-old boy who presented with severe polyuria and polydipsia. Laboratory investigations were consistent with diabetes insipidus (DI). MRI of the brain; confirmed absence of the bright spot in its pituitary gland did not show evidence of tumor or enlargement by inflammation. Moreover, MRI revealed 2 skull lesions and their subsequent biopsy confirmed LCH. Systemic examination and tests including PET scan did not show additional lesions. Since his disease was localized, he received only Desmopressin acetate 120 ug twice daily for his DI without surgery, radiotherapy or chemotherapy. One year later, his disease remained limited to DI and the 2 bonny lesions.

Keywords: bone, diabetes insipidus, pituitary, desmopressin, Langerhans cell histiocytosis.

INTRODUCTION

Diabetes insipidus (DI) is a disorder characterized by polydipsia and polyuria due to lack or ineffective antidiuretic hormone (ADH). The latter is produced by the hypothalamus and stored in posterior lobe of pituitary gland (PG) 1. ADH is released, if serum hyperosmolality is sensed by the hypothalamus, to prevent urinary water loss 2. Hence, the disorder can result from; (a) cranial (PG/hypothalamus) disease and (b) nephrogenic resistance to its action. The most common causes of cranial DI are; (a) idiopathic (30%), (b) malignant or benign tumors of the brain or PG (25%), (c) cranial surgery (20%), and head trauma (16%). Rare etiologies include; genetic familial disorders, toxins, infections, sarcoidosis, hemochromatosis, amyloidosis, autoimmune hypophysitis and Langerhans cell histiocytosis (LCH) 3. The latter is an extremely rare neoplastic disorder of tissue histiocytes with variable phenotypic presentation from localized disease to an overwhelming malignant disease 4. In this case report, we describe a patient presenting with DI and localized bonny lesions and highlight his management.

CASE REPORT:

A 3-year-old boy presented with history of excessive thirst and urination including wetting his bed at night for 1 year. The patient did not have past history of significant medical illness, allergy or chronic intake of medications. He did not have similar problem in his family members. His initial physical examination did not show abnormality. Laboratory tests showed normal peripheral leucocyte and platelets counts. Hemoglobin was normal at 120 g/L. Serum sugar, urea, creatinine, electrolytes and liver function tests showed normal peripheral leucocyte and platelets counts. Hemoglobin was normal at 120 g/L. Serum sugar, urea, creatinine, electrolytes and liver function tests showed normal

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confirming LCH. PET-CT scan did not show additional lesions. He was treated with Minerin melt 120 ug twice
daily. Within few days, he showed significant improvement in his thirst, appetite, enuresis, activity and had gained
adequate weight. One week later; 24 hour urine volume as well as serum and urine osmolality were normal. Since his
DI was controlled and his bonny lesions were small and limited; he was not subjected to surgical excision of lesions,
radiotherapy or chemotherapy. One year later; he remained stable while on Minirin melt. PET-CT scan did not show new
bonny lesions or expansion of the previous ones.

Figure 1: Pre-contrast coronal T1-weighted MRI view showing normal size of the pituitary gland and its stalk (A). Note absence
of posterior pituitary bright spot in sagittal view (B).

Figure 2: Axial Views of MRI of brain T1FS+CFC showing contrast enhancing skull lesions in the (A) right fronto-parietal and (B)
left occipital areas.

Figure 3: Photomicrograph of the bonny lesion showing large histiocytes with reniform and grooved nuclei with indistinct
nucleoli and eosinophilic cytoplasm scattered among an infiltrate of lymphocytes, eosinophils, and macrophages (H&E X 400).
DISCUSSION

Our patient presented with polyuria with low urine osmolality and high serum osmolality indicating DI. The high serum osmolality excluded primary (psychogenic) polydipsia. Moreover, undetectable ADH and correction of polyuria, serum and urine osmolality after Desmopressin therapy confirmed diagnosis of cranial DI. In search for the etiology of cranial DI, MRI was done. In our patient, it confirmed; the absence of the “bright spot” which correlates with the secretory granules in the posterior lobe and ruled out abnormalities in the PG and its stalk induced by tumors and hypophysitis. Moreover, MRI revealed 2 incidental soft tissue masses in his skull. Biopsy of the occipital one disclosed LCH and ruled out bonny metastasis. LCH is a rare disease arising from immature myeloid dendritic cells in the bone marrow rather than from epidermal Langerhans cells. Evidence of its monoclonality was reported more than 20 years ago, supporting the notion that LCH is a neoplastic process. More recently, the identification of oncogenic BRAF V600E mutations in 25% to 64% of cases of LCH has provided additional evidence that LCH is a neoplasm. It occurs most often in children and in white individuals of northern European ancestry. The broad clinical spectrum that is encompassed by LCH is reflected in the many synonyms for this disease, which include eosinophilic granuloma (unifocal LCH), Hand-Shüller-Christian disease (multifocal unisystem LCH), and Letterer-Siwe disease (disseminated multifocal multisystem LCH). In single-system LCH, bone is the most common site of involvement followed by skin, lymph node, spleen, bone marrow and lung. Pulmonary LCH seems to be a distinct entity, as it occurs almost exclusively in smokers and may resolve with cessation of smoking. Liver, spleen, and bone marrow are considered “risk organs involvement” and are associated with high risk of mortality. Management of LCH is tailored to match the sites of involvement and extent of disease. Such strategy was based on the concept of disease induction and progression in relation to exogenous/endogenous stimuli since histiocytes are part of innate immunity system. For treatment purposes, patients with LCH are generally stratified into: (a) single-system or multisystem, and (b) high or low-risk organ involvement. Chemotherapy for patients with low-risk multisystem LCH is 1 year of vinblastine and prednisone compared to an additional mercaptopurine in the high-risk group. On the other hand; children with bonny lesions rarely need more than curettage and may even regress spontaneously. Our patient had central DI which is the most frequent CNS manifestation of LCH, occurring in 10-50% of all patients. However, and on follow up; 30-58% of those patients manifest anterior pituitary hormone deficiencies which seems to be linked to a thickening of the pituitary stalk. Moreover, they can also develop neurodegenerative disease. Our patient had cranial DI and 2 skull lesions. His physical examination, laboratory investigations, radiological and PET scans did not show evidence of other systemic disease. Since he had limited disease; we elected to treat him conservatively and avoid chemotherapy. Fortunately, and after 1 year of follow up; he did not manifest further expansion of his disease.

REFERENCES: