ABSTRACT

Neurofibromatosis (NF) consists of group of genetically inherited disorders all of which is characterized by occurrence of nerve sheath tumors. There are eight different subtypes of neurofibromatosis. Segmental NF, a rare mosaic pattern of NF-type1 (NF1) is diagnosed in individuals who have typical features of NF1 restricted to one part of the body and whose parents are both unaffected. But recently the nomenclature has been changed to mosaic NF.

We report a case of neurofibromatosis following the lines of blaschko. To the best of our knowledge, no such case report has been published till date of generalized mosaic NF following lines of blaschko's bilaterally symmetrically.

KEYWORDS

Blaschko, mosaic, neurofibromatosis, segmental
INTRODUCTION

Neurofibromatosis (NF) consists of group of genetically inherited disorders all of which is characterized by occurrence of nerve sheath tumors. The mode of inheritance is autosomal dominant with a high rate of new mutation and variable expression. There are eight different subtypes of neurofibromatosis. The prevalence of neurofibromatosis type 1 (NF1) is about 1 in 3000. The diagnosis of NF 1 is based on presence of ≥ 2 features out of the seven criteria. It is caused by heterozygous mutations of the NF1 gene located in chromosome 17q11.2.

Segmental neurofibromatosis is a rare mosaic pattern of NF1 which presents with typical features of NF1 restricted to one part of the body and whose parents are both unaffected. The presentation can be limited to a narrow strip of one quadrant to occasionally to whole half of the body.

Here, we report a case of 52 years old male who presented with freckling, multiple lentigines, café-au-lait macules (CALM) and neurofibroma in a blaschkoid pattern bilaterally, which to our knowledge has not been reported till date.

CASE REPORT

A 52 years old male came to dermatology out-patient department with Tinea cruris. But, on further examination for signs of fungal infection elsewhere, multiple café-au-lait macules and freckles along with lentigines were noted.

On further questioning, patient mentioned a few grouped soft nodules over the right abdomen. Patient was not exactly sure but claimed that both the pigmentary changes and the nodules were present for more than 25 years.

There was no history of similar skin lesion in other family members. No history of any bony abnormalities or seizures. Patient was literate but was not able to complete high school because of family problems. He was an overseas foreign worker currently placed in middle east.

No major illness or surgeries were noted in the past.

On examination, there were multiple lentigines and freckles following a pattern extending from the forearm, involving the axilla bilaterally and converging in the mid-abdomen. (Figure 1a) Similarly grouped lentigines following a linear pattern was also noted in the back. (Figure 1b). Multiple freckles were also noted in the axilla bilaterally. (figure 2)

Figure 1a: Multiple lentigines and freckles in the blaschkoid pattern over the trunk

Figure 1a and 1b: Multiple lentigines and freckles in the blaschkoid pattern over the trunk

Figure 2: Axillary freckling (arrowhead) and café-au-lait macules (arrow)

Figure 3: Multiple soft skin-colored nodules on right side of abdomen (arrow). On pressing with the index finger, the buttonhole sign was present.

(Note: the string is of religious significance)
Case Report

Ophthalmologic and neurologic examination did not reveal any abnormalities. Patient neither consented for any radiological examination nor biopsy of skin nodules.

The presence of axillary freckling and multiple CALMs made us label the case as NF 1, as per the diagnostic criteria. But from the presence of the freckles and CALMs along with the soft tumors in a blaschoid pattern, we made the diagnosis of generalized mosaic neurofibromatosis.

Patient was explained on the course and prognosis of the disease and was advised for regular follow-up.

DISCUSSION

Neurofibromatosis is classified into eight subtypes. Among these, NF type V is classified as segmental. Segmental NF is defined as presence of CALMs and neurofibromas to given area of the body with no crossing of midline, absence of any family history of NF and systemic symptoms. Segmental NF is further classified into subtypes which include true segmental, localized with deep involvement, hereditary and bilateral.

Segmental NF is an example of mosaicism, in which localized disease results from a postzygotic NF1 gene mutation. Localized disease is manifested if mutation occurs late in embryogenesis, whereas mutation occurring before tissue differentiation causes generalized disease.

But after recognition of the somatic mutation involved in these subtype, recent suggestion is to use correct term as “mosaic” NF rather than type V or segmental. In 2001, Ruggieri and Huson divided neurofibromatosis with somatic mutation into 2 distinct subsets mosaic localized and mosaic generalized based specifically on the time of mutation.

LIMITATION OF THE STUDY

A skin biopsy which is diagnostic of neurofibroma could not be done. An ultrasound of abdomen and pelvis as well as a chest x-ray could have shown any internal organ involvement which was also not done because patient did not agree for any investigation.

CONCLUSION

Mosaic generalized neurofibromatosis is a rare variant. It should be suspected in cases which presents with unusual presentation of neurofibromas. A better nomenclature of mosaic neurofibromatosis should be encouraged over the previously used segmental neurofibromatosis.

CONFLICT OF INTEREST

none

REFERENCES


Parajuli N et al.