



Long term follow up of recurrent monostotic fibrous dysplasia of humerus treated with non vascularised allogeneous fibula graft

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Abstract : Fifteen year old student who had monostotic fibrous dysplasia at the age of 6 years treated with cancellous bone graft ends up with recurrence. Which was treated with allogeneous fibula strut graft from his mother and followed up for 8 years with excellent functional outcome with radiological incorporation of graft with host bone with no evidence of recurrence.

Keyword : Monoostotic fibrous dysplasia, Allogeneous fibula graft. Bone tumour

Introduction :

Fibrous dysplasia is a benign skeletal disorder of adolescents and young adults in which the medullary canal is replaced and weakened by fibrous tissue⁴. It may affects single bone or multiple bones. Management is mainly to prevent pain ,limit deformity and treat pathological fractures. Curettage and cancellous bone grafting have been used to replace the dysplastic fibroosseous tissue with normal bone, but this is associated with high rate of recurrence¹¹ which can be prevented by use of cortical autograft or allograft. The purpose of the study is to review the long term follow up of clinical , functional, and radiographic results of using intramedullary allogeneous strut graft for large humeral defects created after excision of fibrous dysplasia.

Materials and methods :

15 year old male student with monostotic fibrous dysplasia of left humerus(figure 1 a) at the age of six years presented to our hospital for which through deltopectoral approach curettage and allogeneous cancellous graft taken from his mother's iliac crest used for grafting(figure 1 b) . Histopathological examination showed evidence of fibrous dysplasia. The lesion recurred in one year(figure 2 a) .



fig 1 a- Lytic lesions initial radiograph showing fibrous dysplasia

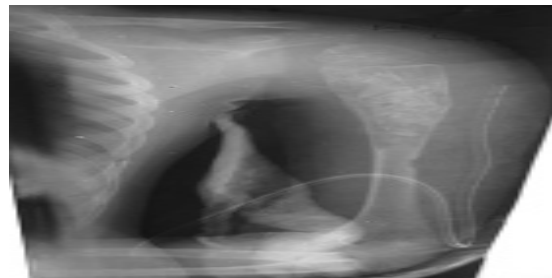
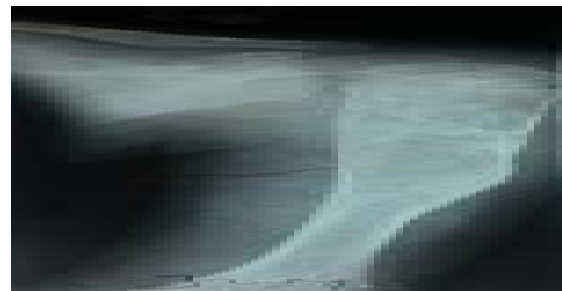


fig 1 b- Immediate post op after curettage and cancellous bone graft fig 2 a- Lytic lesion in proximal humerus



Routine investigations normal. X –ray left arm show lytic lesion in medulla representing recurrence of fibrous dysplasia(figure 2 a).

Operative technique :



Under general anaesthesia patient in supine position through anterolateral approach, pathological bone excised and defect found to be more. Fibula graft taken from his mother used as cortical strut with excess bone telescoped to normal medullary canal and fixed with three 3.5 cortical screws (figure 2 b).

figure 2 b- Immediate post op picture with allogeneous fibula graft

Post operatively patient kept in splint. Suture removal done on 12th day. Passive motion started on 3rd day. Active mobilisation started at 6 weeks. Follow up done with clinical and radiography showed excellent results with good range of movements in shoulder, elbow and hand.

Results :

Eight year follow up shows complete range of movements in shoulder elbow and hand comparable to his normal limb with no pain or deformity (figure 3 a & figure 3 b). Radiography shows incorporation of allograft to host bone with consolidation (figure 2 c & figure 2 d).

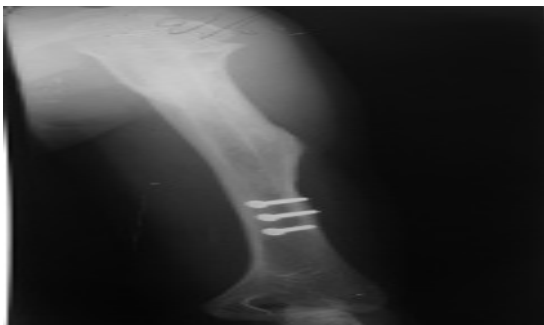


figure 2 c-Two years post op picture with incorporation of allogeneous fibula graft with borders seen



fig 2 d-8 year follow up with incorporation of graft with host bone
fig 3 a- Clinical picture good range of movements
fig 3 b- Clinical picture with good range of movement



Discussion :

Fibrous dysplasia is a benign, pathological condition characterised histologically by poorly oriented osseous trabeculae weakened by replacement with fibrous tissue. Curettage and cancellous bone graft (normally takes place by creeping substitution) causes replacement of graft with poorly formed woven bone which can cause recurrence as in our case. Allogeneous bone graft taken from his mother provides structural stability and also as a cortical graft for healing¹. We conclude that fibrous dysplasia treated with allogeneous non vascularised fibula graft from mother incorporates well in children, providing both mechanical and graft benefits with no immunological reactions and no recurrence.

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