Introduction

Primary splenic lymphoma is a very unusual entity. According to guidelines diagnosis of Primary Splenic Lymphoma should be made when the disease is confined to spleen or at the most involves hilar lymph nodes with no recurrence of disease after splenectomy [1]. Herein, we present such an unusual case of low grade b- cell non-Hodgkin primary Splenic Lymphoma and confirmed by immunohistochemistry in a patient presenting with massive splenomegaly and hypersplenism.