中文題目： 瀰漫性大Ｂ細胞淋巴瘤合併嗜血症候群的個案報告

英文題目：A case of diffuse large B cell lymphoma with initial alarm sign: Hemophagocytic syndrome

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Abstract Hemophagocytic lymphohistocytosis(HLH) is rare but serious phenomenon in clinical practice. Hence, rapid study and prompt treatment are essential to patients. We present a case diagnosed as diffuse large B cell lymphoma primary involvement in bone marrow with initial presentation as HLH. During dealing with this case, initial diagnosis and decision of treatment are two major dilemmas.

Introduction HLH is a presentation of impaired immune system. Primary HLH predominantly manifest in infant and are most caused by infectious agents. Secondary HLH is triggered by underlying disease including malignancies, infection, autoimmune disease and acquired immune deficiency. As to malignancies, most are caused by leukemia or lymphoma. Most are T cells entities, and there is low incidence of B cell lymphoma. Primary bone marrow large B cell lymphoma is rare and the prognosis is poor.