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Apr 18, 2014 · *Epidemiology* Acquired hemophilia A (AHA) is a rare but clinical significant entity. The reported incidence ranges between 1.20 and 1.48 cases per million/years and was reported from three cohorts: a large Australian hemophilia center cohort collected over 12 years (1997–2008) based in South Australia, a cohort of patients diagnosed with AHA in South and West Wales between 1996 and 2002, ...

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Myelofibrosis and acquired hemophilia A: a case report

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AHA should be suspected immediately in patients who have a malignant disease, bleeding, and isolated prolongation value of APTT. Surgery could be a precipitating cause of acquired hemophilia. This syndrome is remarkable for its abrupt onset within days of surgery, dramatic bleeding, subsequent persistence, but potential reversal by ...

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Acquired hemophilia A: a case report – itjem

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Oct 17, 2017 · Abstract. Acquired hemophilia A (AHA) is a rare bleeding disorder with an incidence of approximately 1.5 cases/million/year¹ and is characterized by autoantibodies directed against circulating coagulation FVIII. These autoantibodies constitute the most common spontaneous inhibitor to any coagulation factor and may induce spontaneous bleeding in patients with no previous history of a ...

Myelofibrosis and acquired hemophilia A: a case report

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- What is acquired hemophilia? ▾
- How rare is acquired haemophilia? ▾
- How is hemophilia classified? ▾
- Is hemophilia A recessive or dominant genetic disorder? ▾

Name of Journal: *World Journal of Clinical Cases*

Manuscript NO: 61887

Manuscript Type: CASE REPORT

Acquired haemophilia in patients with malignant disease: A case report

Acquired haemophilia in malignancy

Abstract

BACKGROUND

Acquired haemophilia is a rare coagulation disorder characterized by autoantibodies against coagulation factor VIII leading to severe and potentially life-threatening haemorrhages. The underlying disorder causing the development of an autoimmune phenomenon is not always known, however, 10-15% could be linked to malignancies. Patients with cancer who require surgical resection represent a treatment challenge not solely due to increased risk of bleeding but also due to adverse events of immunosuppressive therapy.

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The development of factor VIII inhibitors in non-hemophilic patients is rare and may occur in healthy individuals, mostly elderly and women in postpartum period, and in patients with malignant neoplasia