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Name of Journal: *World Journal of Clinical Cases*

Manuscript NO: 55851

Manuscript Type: CASE REPORT

Primary myelofibrosis with concurrent *CALR* and *MPL* mutations: A case report

Zhou FP *et al.* Co-mutation of *CALR* and *MPL* genes

Feng-Ping Zhou, Cheng-Cheng Wang, Hua-Ping Du, Shan-Bo Cao, Jin Zhang



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Primary autoimmune myelofibrosis: a case report and review of the literature. ... Autoimmune myelofibrosis is a rare, distinct clinicopathological entity that can occur in isolation (primary) or in association with systemic autoimmune disorders (secondary), such as systemic lupus erythematosus and Sjogren's syndrome. ... CALR, and MPL mutation ...

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Author: Ayalew Tefferi

Publish Year: 2018

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Abstract. **Somatic mutations** in **CALR gene** have been reported in 60%–88% of patients with **essential thrombocythemia (ET)** and **primary myelofibrosis (PMF)** who are negative for JAK2 and **MPL mutations**. Most of the **CALR mutations** analyzed to date are **heterozygous mutations** in **exon 9** of the gene.

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