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Case Control Study

Relationship between granulomatous lobular mastitis and methylene tetrahydrofolate reductase gene polymorphism

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Abstract

BACKGROUND

Variations in the methylene tetrahydrofolate reductase (*MTHFR*) gene have been reported as risk factors for numerous conditions, including cardiovascular disease, thrombophilia, stroke, hypertension and pregnancy-related complications. Moreover, it was reported there is an association between breast

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3. Discussion. In 1972, Kessler and Wolloch first defined **granulomatous mastitis** (GM) [1, 2]. GM can be idiopathic (primary) and specific (secondary) [2, 3]. Secondary GM involves caseation necrosis and emerges with a variety of infectious conditions such as vasculitis, sarcoidosis, tuberculosis, actinomycosis, and blastomycosis filariasis [2, 3]. IGM is detected in less than 1% of breast ...

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An association was also reported **between** breast cancer and mutations in MTHFR-C 677 T, PAI-1, ACE genes. Genetic **polymorphisms** may involve in the development of IGM as it was seen in our case.

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diagnosed with IGM [1]. It is also called idiopathic **granulomatous lobular mastitis** or idiopathic **granulomatous lobulitis**. It is more frequent **between** 2-4 decades in reproductive ages and in the first 6 years after childbirth [3,4]. It can also rarely be seen in men ...

Author: Sabahattin Destek

Publish Year: 2018

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Granulomatous lobular mastitis is a rare chronic breast disease, firstly described by Kessler and Wolloch in 1972. In this article we present a 35-year-old patient with **granulomatous lobular** ...

Idiopathic granulomatous mastitis: a disease mimics breast ...



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3. Discussion. In 1972, Kessler and Wolloch first defined **granulomatous mastitis** (GM) [1, 2]. GM can be idiopathic (primary) and specific (secondary) [2, 3]. Secondary GM involves caseation necrosis and emerges with a variety of infectious conditions such as vasculitis, sarcoidosis, tuberculosis, actinomycosis, and blastomycosis filariasis [2, 3]. IGM is detected in less than 1% of ...

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It is also called idiopathic **granulomatous lobular mastitis** or idiopathic **granulomatous** lobulitis. It is more frequent **between** 2-4 decades in reproductive ages and in the first 6 years after childbirth [3,4]. It can also rarely be seen in men . The etiology of the IGM is not fully understood.

Author: Sabahattin Destek

Publish Year: 2018

[CASE STUDY 7.1 CHRONIC MASTITIS ANSWERS](#)

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Case diagnosis was based on chronic findings, after extensive work-up ruled out malignancy and known causes of **granulomatous mastitis**. Eleven patients representing 1. Associations **between** single nucleotide **polymorphism** SNP genotypes with CMB and breeding values for milk and protein yield were



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Idiopathic **granulomatous mastitis** (IGM) is a rare and chronic inflammatory disorder. ... variants in the **methylene tetrahydrofolate reductase** (MTHFR) and thymidylate synthase (TS) **genes** may be ...

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We studied the -930A/G **polymorphism** of the CYBA **gene** promoter, the apolipoprotein E (APOE) genotype and the **methylene-tetrahydrofolate reductase** (MTHFR) **gene** C677T **polymorphism** in ...

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Jan 01, 2016 - The enzyme **methylene tetrahydrofolate reductase** generates 5-**methyl tetrahydrofolate** to serve as a carbon donor for methionine synthase. Homozygous deficiencies of certain **genes** involved in methionine metabolism including cystathionine synthase or **methylene tetrahydrofolate reductase** results in the condition homocystinuria.

Author: J.R. Stone **Publish Year:** 2016

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Aim of the work To detect **methylene-tetra-hydrofolate reductase** (MTHFR) 677C/T and 1298A/C **gene** polymorphisms in RA patients treated with MTX and to investigate the **relationship** with serum OPN ...

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Title: Association **between** the **methylene tetrahydrofolate reductase gene** C677T mutation and