

**Clinical and molecular studies of patients with characteristics of Opitz G/BBB syndrome shows a novel MID1 mutation**

Hsieh EWY, Vargervik K, Slavotinek AM.

American journal of medical genetics. Part A

2008; 146A(18):2337-2345

**ARTICLE IDENTIFIERS**

DOI: 10.1002/ajmg.a.32368

PMID: unavailable

PMCID: not available

**JOURNAL IDENTIFIERS**

LCCN: 2004212588

pISSN: 1552-4825

eISSN: 1552-4833

OCLC ID: 51644587

CONS ID: not available

US National Library of Medicine ID: not available

This article was identified from a query of the SafetyLit database.