

## Gene Section

### Short Communication

# MDS2 (myelodysplastic syndrome 2 translocation associated)

Jean-Loup Huret, Sylvie Senon

Genetics, Dept Medical Information, UMR 8125 CNRS, University of Poitiers, CHU Poitiers Hospital, F-86021 Poitiers, France (JLH, SS)

Published in Atlas Database: July 2003

Online updated version: <http://AtlasGeneticsOncology.org/Genes/MDS2ID476.html>  
DOI: 10.4267/2042/37984

This work is licensed under a Creative Commons Attribution-Noncommercial-No Derivative Works 2.0 France Licence.  
© 2003 Atlas of Genetics and Cytogenetics in Oncology and Haematology

## Identity

HGNC (Hugo): MDS2

Location: 1p36

Location (base pair): 23024



Probe(s) - Courtesy Mariano Rocchi, Resources for Molecular Cytogenetics.

## DNA/RNA

### Description

Spans 13 kb; 7 exons.

### Transcription

4 alternate splicings; major splice: 281 bp, exons 1, 4, 6, 7 are constantly used.

## Protein

### Description

The 4 alternate splicings give rise to 2 proteins, of 82 and 140 amino acids.

### Expression

Wide.

## Implicated in

### *t(1;12)(p36;p13) in myeloid disorders*

#### Prognosis

Unknown so far.

#### Hybrid/Mutated gene

5' ETV6 - 3' MDS2.

#### Abnormal protein

Truncated ETV6.

## References

Odero MD, Vizmanos JL, Román JP, Lahortiga I, Panizo C, Calasanz MJ, Zeleznik-Le NJ, Rowley JD, Novo FJ. A novel gene, MDS2, is fused to ETV6/TEL in a *t(1;12)(p36.1;p13)* in a patient with myelodysplastic syndrome. *Genes Chromosomes Cancer*. 2002 Sep;35(1):11-9

*This article should be referenced as such:*

Huret JL, Senon S. MDS2 (myelodysplastic syndrome 2 translocation associated). *Atlas Genet Cytogenet Oncol Haematol*. 2003; 7(3):174.