

# Leukaemia Section

## Short Communication

### t(11;19)(q13;p13) FSTL3/CCND1

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Published in Atlas Database: February 2015

Online updated version : <http://AtlasGeneticsOncology.org/Anomalies/t1119q13p13FSTL3-CCND1ID1527.html>

Printable original version : <http://documents.irevues.inist.fr/bitstream/handle/2042/62528/02-2015-t1119q13p13FSTL3-CCND1ID1527.pdf>

DOI: 10.4267/2042/62528

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#### Abstract

Review on t(11;19)(q13;p13) FSTL3/CCND1, with data on clinics, and the genes involved.

#### Keywords

t(11;19)(q13;p13); FSTL3; CCND1; chronic lymphocytic leukemia

#### Clinics and pathology

##### Disease

B-cell chronic lymphocytic leukemia (B-CLL)

##### Epidemiology

Only one case to date, a 67 year-old female patient (Rimokh et al., 1993; Hayette et al., 1998).

##### Prognosis

No data

#### Cytogenetics

##### Cytogenetics morphological

Complex karyotype.

#### Genes involved and proteins

##### CCND1

##### Location

11q13.3

##### Protein

Binds and activates the G1 cyclin dependent kinases; Phosphorylates SMAD3 and inhibits its transcriptional activity and antiproliferative

function.

##### FSTL3

##### Location

19p13.3

##### Protein

Binds activin A and with lower affinity several other members of the TGF beta family; interacts with ADAM12, FN1 (fibronectin type 1.), and MLLT10 (Grusch, 2012).

#### Result of the chromosomal anomaly

##### Hybrid gene

##### Description

The breaks occurred at nucleotide 2276 in the 3' untranslated region of the last CCND1 exon, and about 7 kbp upstream of FSTL3.

#### References

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*This article should be referenced as such:*

Huret JL. t(11;19)(q13;p13) FSTL3/CCND1. Atlas Genet Cytogenet Oncol Haematol. 2016; 20(3):162-.