

# Solid Tumour Section

## Short Communication

### t(2;17)(p23;q23)

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## Clinics and pathology

### Disease

Inflammatory myofibroblastic tumors.

### Clinics

Rare soft tissue tumour found in children and young adults.

### Pathology

Spindle cell proliferation with myofibroblastic differentiation and an inflammatory infiltrate.

### Prognosis

Low malignant potential and good prognosis.

## Genes involved and proteins

### ALK

#### Location

2p23

#### Protein

1620 amino acids; 177 kDa; glycoprotein (200 kDa mature protein) ; membrane associated tyrosine kinase receptor.

### CLTC (*clathrin heavy polypeptide*)

#### Location

17q23

#### Protein

1675 amino acids, 191 kDa; mediate endocytosis of transmembrane receptors.

## Result of the chromosomal anomaly

### Hybrid Gene

#### Description

5' CLTC - 3' ALK

### Fusion Protein

#### Description

2196 amino acids. 1634 N-term amino acid from CLTC, containing nearly all of CLTC, including the triskelion assembly domain, fused to the 562 C-term amino acids from ALK (i.e. the entire cytoplasmic portion of ALK with the tyrosine kinase domain).

## References

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