Open Peer Review on Qeios

T(10;17)(q22;p13)

National Cancer Institute

Source

National Cancer Institute. t(10;17)(q22;p13). NCI Thesaurus. Code C127049.

A cytogenetic abnormality that refers to the translocation of the long arm (q22) of chromosome 10 and the short arm (p13) of chromosome 17. It results in an YWHAE-FAM22 fusion. It has been described in high grade endometrial stromal sarcomas and a subset of clear cell sarcomas of the kidney.