FISH(fluorescence *in situ* hybridization)를 이용하여 분석한 크롬에 의해 유발된 염색체 이상

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Detection of chromosomal rearrangements by chromium in human lymphocyte using fluorescence in situ hybridization(FISH) with triple combination of composite whole chromosome specific probe

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Abstract

Chromosome rearrangements induced in human lymphocyte after in vitro exposure to chromium were analysed by the use of fluorescence in situ hybridization(FISH) with triple combination of composite whole chromosome-specific probe for chromosome 1, 2 and 4. Chromosome aberrations was scored by the Protocol for Aberration Identification and Nomenclature Terminology(PAINT).

Stable translocation was the most frequent type of aberrations and dicentrics and insertions were also observed. Chromium treatment enhanced the frequencies of stable translocations and color junctions in a

dose-dependent manners, but no distinct increase of dicentrics and insertions was seen.

The ratio of the yields of translocation to the yields of dicentric varied between 13 to 27.

The presents results demonstrate fluorescence in situ hybridization(FISH) are useful for detecting chromosomal rearrangements induced by chromium.

key words: chromium, fluorescence *in situ* hybridizaton(FISH), chromosome rearrangements