

MRP1 polymorphisms (T2684C, C2007T, C2012T, and C2665T) are not associated with multidrug resistance in leukemic patients

F. Mahjoubi, S. Akbari, M. Montazeri and F. Moshyri

Clinical Genetics Department, National Institute of Genetic Engineering and Biotechnology, Tehran, Iran

Corresponding author: F. Mahjoubi E-mail: Frouz@nigeb.ac.ir

Genet. Mol. Res. 7 (4): 1369-1374 (2008) Received September 12, 2008 Accepted October 13, 2008 Published December 9, 2008

ABSTRACT. One of the main problems in treating cancer patients is that cancer cells can develop drug resistance. Resistance to multiple anticancer drugs, so called multidrug resistance (MDR), most likely involves a nonspecific mode of resistance, through drug-efflux transporters. One of the most extensively studied genes involved in MDR is multidrug resistance protein 1 (MRPI). We investigated a possible association between the expression level of MRPI and the occurrence of MDR in leukemic patients, and we tested the hypothesis that MRPI polymorphisms are predictive of MDR in patients with acute leukemia. The mRNA level of MRPI was determined in 111 patients with acute leukemia (including 52 patients with acute myeloid leukemia and 59 patients with acute lymphoblastic leukemia), by quantitative real-time PCR, to determine how it af-

fected the response to chemotherapy. We typed T2684C, C2007T, C2012T, and C2665T *MRP1* polymorphisms in 111 patients classified as either drug-resistant or drug-responsive. We found that high expression of *MRP1* was associated with the MDR phenotype in both acute myeloid leukemia and acute lymphoblastic leukemia patients. There was no effect of a particular genotype on the expression level of the *MRP1* gene. We found no significant differences in chemosensitivity among any of these genotypes.

Key words: Multidrug resistance protein; Multidrug resistance; Single nucleotide polymorphism