analysis revealed that the downstream rfaZ gene consists an operon with the rfaY. SoxS bound upstream of the -35 box and directly activated rfaY as judged by gel shift and in vitro transcription assay. To elucidate the function of rfaYZ genes in oxidative stress response, we disrupted rfaYZ operon. The DrfaYZ mutant was more sensitive to the oxidative stress. In addition, SoxR became more sensitized in the mutant to the quinone-type redox cycling agents, such as menadione and plumbagin. When DrfaYZ mutation was introduced into rpoHp3-lacZ single copy fusion strain, expression from rpoHp3 promoter increased 5.4 fold, implying rfaYZ mutation cause extracytoplasmic stress. The phenotype of SoxR-sensitization in the mutant was complemented by overexpression of the rfaYZ genes. However, increase in rpoH expression was not rescued even with the overexpression of rfaYZK genes.

F801

(GT)n Repeats Polymorphism of the Third Intron of T Cell Receptor Delta (TCRD) among Koreans

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The gamma/delta T cells recognize unprocessed or native antigens independently of the MHC, rapidly enhancing the immunity defense mechanism. The gamma/delta T cells have cytotoxic effects and produce large amounts of Th1 cytokines. The TCRD is located on the chromosome 14q11.2 and consists of three variables, three diversities, and three joining and constant genes. We analyzed the (GT)n repeats polymorphism of the third intron of TCRD using 6% denaturing PAGE in two hundred thirty-one unrelated Korean subjects. The number of GT repeat was

estimated for repeat polymorphism by auto-sequencing. Six alleles ranged in size from TCRD*13 (116 bp) to TCRD*19 (128 bp). The allele frequencies, TCRD*13, TCRD*15, TCRD*16, TCRD*17, TCRD*18, and TCRD*19 were 0.002, 0.424, 0.273, 0.026 0.253, and, 0.022 respectively. No deviation from the expectations according the Hardy-Weinberg equilibrium was found. The heterozygosity and polymorphism information content were 0.68 and 0.62. The highest allele frequency among Koreans was TCRD*15 (0.424), while TCRD*18 (0.308) among Caucasians.

F802

Single Nucleotide Polymorphisms in the Non-coding region of *CYP2E1* among Koreans

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Cytochrome P4502E1 (CYP2E1) is a major component of the microsomal ethanol-oxidizing system involved in the metabolism of a variety of fcreign compounds including carcinogens. CYP2E1 has been implicated in alcohol-related liver diseases and several forms of cancer because of its contribution to oxidative stress. From the genomic DNA of two hundred forty-eight Koreans who are not relatives, single nucleotide polymorphisms at -1293 and -1053 nucleotide of the 5' flanking region and at 7623 nucleotide of intron 6 of CYP2E1, which are contribute to regulation of CYP2E1 expression, were screened using PstI, RsaI and DraI RFLP. The genotype frequencies of CYP2E1*c1/*c1, CYP2E1*c1/*c2 and CYP2E1*c2/*c2 of PstI/RsaI RFLP in the 5' flanking region were 69.8%, 25.8%, and 4.4% respectively; the allele frequenc CYP2E1*c1 was 0.827; the genotype

frequencies of CYP2E1*C/*C, CYP2E1*C/*D and CYP2E1*D/*D of DraI RFLP in intron 6 were 3.2%, 25.8% and 71.0% respectively; the allele frequency of CYP2E1*D was 0.839;and the heterozygosities of PstI/RsaI and the DraI SNPs of CYP2E1 were both at 0.258. The test for Hardy-Weingberg equilibrium showed significant deviation from expectations. The allele frequencies in the 5' flanking region and intron 6 of CYP2E1 among Koreans were proven to be similar to those of other Oreintals, and the allele frequency of CYP2E1*c2 in the 5' flanking region in Caucasians is lower than that of Orientals including Koreans.

F803

Three Tetranucleotide Repeat Polymorphism at D16S539, D7S820 and D13S317 Loci in Korean Population

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The markers of microsatellite are polymorphic and are mostly based on loci with tetranucleotide repeats, each locus commonly having between 5 and 15 alleles. Allele and genotype frequencies of D16S539, D7S820 and D13S317 were analyzed using PCR and denaturing polyacrylamide gel electrophoresis followed by silver staining visualization. DNA extracts were obtained from about 80 unrelated Koreans. The frequency of allele 11 at D165539 locus in Korean was the lowest among some racial populations. The frequency of allele 11 at D13S317 in Korean was similar to that of Asian populations. Comparing the results of present study with other racial populations, allele frequencies of three loci were different according to the loci. This study confirms that the irregularity in distribution of microsatellite alleles in different populations

with the predominance of two or three alleles on these three investigated microsatellite loci.

F804

A Dinucleotide Repeat Polymorphism at the Calcium Sensing Receptor (CASR) Locus in Korean Population

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Calcium sensing receptor (CASR) in parathyroid gland regulates ca'cium homeostasis by sensing decreases in extracellular calcium levels and affecting an increase in secretion of parathyroid hormone and calcium reabsorption in the kidney. We analyzed dinucleotide repeat polymorphism at CASR locus in normal individuals and osteoporosis patients by polymerase chain reaction and polyacrylamide gel electrophoresis. Eight alleles were detected and allele frequencies of A3 and A9 were 0.46 and 0.49 in the normal individuals. In osteoporosis patients, allele frequencies of A3 and A9 were 0.48 and 0.46, respectively. The observed heterozygosities were 0.552 in normal sample and 0.503 in osteoporosis one. Allele distribution and frequencies of normal group were similar to those of osteoporosis one in Koreans. Comparing this result with a Japanese population, some differences were found in the frequencies of main alleles, A3 (0.32) and A9 (0.57). It is concluded that there is significant difference in the allele frequency for normal groups between Korean and Japanese populations.

F805

Factor V Leiden (G1691A) and G20210A Prothrombin Variant by Multiplex PCR-RFLP