Polymorphisms in promoter sequences of \textit{MDM2}, \textit{p53}, and \textit{p16^{INK4a}} genes in normal Japanese individuals

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Abstract

Research has been conducted to identify sequence polymorphisms of gene promoter regions in patients and control subjects, including normal individuals, and to determine the influence of these polymorphisms on transcriptional regulation in cells that express wild-type or mutant p53. In this study we isolated genomic DNA from whole blood of healthy Japanese individuals and sequenced the promoter regions of the \textit{MDM2}, \textit{p53}, and \textit{p16^{INK4a}} genes. We identified polymorphisms comprising 3 nucleotide substitutions at exon 1 and intron 1 regions of the \textit{MDM2} gene and 1 nucleotide insertion at a poly(C) nucleotide position in the \textit{p53} gene. The Japanese individuals also exhibited \textit{p16^{INK4a}} polymorphisms at several positions, including position -191. Reporter gene analysis by using luciferase revealed that the polymorphisms of \textit{MDM2}, \textit{p53}, and \textit{p16^{INK4a}} differentially altered luciferase activities in several cell lines, including the Colo320DM, U251, and T98G cell lines expressing mutant p53. Our results indicate that the promoter sequences of these genes differ among normal Japanese individuals and that polymorphisms can alter gene transcription activity.

Key words: Murine double minute 2, polymorphism, \textit{p16^{INK4a}}, \textit{p53}, transcription.

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Introduction

Different populations exhibit sequence polymorphisms of the murine double minute 2 (\textit{MDM2}) gene (Atwal \textit{et al.}, 2007), a target gene of the transcription factor p53. A promoter polymorphism of the \textit{MDM2} gene, SNP309, is located in the intron 1 region of the gene and influences transcriptional regulation in a cell line expressing wild-type p53 (Bond \textit{et al.}, 2004). Non-cancerous control subjects frequently have a polymorphism of p53 at codon 72 (Wu \textit{et al.}, 1995; Minaguchi \textit{et al.}, 1998), and this common polymorphism differentially alters promoter activity of the \textit{MDM2} gene that contains the SNP309 polymorphism (Yang \textit{et al.}, 2007). Cytosine-phospho-guanine (CpG) dinucleotides are methylated by DNA methyltransferase (DNMT) (Siedlecki and Zielenkiewicz, 2006), and the methylated form interacts with methyl-CpG-binding proteins (Fan and Hutnick, 2005), which serve as modulators of gene transcription. CpG methylation in the promoter region of p53 decreases promoter activity of the gene (Schroeder and Mass, 1997). Genomic DNA obtained from blood of people exposed to arsenic has been reported to exhibit methylation at the promoter regions of p53 and \textit{p16^{INK4a}}, a cycline-dependent kinase inhibitor, and the DNA methylation status of these promoters differs among people (Chanda \textit{et al.}, 2006). Normal populations have polymorphisms in \textit{DNMT}; a polymorphism \textit{DNMT3L} affects the ability of this gene to stimulate DNA methylation (El-Maarri \textit{et al.}, 2009). Nucleotide substitutions in the p53 promoter at 4 mutated positions, including position -250, and in the \textit{p16^{INK4a}} promoter at positions -735, -493, and -191 are found not only in Taiwanese patients with uterine leiomyoma (Hsieh \textit{et al.}, 2007) and frequently in melanoma families from other populations (Harland \textit{et al.}, 2000) but also in control groups; p53 promoter polymorphism at position -250 and \textit{p16^{INK4a}} promoter polymorphism at position -191 are located within CpG dinucleotides. Thus, control subjects, including normal populations, exhibit gene promoter polymorphisms. However, no studies have investigated differences at other nucleotide positions in \textit{MDM2}, p53, and \textit{p16^{INK4a}} promoter sequences among healthy individuals. In addition, it has not been determined whether known polymorphisms in the promoter

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sequence of these genes are present in other normal populations and alter gene promoter activity in other cell lines.

A number of polymorphisms have been identified in 10 ENCODE (Encyclopedia of DNA Elements) regions in the human genome of 48 individuals from 4 populations, including 8 Japanese individuals. Moreover, the frequency distributions of these polymorphisms have been investigated among different populations (International HapMap Consortium, 2005). In the present study, we sequenced the MDM2 and p53 promoters and the p16\(^{INK4a}\) promoter nucleotides at positions -735, -493, and -191 from genomic DNA extracted from whole blood samples obtained from healthy Japanese individuals and determined whether these promoter polymorphisms affect gene promoter activity in cell lines expressing mutant or wild-type p53. We found that normal Japanese individuals exhibit polymorphisms in these regions of the MDM2, p53, and p16\(^{INK4a}\) genes and that these polymorphisms alter promoter activity in some cell lines.

### Materials and Methods

#### Extraction of genomic DNA

Human peripheral blood was obtained from 17 healthy Japanese students, who consented to have their DNA sequenced for identification of polymorphisms. Genomic DNA was extracted from whole blood by using a QIAamp DNA Blood Mini kit (Qiagen, Hilden, Germany). DNA analysis of the samples showed that the frequency of commonly known polymorphisms (Kadowaki et al., 1995; Eguchi-Ishimae et al., 2005) was similar to that previously identified in control Japanese subjects.

#### Amplification of promoter regions by polymerase chain reaction

Promoter sequences of the MDM2 genes were amplified by polymerase chain reaction (PCR) in a reaction mixture containing genomic DNA (0.1 μg) and MDM2 primers (Table 1) in the presence or absence of 5% dimethyl sulfoxide with DNA polymerase (KOD-Plus DNA polymerase (Cline et al., 1997); Toyobo, Osaka, Japan) according to the manufacturer’s instructions. The amplification was performed in a thermal cycler (Takara PCR thermal cycler MP; Takara, Osaka, Japan) under the following conditions: denaturation at 94 °C for 15 s, annealing at (melting temperature [Tm]-5) °C for 30 s, and extension at 68 °C for 2 min. p53 and p16\(^{INK4a}\) promoter sequences were amplified using p53 and p16\(^{INK4a}\) primers (Table 1) and DNA polymerase (PfuTurbo DNA polymerase (Cline et al., 1997); Stratagene, La Jolla, CA) according to the manufacturer’s instructions under the following conditions: denaturation at 94 °C for 30 s, annealing at (Tm-5) °C for 1 min, and extension at 72 °C for 2 min.

#### Determination of nucleotide sequences

Sequence reactions were performed using an ABI PRISM Dye Terminator Cycle Sequencing kit (Perkin-Elmer Biosystems, Foster City, CA), and nucleotide sequences were determined using the ABI PRISM 377 automated DNA sequencer (Perkin-Elmer Biosystems).

### Table 1 - Primers used for PCR and sequencing

<table>
<thead>
<tr>
<th>Primer position</th>
<th>Oligonucleotide primer</th>
</tr>
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<tbody>
<tr>
<td>MDM2 gene</td>
<td></td>
</tr>
<tr>
<td>-725 to -704</td>
<td>5'-TCTGACCGGAGATCTGCTGCTTT-3'</td>
</tr>
<tr>
<td>-465 to -442</td>
<td>5'-TCTATCGGTTGTTCCACGGCTTCG3'</td>
</tr>
<tr>
<td>-310 to -289</td>
<td>5'-TCCGGACGGCCTTCGCGGCGG-3'</td>
</tr>
<tr>
<td>+96 to +175</td>
<td>5'-GACTGACAAGAAGTCTGCTG-3'</td>
</tr>
<tr>
<td>-725 to -702</td>
<td>5'-TATGGTACCTCGAGGACGCTCTGCTG3'</td>
</tr>
<tr>
<td>+99 to +176</td>
<td>5'-TAGTAGACTGCTAAAGTCACAGCAAGTG-3'</td>
</tr>
</tbody>
</table>

Nucleotide positions have been numbered by considering the positions of nucleotide C at the 5' end of exon 2 (Zauberman et al., 1995) in the MDM2 gene (accession number: U39736.1), nucleotide G at the initiation site for translation in the p16\(^{INK4a}\) gene (accession number: X54156.1) in the gene (Hara et al., 1996), and nucleotide G at the KpnI site of the PGV-B2 vector (sequence identical to that of pGL3-basic vector; GenBank accession number, U47295) as +1. The underlines indicate the recognition sites of the KpnI (single) and BglII (doublet) restriction enzymes.

#### Construction of reporter plasmid vectors

Regions from positions -725 to +99 of the MDM2 gene, positions -920 to +184 of the p53 gene, and positions -1703 to -93 of the p16\(^{INK4a}\) gene were amplified by PCR with primers bearing a cleavage site of the restriction enzyme KpnI or BglII. The amplified fragments were digested...
Polymorphisms in normal individuals

and separated by agarose gel electrophoresis. DNA fragments were extracted using a gel extraction kit (Concert Gel Extraction System; Invitrogen, Carlsbad, CA), precipitated with ethanol, and ligated between the \textit{KpnI} and \textit{BglII} sites of the PicaGene Basic Vector 2 (PGV-B2) reporter plasmid (Nippon Gene, Tokyo, Japan). The constructed vectors were cloned in \textit{Escherichia coli} DH5\alpha (Toyobo) and then purified using a QIAfilter Plasmid Midi kit (Qiagen). Nucleotide sequences were confirmed using primers (Table 1), including the PGV-B2 vector primers.

Analysis of nucleotide sequences and polymorphisms

The nucleotide sequences obtained for 17 individuals (or 9 individuals in the case of \textit{p16\textsubscript{INK4a}}) were compared with DNA sequences (accession numbers U39736.1, U28935.1, X54156.1, J04238.1, M13111.1, U94788.1, AF022809, and X94154.1) in the DNA database GenBank at the National Center for Biotechnology Information by using the program BLAST (basic local alignment search tool). For detection of polymorphisms, nucleotides were compared among individuals by using the sequence alignment editor program BioEdit.

Cell culture

Human colon adenocarcinoma Colo320DM (American Type Culture Collection [ATCC], Manassas, VA) and breast carcinoma YMB-1 (Japanese Collection of Research Bioresources [JCRB], Osaka, Japan) cell lines were grown in RPMI 1640 medium (Nissui Pharmaceutical, Tokyo, Japan). The human glioma U251 (Riken Cell Bank [RCB], Tsukuba, Japan) cell line was grown in Dulbecco’s modified Eagle’s medium (Nissui Pharmaceutical). Human glioblastoma T98G and cervix epithelioid carcinoma HeLa cell lines (RCB) were cultured in Eagle’s minimum essential medium (Nissui Pharmaceutical) containing nonessential amino acids and sodium pyruvate (Invitrogen). These cells were grown at 37 °C in media supplemented with 10% fetal bovine serum (Trace Scientific, Melbourne, Australia) in a humidified atmosphere of 95% air and 5% CO_2.

Transfection and reporter gene analysis

The cultured cells were plated at a density of 1.04 x 10^4 cells/cm^2 and then transfected with 1 μg of vector DNA using Lipofectamine 2000 (Invitrogen) according to the manufacturer’s instructions. The transfected cells were cultured and then lysed after 24 h in a reporter lysis buffer (Promega, Madison, WI). Luciferase was used as a reporter gene, and luciferase activity was determined using the Luciferase Assay System (Promega) in a luminometer (Monolight 2010; Analytical Luminescence Laboratory, San Diego, CA). For correction of variations in the amount of vector DNA incorporated into cells, the cultured cells were co-transfected with pCMVβ (Clontech, Palo Alto, CA), a β-galactosidase expression vector. Luciferase activity was normalized according to the β-galactosidase activity.

Statistical analysis

Comparisons among multiple groups were performed using analysis of variance (ANOVA) with a post hoc test. Differences between two groups were evaluated using unpaired Student’s t test. Statistical significance was defined as p < 0.01.

Results

Polymorphisms in promoter regions of the \textit{MDM2}, \textit{p53}, and \textit{p16\textsubscript{INK4a}} genes

We amplified the regions containing \textit{MDM2} and \textit{p53} promoters (Tuck and Crawford, 1989; Zauber et al., 1995) in genomic DNA isolated from 17 healthy individuals by PCR and determined the nucleotide sequences of the amplified regions. Comparisons of the nucleotide sequences among individuals revealed that nucleotide A was substituted with nucleotide G at position -628 in exon 1 (Figure 1A) and C with T at -466 (Figure 1B) and T with G at -215 (Figure 1C) in intron 1 of the \textit{MDM2} gene. Nucleotide C was inserted at positions -824 to -818 (C-to-C insertion) of the \textit{p53} gene (Figure 2). We identified 9 genotypes related to \textit{MDM2} and \textit{p53} polymorphisms containing nucleotide alterations (I-IX; Table 2); genotypes I, II, III, VI, and VIII were observed in 1 individual each, types V and IX were observed in 2 individuals each, and types IV and VII were observed in 4 individuals each. Promoter nucleotides of the \textit{p16\textsubscript{INK4a}} gene at positions -735, -493, and -191 were examined using samples obtained from 9 individuals randomly selected from the 17 Japanese individuals. We observed substitution of nucleotide A with nucleotide G in \textit{p16\textsubscript{INK4a}} at position -191 (Figure 3); no nucleotide substitutions were observed at positions -735 and -493. This nucleotide alteration was observed in 3 individuals who were homozygous A/A, 4 individuals who were heterozygous A/G, and 2 individuals who were homozygous G/G at -191.

The polymorphic \textit{MDM2}, \textit{p53}, and \textit{p16\textsubscript{INK4a}} sequences were cloned into PGV-B2 vectors, and 3 constructs of \textit{MDM2} PGV-ACT, PGV-ACG, and PGV-GTT (Table 3), 2 constructs of \textit{p53} PGV-7C and PGV-8C (Table 4), and 2 constructs of \textit{p16\textsubscript{INK4a}} PGV-A (containing nucleotide A at -191) and PGV-G (containing nucleotide G at -191) were obtained. Each of the cloned sequences was consistent with the nucleotide sequences determined by directly sequencing the PCR products (Figure 4A and other data not shown). In confirmation of the cloned sequences of \textit{p16\textsubscript{INK4a}}, we also observed 2 alleles of \textit{p16\textsubscript{INK4a}} containing a substitution of nucleotide C with T at position -1602 (Figure 4A), and substitutions of C with T at -871 (Figure 4B), T with A at -315 (Figure 4D), and A with G at -191 (Figure 4E) and deletion of nucleotide C at positions -862 to -858 (Figure 4C). These nucleotide alterations at positions
-1602, -871, -862 to -858, and -315 were ascertained by direct sequencing of PCR products (Figure S1). The MDM2 nucleotides at positions -339, -333, and -218 and positions -237 to -234, -231 to -228, and -223 to -221 (G-to-G insertion) and the p53 nucleotides at positions -512 to -511 (G-to-G deletion) and -479 to -477 (T-to-T insertion) dif-

**Figure 1** - Promoter polymorphisms of MDM2 at positions -628, -466, and -215. (A-C) Genomic DNA was amplified by PCR with a forward primer at positions -725 to -704 and a reverse primer at +96 to +75 of the MDM2 gene, and nucleotide sequences were determined by directly sequencing the PCR products. These sequencing reactions were performed using primers at positions -725 to -704 and -310 to -289. (*) shows nucleotides at positions -628 (A), -466 (B), and -215 (C) in the MDM2 promoters, where “N” indicates heterozygous nucleotides. Nucleotides around these positions (*) are indicated.

**Figure 2** - A poly(C) polymorphism of p53 at positions -824 to -818. (A-C) Genomic DNA was amplified by PCR with a forward primer at positions -918 to -899 and a reverse primer at -669 to -690 of the p53 gene. Nucleotide sequences were determined by directly sequencing the PCR products; these sequencing reactions were performed using a primer at positions -918 to -899. (*) shows homozygous (A and C) or heterozygous (B) nucleotides at positions -824 to -818 in the p53 promoters, where “NNCN” in (B) indicates sequences including “GCCG” and “CGCC,” which result from nucleotide C insertion. Nucleotides around the poly(C) polymorphic position (*) are indicated.

**Figure 3** - A polymorphism of p16INK4a at position -191. Genomic DNA was amplified by PCR with a forward primer at positions -485 to -465 and a reverse primer at +214 to +194 of the p16INK4a gene, and nucleotide sequences were determined by directly sequencing the PCR products. These sequencing reactions were performed using a primer at positions -248 to -228. (*) shows nucleotides at position -191 in the p16INK4a promoters, where “N” indicates heterozygous nucleotides. Nucleotides around the polymorphic position (*) are indicated.
fered from the nucleotides at the corresponding positions in the \textit{MDM2} and \textit{p53} sequences deposited in the GenBank database (accession numbers U39736.1, U28935.1, X54156.1, J04238.1, and M13111.1) (Tables 3 and 4).

However, these nucleotides were identical among all study individuals (Tables 3 and 4). The \textit{p53} nucleotides at positions -512 to -511 (G-to-G deletion) and -479 to -477 (T-to-T insertion) in the healthy individuals were consistent with the corresponding nucleotides in the \textit{p53} sequence in the GenBank database (accession number U94788.1, Table 4).

Influence of promoter polymorphisms on reporter gene activity in different cell lines

Several cell lines, including the colo320DM, U251, and T98G cell lines, were transfected with vector constructs containing \textit{MDM2}, \textit{p53}, and \textit{p16\textsuperscript{INK4a}} polymorphisms, and the activity of the luciferase reporter gene in the transfected cells was examined to assess whether these polymorphisms alter gene promoter activity. The level of luciferase activity increased in cell lines transfected with the \textit{MDM2} construct PGV-ACT (containing nucleotides A at position -628, C at -466, and T at -215) (Figure 5); slight luciferase activity was detected in cell lines transfected with the empty vector PGVB2 (PGV-(-)). A significant decrease or increase was found in the luciferase activity of Colo320DM or U251 cells transfected with the \textit{MDM2} construct PGV-GTT (containing an A-to-G substitution at -628 and a C-to-T substitution at -466) compared to the activity in cells transfected with PGV-ACT (Figure...
5A, B). Similar increase in luciferase activity was also observed in YMB-1 (Figure 5D) or HeLa (data not shown) cells transfected with the PGV-GTT construct. The MDM2 construct PGV-ACG (containing a T-to-G substitution at -215) did not significantly alter luciferase activity in the transfected Colo320DM and U251 cells (Figure 5A, B) and increased the activity in the transfected YMB-1 cells (Figure 5D). PGV-ACT, PGV-ACG, and PGV-GTT T98G transfectants had similar luciferase activities (Figure 5C). In the case of Colo320DM, U251, and T98G cells, the luciferase activities of cells transfected with p53 PGV-8C (containing nucleotide C insertion) and cells transfected with p16INK4a PGV-G were significantly lower and higher than the activities of cells transfected with p53 PGV-7C and cells transfected with p16INK4a PGV-A, respectively (Figures 6 and 7). The p53 PGV-8C and p16INK4a PGV-G vectors did not decrease and increase the luciferase activity in the transfected YMB-1 and HeLa cells, respectively (data not shown).

Discussion

In the present study, we found that normal Japanese individuals exhibit polymorphisms in the MDM2, p53, and p16INK4a promoter regions (Figures 1-4 and Figure S1) and that these individuals harbor several nucleotides in the MDM2 and p53 promoters that are different from those in the GenBank database (Tables 3 and 4); however, these nucleotides were identical among all the Japanese individuals included in this study. This result implies that the MDM2 and p53 promoter nucleotides differ among different populations, including individuals within the same population. The frequency of SNP309, a polymorphism involving nucleotide G, in intron 1 of the MDM2 gene is low among African Americans and high in other populations (Dharel et al., 2006; Hu et al., 2006; Millikan et al., 2006; Park et al., 2006; Atwal et al., 2007), including Ashkenazi Jewish, White, and Japanese populations. The polymorphism that we identified at position -215 in the intronic MDM2 promoter was polymorphism SNP309 of MDM2. The G allele
frequency of SNP309 at -215 in our Japanese samples was high (44.1%; 15 of 34 alleles) and similar to that in healthy individuals who visited a Japanese hospital (54%; 52 of 96 alleles) (Dharel et al., 2006). African Americans, Caucasians, and Ashkenazi Jewish populations have other MDM2 polymorphisms involving nucleotide substitutions from A

Figure 6 - Effects of the p53 poly(C) polymorphism on luciferase activity in different cell lines. (A-C) Colo320DM (A), U251 (B), and T98G (C) cells were transfected with the indicated vector constructs of PGV(-), PGV-7C, and PGV-8C. The transfected cells were cultured for 24 h, and luciferase activity was determined. Data are shown as percentages of values obtained from the PGV-7C transfectants and are expressed as the mean ± SEM (n = 8). *: Significant difference compared with the PGV-7C transfectant.

Figure 7 - Effects of the p16INK4a polymorphism at position -191 on luciferase activity in different cell lines. (A-C) Colo320DM (A), U251 (B), and T98G (C) cells were transfected with the indicated vector constructs of PGV(-), PGV-A, and PGV-G. The transfected cells were cultured for 24 h, and luciferase activity was determined. Data are shown as percentages of values obtained from the PGV-A transfectants and are expressed as the mean ± SEM (n = 8). *: Significant difference compared with the PGV-A transfectant.
to G (rs937283) and C to T (rs2870820), which are registered in the dbSNP database, at rates of 27%, 34%, and 25% and of 6%, 34%, and 0%, respectively (Atwal et al., 2007). These polymorphisms are located at position -628 in exon 1 (rs937283) and position -466 in intron 1 (rs2870820) of the MDM2 gene and corresponded to the nucleotides observed in our samples. The frequency of the G or T allele in healthy Japanese individuals was 29.4% each (10 of 34 alleles). The T allele frequency in Japanese individuals (29.4%) was higher than that in African American (6%) and Ashkenazi Jewish (0%) populations. Families with Li-Fraumeni (LF; an autosomal dominant cancer-predisposition syndrome) and of 6%, 34%, and 0%, respectively (Atwal et al., 2007). These polymorphisms were not observed in our healthy Japanese samples, and the poly(C) polymorphism of p53 at positions -824 to -818 (C-to-C insertion) found in our samples was observed at a high frequency of 41.2% (14 of 34 alleles). It has been reported that the nucleotides at positions -735, -493, and -191 of p16INK4a are substituted not only in melanoma families in the United Kingdom, the United States, Italy, and Australia but also frequently in control populations (Harland et al., 2000). In these control populations, the frequencies of homozygous A/A, heterozygous A/G, and homozygous G/G polymorphisms at position -191 of p16INK4a are 38%, 48%, and 14%, respectively (Harland et al., 2000). The corresponding frequencies of these polymorphisms in our samples were 33.3%, 44.4%, and 22.2%. In addition, the frequency of polymorphisms C/C, C/C, 5C/5C, and T/T at positions -1602, -871, -862 to -858, and -315 of p16INK4a was 44.4%; that of C/C, C/T, 4C/4C, and A/A was 22.2%; that of C/T, C/C, 5C/5C, and T/T was 22.2%; and that of C/T, C/C, 5C/5C, and T/T was 11%. Thus, the promoter polymorphisms in MDM2 at positions -215, -628, and -466, in p53 at positions -342, -250, -216, -103, and -33, and in p16INK4a at positions -735, -493, and -191 were differentially distributed in some normal and control populations. In this study, polymorphisms were observed among healthy Japanese individuals in the sequence of MDM2 at positions -628 and -466, of p53 at positions -824 to -818, and of p10INK4a at positions -1602, -871, -862 to -858, -315, and -191.

The promoter region of MDM2 includes multiple transcription-factor response elements (Phelps et al., 2003; Bond et al., 2004), including the p53-responsive element (positions -93 to -74 and -55 to -36) (Zauberman et al., 1995). Reporter gene activity is increased in wild-type p53-expressing HeLa cells transfected with the G allele of MDM2 SNP309 (Bond et al., 2004); the polymorphism SNP309 is located adjacent to a binding site of the transcription factor SP1 in the MDM2 promoter and can alter SP1 binding at this site. The MDM2 G allele of SNP309 observed at -215 similarly increased reporter activity in HeLa cells (data not shown) and YMB-1 cells (Figure 5D). In contrast, the reporter luciferase activity of Colo320DM, U251, and T98G cells transfected with SNP309 at -215 was not altered (Figure 5A, B, C). We also observed a decrease, increase, or no change in luciferase activity in transfected Colo320DM, T98G, U251, HeLa, or YMB-1 cells that harbored the MDM2 polymorphism at both positions -628 and -466 (Figure 5). Deletion of a promoter region of MDM2 at positions -725 to -311, which include the -628 and -466 polymorphic positions, did not alter luciferase activity in Colo320DM cells and increased luciferase activity in other cells (Figure S2). This deletion experiment indicated that luciferase activity in Colo320DM cells may not be regulated constitutively by transcription factors through the MDM2 promoter sequence containing the polymorphic sites -628 and -466 and that luciferase activity in other cells appears to be negatively regulated by these factors via the promoter containing the polymorphic sites. The reporter gene activity induced by polymorphism is thought to be caused by cell type-dependent regulation of the promoter machinery. A search for transcription factors that can bind to sequences around the polymorphic sites was carried out in the TRANSFAC database by using the MatInspector program (Wingender et al., 2000). Database analysis showed that the -466 MDM2 polymorphism affected the matrix similarity of the transcription factor MZF1. The MDM2 polymorphism at both positions -628 and -466 may alter promoter activity via a known regulatory factor at position -466.

Patients with diseases such as cancer show not only nucleotide deletions and mutations (Pollock et al., 2001; Attwooll et al., 2002; Tanai et al., 2002; Fombonne et al., 2005; Hsieh et al., 2007) but also nucleotide methylation (Herman et al., 1996; Narimatsu et al., 2004; Amatya et al., 2005) in the promoter regions of p53 and p16INK4a. The deletion at positions -346 to -338 (C/EBP site) of p53 in LF (Attwooll et al., 2002) and the mutation at position -250 of p53 in patients with uterine leiomyoma (Hsieh et al., 2007) are located within CpG dinucleotides. The locus of the p53 poly(C) polymorphism at positions -824 to -818 was adjacent to CpG sequences, which are located at positions -826 to -825 and -818 to -817, and the -191 nucleotide G of p16INK4a was located within CpG sites. Methylation of the p16INK4a promoter has been found in noncancerous tissues in 3 of 21 (14%) individuals with non-cirrhotic virus-positive livers (Narimatsu et al., 2004). Exposure to arsenic differentially stimulates the methylation of p53 and p16INK4a promoters (Chanda et al., 2006). We did not observe nucleotide alterations in these methylated nucleotides.
in the p53 and p16INK4a promoters among normal Japanese individuals.

Some transcription factors interact with p53 and p16INK4a promoters and regulate gene promoter activity through these promoter regions (Furlong et al., 1996; Kirch et al., 1999; Myöhänen and Baylin, 2001; Xue et al., 2004; Wu et al., 2007). The polymorphic sites identified in these promoters in the present study were not located within their transcription factor-binding sites. Promoter methylation of p53 is induced in the T98G cell line, and expression of the p53 gene is upregulated by treatment of the cell line with 5-aza-2’-deoxycytidine, a methylation inhibitor, (Amatya et al., 2005). Luciferase reporter gene activity was altered in T98G cells harboring the p53 poly(C) polymorphism (Figure 6C). Chromatin immunoprecipitation with an antibody against the methyl-CpG binding domain protein (MBD) 2, an MBD family protein (Hendrich and Bird, 1998), revealed that the MBD2 complexes extracted from human colon cell lines bind to the p16INK4a promoter region (from position -494 to -101) and that this binding is abolished by treatment with the methylation inhibitor 5-aza-2’-deoxycytidine (Magdinier and Wolffe, 2001). It was also revealed that although HeLa cells express MBD2 (Billard et al., 2002), MBD2 does not bind to the p16INK4a promoter region (Magdinier and Wolffe, 2001). Database analysis with the TFSEARCH program showed that several transcription factors, including those of SP1, CREB, and NF-kB, can bind to the p53 poly(C) and p16INK4a -191 sites and that nucleotide alterations at these sites affect the similarity of sequences of the substrates for the transcription factors activator protein 2 and ATF or NF-kB, etc., respectively. The regions around and within the p53 poly(C) sequence are rich in guanine and cytidine nucleotides and are similar in part to the consensus sequences for transcription factors such as activating enhancer-binding protein 2 (Hilger-Eversheim et al., 2000) whose gene includes the p53-responsive sequence (Li et al., 2006). p53 can form complexes with SP1, CREB-binding protein, and NF-kB protein (Borellini and Glazer, 1993; Huang et al., 2007). The p53 poly(C) and p16INK4a -191 polymorphisms did not decrease and increase luciferase activity in the YMB-1 and HeLa cells expressing wild-type p53, respectively. In contrast, these polymorphisms altered luciferase activity in the Colo320DM, U251, and T98G cells (Figures 6 and 7) that harbored p53 mutants at positions codon 248, 273, and 237, respectively. In each case, the polymorphism-induced reporter gene activity seems to be attributable to cellular differences in promoter regulation that occurs through the binding of DNA-interacting proteins; the regulation may result in a decrease or increase in reporter activity and may depend on cellular methylation of CpG nucleotides and on the ability of transcription factors.

Patients with cancers exhibit differential MDM2 and p53 protein expression in gliomas and breast cancers, which contain the G and/or T alleles of SNP309 (Tsuiki et al., 2007; Krekac et al., 2008), and differential p16INK4a protein expression in gastric cancers in the absence of methylation (Tsuje et al., 2000). Gene expression analysis performed using the TCGA database revealed that MDM2 and p16INK4a gene expression levels differ among patients with cancers, including those with glioblastoma multiforme; it is not known whether the regulatory regions of these genes have polymorphisms. Chemotherapeutic drugs increase the transcript level of p53 target genes in cells expressing wild-type MDM2, and the increased levels decrease in cells expressing the SNP309 G allele (Arva et al., 2005). In addition, MDM2 promoter activity is increased through the p53-binding site in the MDM2 gene in cells expressing the thyroid hormone receptor (Qi et al., 1999) and through multiple transcription-factor response elements in estrogen receptor (ER) α-positive breast cancer cells (Phelps et al., 2003), and p16INK4a reporter activity is increased in senescent cells in comparison with that in young cells (Wang et al., 2001). Breast cancer YMB-1 and cervix epithelioid carcinoma HeLa cell lines are ERα-positive and -negative cells, respectively. The region around the MDM2 polymorphic site at position -466 was a possible binding site of the transcription factor GATA (Phelps et al., 2003) whose gene expression is increased in ERα-positive cells treated with estradiol (Eeckhoute et al., 2007). The normal Japanese individuals who participated in our study included 13 male and 4 female students. Further studies are needed to clarify the significance of promoter polymorphisms in the variations in gene expression among individuals, in responses to chemical and hormone stimuli in individuals of different genders and ages, and under normal and aberrant conditions.

Research has been conducted on the construction of common patterns of sequence variations in the human genome (International HapMap Consortium, 2005). Polymorphism SNP309 is present in linkage disequilibrium with other MDM2 intronic polymorphisms, and the linkage differs between different populations: Ashkenazi Jewish and Caucasian populations or African Americans (Atwal et al., 2007). Exon 1 of the p14ARF gene (Robertson and Jones, 1998) is located approximately 20 kb upstream of the first p16INK4a exon 1 and splices into the common exons 2 and 3 shared with p16INK4a in a different reading frame (Mao et al., 1995). A Korean population has a promoter polymorphism of p14ARF at position -1477, which is observed in strong linkage disequilibrium in a haplotype block, and the -1477 A allele of p14ARF is associated with the methylation status of the p14ARF promoter and alters the level of p14ARF mRNA in colorectal tumor tissues (Kang et al., 2008). Japanese populations frequently exhibited p14ARF polymorphism at position -1477 (Figure S3). p16INK4a polymorphisms at positions -871, -862 to -858, and -315 or MDM2 polymorphisms at positions -628 and -466 appeared to be possible candidates for linkage disequilibrium. Single-nucleotide polymorphisms (SNPs) (Brookes, 1999) are the
most frequently occurring sequence variation in a population and are associated with a predisposition to disease and susceptibility or tolerance to medication. They are useful as markers of individual constitution and as information for personalized medicine. A population-based study showed that the SNP309 polymorphism in MDM2 and/or the common polymorphism at codon 72 in p53 are associated with predispositions to carcinomas in populations who visited Asian hospitals (Kawaguchi et al., 2000; Hong et al., 2005; Dharel et al., 2006). Other studies showed that the MDM2 SNP309 polymorphism or the p53 common polymorphism at codon 72 is not associated with or only weakly associated with predisposition to carcinomas in other populations (Millikan et al., 2006; Krekac et al., 2008; Tornesello et al., 2009; Zubor et al., 2009). The p16\textsuperscript{INK4a} -191 polymorphism does not segregate with disease in melanoma kindreds in the United Kingdom (Harland et al., 2000). It is of interest to determine whether the promoter polymorphisms of MDM2, p53, and p16\textsuperscript{INK4a} identified in this study are present in linkage disequilibrium in a haplotype block and influence predisposition to diseases; our findings provide information for studies on the construction of common polymorphism patterns and for studies on the frequency distribution among different populations, including diseased patients. In the current study, we identified promoter polymorphisms in healthy Japanese individuals and found that gene promoter activity was altered by these polymorphisms in different cell lines. Our results indicate that promoter sequences of the MDM2, p53, and p16\textsuperscript{INK4a} genes differ among normal Japanese individuals.

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Supplementary Material
The following online material is available for this article:
Figure S1 - Promoter polymorphisms of p16INK4a at positions
-1602, -871, -862 to -858, and -315.
Figure S2 - Effects of the deletion of a region in the MDM2 pro-
moter on luciferase activity in Colo320DM, U251, T98G,
YMB-1, and HeLa cells.
Figure S3 - A promoter polymorphism of p14ARF at position -1477
in normal Japanese individuals.
This material is available as part of the online article from
http://www.scielo.br/gmb.

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Figure S1 - Promoter polymorphisms of p16INK4a at positions -1602, -871, -862 to -858, and -315. (A-D) Genomic DNA was amplified by PCR with forward primers at positions -2028 to -2009, -1198 to -1174, and -485 to -465 and reverse primers at -1173 to -1192, -475 to -498, and +214 to +194 of the p16INK4a gene; nucleotide sequences were determined by directly sequencing the PCR products. These sequencing reactions were performed using primers at positions -1706 to -1684, -950 to -931, and -381 to -361. (*) shows nucleotides at positions -1602 (A), -871 (B), -862 to -858 (C), and -315 (D) in the p16INK4a promoters, where “N” indicates heterozygous nucleotides. “NNNC” in (C) indicates sequences including “CGTC” and “GTCC,” which result from nucleotide C deletion. Nucleotides around these positions (*) are indicated.
Figure S2 - Effects of the deletion of a region in the MDM2 promoter on luciferase activity in Colo320DM, U251, T98G, YMB-1, and HeLa cells. Colo320DM (A), U251 (B), T98G (C), YMB-1 (D), and HeLa (E) cells were transfected with the indicated vector constructs of PGV-ACT (containing nucleotide A at position -628, C at -466, and T at -215), PGV-ACT(Del) (nucleotides at -725 to -311 deleted), and PGV(-). The transfected cells were cultured for 24 h, and luciferase activity was determined. Data are shown as percentages of values obtained from the PGV-ACT transfectants and are expressed as the mean ± SD (n = 3). *: Significant difference compared with the PGV-ACT transfectant.
Figure S3 - A promoter polymorphism of p14ARF at position -1477 in normal Japanese individuals. Genomic DNA was amplified by PCR with a forward primer at positions -1649 to -1627 and a reverse primer at -1090 to -1112 of the p14ARF gene; the nucleotide sequences were determined by directly sequencing the PCR products. These sequencing reactions were performed using a primer at position -1649 to -1627. (*) shows nucleotides at position -1477 in the p14ARF promoters, where “N” indicates heterozygous nucleotides containing G and A. Nucleotides around the polymorphic position (*) are indicated. The -1477 polymorphic frequency in 9 healthy Japanese individuals was 33.3% for homozygous G/G, 44.4% for heterozygous G/A, and 22.2% for homozygous A/A. Nucleotide positions are numbered with respect to nucleotide G (Robertson and Jones, 1998) at the initiation site of transcription in the p14ARF gene (accession number: AF082338).