A wild-derived inbred mouse strain, MSM/Ms, provides insights into novel skin tumor susceptibility genes

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Abstract: Cancer is one of the most catastrophic human genetic diseases. Experimental animal cancer models are essential for gaining insights into the complex interactions of different cells and genes in tumor initiation, promotion, and progression. Mouse models have been extensively used to analyze the genetic basis of cancer susceptibility. They have led to the identification of multiple loci that confer, either alone or in specific combinations, an increased susceptibility to cancer, some of which have direct translatability to human cancer. Additionally, wild-derived inbred mouse strains are an advantageous reservoir of novel genetic polymorphisms of cancer susceptibility genes, because of the evolutionary divergence between wild and classical inbred strains. Here, we review mapped Stmm (skin tumor modifier of MSM) loci using a Japanese wild-derived inbred mouse strain, MSM/Ms, and describe recent advances in our knowledge of the genes responsible for Stmm loci in the 7,12-dimethylbenz(a)anthracene (DMBA)/12-O-tetradecanoylphorbol-13-acetate (TPA) two-stage skin carcinogenesis model.

Key words: 7,12-dimethylbenz(a)anthracene (DMBA)/12-O-tetradecanoylphorbol-13-acetate (TPA) two-stage skin carcinogenesis, MSM/Ms, skin tumor susceptibility gene, Stmm loci, wild-derived inbred mouse strain

Introduction

Identifying specific genetic variants responsible for increased susceptibility to familial or sporadic cancers has significant implications for predicting individual cancer risk and developing strategies for prevention or targeted therapy [1, 2]. Mouse models of cancer have been extensively used to analyze the genetic basis of cancer susceptibility and have led to the identification of multiple loci that confer, either alone or in specific combinations, an increased susceptibility to cancer [3–8].

One of the most commonly used experimental inflammatory cancer models is the DMBA/TPA two-stage skin carcinogenesis model. The pathology of this model is almost identical to that of human skin cancer, and it offers an ideal model to research skin cancer initiation and growth [9, 10]. The first step is to treat mice with a low dose of the mutagenic substance 7,12-dimethylbenz(a)anthracene (DMBA) to initiate carcinogenesis. In the second step, mice are continuously treated with 12-O-tetradecanoylphorbol-13-acetate (TPA) to stimulate epidermal tumor growth. During tumor promotion, papillomas are thought to be caused by an inflammatory response owing to additional TPA treatment. After long-term treatment (approximately 20 weeks), some benign tumors (papillomas) progressed to squamous cell carcinoma (SCC; Fig. 1). The role of various genes and cellular signaling pathways involved in developing skin tumors can be investigated in this two-stage skin carcinogenesis model by using a genetically engineered mouse model [10]. Genetically modified mice have been used to analyze various genes (i.e., oncogenic transcription factors, constitutive centromere-related network proteins) [11–15].
Inbred strains recently obtained from wild-derived mice are often more resistant to carcinogens and some pathogens than commonly used inbred strains. Compared with classical inbred strains, wild-derived mice show genetic polymorphisms every 100–200 base pairs in the genome [16]. Investigating the molecular basis of these traits provides insights into how polymorphisms contribute to cancer susceptibility. Most wild-derived strains reproduce with classic inbred mice, representing a rich source of evolutionarily significant diversity for forward genetics studies [16, 17]. Although still mostly unexplored, these mice are the new models for identifying and studying novel cancer susceptibility genes. In addition, these models can be combined with the latest innovations in next-generation sequencing and genome editing technologies to study single nucleotide polymorphism (SNP) levels in cancer susceptibility genes.

Here, we review mapped Stmm (skin tumor modifier of MSM) loci using a Japanese wild-derived inbred mouse strain, MSM/Ms, and describe recent advances in our knowledge of the genes responsible for Stmm loci in the DMBA/TPA two-stage skin carcinogenesis model.
Mapping of Stmm Loci Using a Forward Genetics Approach

The tumor susceptibility of the DMBA/TPA two-stage skin carcinogenesis model varies among mouse strains, and a genetic approach has been employed to identify genes related to tumor susceptibility [18, 19]. Using the genetic approach, several skin tumor-susceptibility loci were identified using commonly inbred strains or wild-derived strains [20–30], some of which have clear translatability to human cancer susceptibility [31–33]. Therefore, the DMBA/TPA skin carcinogenesis model is a useful model for identifying modifier genes.

The MSM/Ms mouse strain was generated in Mishima, Shizuoka, Japan, from wild-caught mice of the Mus musculus molossinus subspecies [34]. Breeding for the MSM/Ms strain began in 1978, after six mice were donated to the Cytogenetics Department of the National Institute of Genetics. After repeated sister-brother mating for the past 40 years, inbreeding has reached generation N100, and these mice are now recognized as an inbred mouse strain like other commonly inbred mouse strains. Due to its genetic divergence from standard inbred mouse strains mainly established from M. m. domesticus, the MSM/Ms strain has been broadly used in linkage studies, mainly in Japan. The MSM/Ms strain is also useful as a genetic resource, with complete genome sequence information (https://shigen.nig.ac.jp/mmdbj/top.jsp) [35, 36], a full set of mouse consomic strains (MSM/Ms as the chromosome donor and C57BL/6J as the host strain) [37], a BAC clone library (https://dna.brc.riken.jp/en/cloneeset/mms_mms_bac_en) [38], a microsatellite database (https://shigen.nig.ac.jp/mouse/mmdbj/top.jsp) [39], and ES cells [40, 41] now available. In addition, MSM/Ms mice have a very low incidence of development of several types of tumors [42–45], age-dependent hearing loss [46–50], and unique behavioral traits such as a very high locomotor activity [51–56], compared with classical inbred mice.

We first investigated DMBA/TPA skin carcinogenesis using MSM/Ms and FVB/N mice as susceptible strains. In these experiments, MSM/Ms and (FVB/N × MSM/ Ms) F1 mice showed strong resistance to skin carcinogenesis, and their phenotype was dominant [57]. Therefore, in order to identify the tumor susceptibility modifier gene, we used the following protocol. 1) Using p53 tumor suppressor gene knockout mice, we created F1 backcross mouse groups of p53 wild-type (p53+/–) and heterozygous knockout mice (p53+/-). We also investigated the dependence of the loci on the p53 gene. 2) We made linkage analysis possible at each stage of carcinogenesis by measuring the diameters of all developing tumors over time. Combining these improvements, 228 F1 backcross mice (121 with the p53+/– background and 107 with the p53+/- background) were subjected to the DMBA/TPA chemical carcinogenesis protocol (Fig. 2). The number of papillomas at 20 weeks and the presence of carcinoma at 40 weeks were recorded in each mouse. Additionally, papillomas were further categorized into three groups based on size (<2 mm, 2–6 mm, and >6 mm in diameter; Fig. 2). All mice were genotyped using 107 SNP markers distributed evenly throughout the genome.

In a population of 228 F1 backcross mice, highly significant QTLs for papilloma development with a maximum LOD score of 7.0 were mapped to chromosomes 7 and 4. A suggestive QTL for carcinoma development was mapped to chromosome 5. We conducted linkage analysis for each of the three categories of tumors, which were specified based on tumor diameter. As a result of linkage analysis, Stmm1 and Stmm2 were mapped around 50 cM of chromosome 7 as resistance loci for early and mid-stage papillomas (diameter <2 mm, 2–6 mm). We succeeded in mapping Stmm3 as resistance to late-stage papillomas (>6 mm) locus around 40 cM on chromosome 4. Notably, Stmm3 was not detected by linkage analysis of the F1 group of p53+/-, suggesting that the gene responsible for Stmm3 is strongly dependent on the p53 gene. We succeeded in mapping Stmm1-Stmm12 on mouse chromosomes, including the less effective loci (Table 1). Previously identified cancer susceptibility loci, including other types of cancer, are also present in these candidate regions (Table 1). Further studies will be necessary to narrow down the candidate regions, by combining this information with human GWAS data and congenic mapping.

Refinement of Candidate Regions for Stmm Loci by Congenic Mapping

On the other hand, for Stmm1a/b and Stmm3, multiple congenic mouse strains were prepared, and mapping was performed to narrow down the candidate regions [58, 59]. We screened stage-specific papilloma modifier loci based on size (Table 1). Stmm1 and Stmm2 were identified on chromosome 7 as modifier loci conferring resistance to smaller-sized papillomas. Nagase et al. mapped the loci of Skts1 and Skts2 loci, which had previously been identified using a wild-derived inbred mouse strain, Mus spretus [25, 26]. Congenic analysis narrowed down the candidate genetic region of Skts1 to approximately 15 Mb, which is proximal to Igf1r [70].

Stmm2, identified near Skts2 and Hras on chromosome 7, has the same effect on the allelic specificity of the Hras mutation as Skts2, suggesting that the same gene
may be responsible for Skts2 and Stmm2 [58]. Skin carcinogenesis experiments using congeneric mice with Skts2 have not yet been reported; data presented in our report suggest that Stmm2 and Skts2 genes may have a relatively mild effect on papilloma multiplicity compared with Stmm1 [58].

As previously shown, the vicinity of the Stmm3 candidate region contains several tumor modifier loci, including Skts7 and Skts-fp1, mapped for chemically induced skin papilloma susceptibility by analysis of an NIH and Mus spretus cross and an FVB/N and PWK cross, respectively [23, 25] (Table 1). PWK, Mus spretus, and MSM/Ms are wild-derived inbred strains that share common haplotypes. A gene common to Stmm3 might be involved in the susceptibility of these two strains through mating.

Eventually, Stmm1 on chromosome 7 was subdivided into Stmm1a (about 0.24 Mb) [unpublished data] and Stmm1b (about 4.7 Mb) [59]. In the case of Stmm3 on chromosome 4, the candidate region was narrowed down

Fig. 2. Investigation of the Stmm loci by the forward genetics approach. (A) To identify genetic loci that control susceptibility to skin carcinogenesis, we subjected 228 FVB/N × (FVB/N × MSM/Ms) F1 backcross mice, 121 on a p53+/+ background and 107 on a p53+/− background, to the 7,12-dimethylbenz(a)anthracene (DMBA)/12-O-tetradecanoylphorbol-13-acetate (TPA) chemical carcinogenesis protocol. We monitored their tumor development for 40 weeks. For each mouse, we documented the number of papillomas at 20 weeks after initiation as well as the presence of carcinomas at 40 weeks. In addition, papillomas were further categorized into three groups based on size (<2 mm, 2–6 mm and >6 mm in diameter). (B) A schematic drawing of suppressive functions of Stmm loci in the development of a chemically induced skin tumor. Stmm1a is the most effective and is involved in resistance to early-stage papilloma genesis. Stmm1b affects papilloma formation from early to late stages, and Stmm3 is involved in p53 gene-dependent resistance to the malignant transformation from late-stage papillomas to squamous cell carcinomas (SCCs).
Table 1. Genetic candidate regions of Stmm loci and other cancer susceptibility loci in the vicinity

| Locus name | Chromosome No. | Genetic marker | rs No. | Candidate region (bp) | Tumor category | Effect of p53 | Other cancer susceptibility QTLs in the vicinity | References |
|------------|----------------|----------------|--------|-----------------------|---------------|-------------|-----------------------------------------------|------------|
| Stmm1a     | 7              | D7Mit351–D7Mit419 | –      | 96,776,254–99,188,960 | Early papilloma | Independent | -                             | [57], Okumura et al. (Unpublished data) |
| Stmm1b     | 7              | D7Mit53–D7Mit23 | –      | 110,583,818–115,356,689 | Early-middle papilloma | Independent | -                             | [57–59]    |
| Stmm2      | 7              | D7Mit255–D7Mit259 | –      | 124,705,587–144,566,894 | Early-middle papilloma | Independent | Skin2                | [25, 26, 57, 58] |
| Stmm3      | 4              | D4Mit26–D4Mit116 | –      | 88,535,666–93,366,888 | Late papilloma, SCC | Dependent | Skin7, Skts-fp1 | [22, 25, 57, 60, 61] |
| Stmm4      | 6              | D6SNP511–D6SNP51512 | rs30149679–rs38226370 | 55,547,013–108,360,689 | Middle papilloma | Independent | Skts11          | [25, 57]    |
| Stmm5      | 5              | DSSNP12–DSSNP6 | rs13478361–rs3023057 | 83,810,839–126,154,436 | Middle-late papilloma, SCC | Dependent | HeX5, Thy12 | [57, 62, 63] |
| Stmm6      | 3              | D3SNP519* | rs46831546 | 45,778,132–10,000,000 | Middle papilloma | Unknown | -                             | [57]        |
| Stmm7      | 3              | D3SNP504* | rs3053537 | 111,903,098–20,000,000 | Middle papilloma | Unknown | Raml2             | [57, 64]    |
| Stmm8      | 12             | D12SNP12* | rs3682985 | 70,353,327–20,000,000 | Middle papilloma | Unknown | Par3              | [57, 65]    |
| Stmm9      | 13             | D13SNP12* | rs3684241 | 71,839,279–20,000,000 | Middle papilloma | Unknown | Par10             | [57, 66]    |
| Stmm10     | 17             | D17SNP19–D17SNP510 | rs3705342–N.D. | 68,392,183–94,100,159 | Middle papilloma | Unknown | Skin10            | [25, 57]    |
| Stmm11     | 11             | D11SNP8* | rs3023216 | 117,028,591–15,000,000 | Middle-late papilloma, SCC | Dependent | Skin10             | [57, 67, 68] |
| Stmm12     | 1              | D1SNP11–D1SNP12 | rs3688428–rs36897954 | 4,137,358–32,741,433 | Early papilloma | Unknown | Locc1              | [57, 69]    |

Early papilloma, tumor diameter of <2 mm; middle papilloma, tumor diameter of 2–6 mm; late papilloma, tumor diameter of >6 mm; SCC, squamous cell carcinoma; N.D., not detected. *Linkage peak. #Candidate regions are based on MGI data.

...to approximately 4.8 Mb by congenic mapping [61, 62]. As a result of an in silico search for a gene that is functionally dependent on the p53 gene of Stmm3, Cdkn2a was extracted. The two tumor suppressor genes p16^{ink4a} and p16^{ink6-a} are encoded by Cdkn2a through the sharing of exons, and both proteins have amino acid substitutions in one polymorphism between FVB/N and MSM/Ms. Based on these results, the most promising candidate gene for Stmm3 was considered to be Cdkn2a.

On the other hand, there are 15 protein-encoding genes in Stmm1a, and their details are still under investigation. There were 28 protein-encoding genes in Stmm1b. Pth (parathyroid hormone) was extracted as a gene with an amino acid substitution between FVB/N and MSM/Ms. Therefore, we focused on Pth as one of the candidate genes for Stmm1b.

**Stmm1b: Parathyroid Hormone is a Novel Regulator of Early-stage Skin Carcinogenesis**

PTH is a hormone that, along with vitamin D, is essential for calcium homeostasis in the body [71, 72]. It is well known that the skin acts as a neuroendocrine organ. Almost all the elements controlling the hypothalamic-pituitary-adrenal axis activity are expressed in the skin, as previously described in detail [73]. PTH and PTH-related peptide (PTHrP) also influence the proliferation and differentiation of epidermal cells [74–77] via paracrine and intracrine routes [78], but the role of PTH in skin carcinogenesis is poorly understood. Surprisingly, a comparison of serum intact PTH (iPTH) between the two mouse strains showed that MSM/Ms had significantly higher iPTH levels than FVB/N (5–10 times higher), while there was no difference in calcium and vitamin D levels [59]. Furthermore, the amount of iPTH in serum was approximately twice as high in Stmm1b sub-congenic mice compared with control mice [59].

According to a report by Kalu and Hardin, the basal circulating PTH level in sera is higher in rodents than in humans. Besides, there are differences between rodent strains, with the rat F344 strain having the highest PTH levels and the Wistar and Sprague Dawley rats having low PTH levels [79]. Therefore, differences in serum PTH levels, such as those seen between MSM/Ms and FVB/N, are also observed between and within species.

Next, a highly PTH-expressing Pth^{Msm-Tg} was prepared by introducing an MSM/Ms-BAC clone containing a Pth genetic region into FVB/N mice, and we then carried out a skin carcinogenesis experiment. The results showed that Pth^{Msm-Tg} was resistant to skin carcinogenesis. In addition, based on the results of carcinogenesis experiments using Pth heterozygous knockout mice (Pth^{+/–}), it was clarified that mice with low iPTH levels were susceptible to skin carcinogenesis, and it was possible to show for the first time that PTH is a skin cancer modifier (Fig. 3) [59].

We then detected a well-conserved Val/Met polymorphism (rs51104087) in the Pro-PTH region (amino acid position 28) of the mouse Pth gene, between FVB/N (i.e., valine) and MSM/Ms (i.e., methionine). The Pro-PTH sequence is conserved in mammals and is essential for the signal peptide sequence in vitro [80] and in vivo [81]. Based on a MoG+ database search, this SNP in Pth exon 1 was found only in wild-derived strains and was well conserved in *musculus* subspecies (Table 2).
We hypothesized that the cSNP in Pro-PTH could influence protein posttranslational modifications and increase secreted PTH levels. Nascent PTH is translated as a pre-pro protein. Thus, analysis of the effect of this SNP in in vitro experiments confirmed that the intracellular stability of PTH was improved, and the amount of extracellular secretion was increased [59]. It was also shown that secreted PTH acts to suppress skin cancer by inducing the suppression of epidermal cell proliferation, an increase in intracellular calcium, and accompanying cell pro-differentiation, in a quantity-dependent manner (Fig. 3) [59].

These results showed that Pth accounts, at least in part, for the effects attributed to Stmm1b. Our data demonstrating the association between high serum PTH levels and skin tumor suppression in a two-stage skin

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**Table 2.** Variant table for **Pth** and **p19Arf** in wild-derived strain mice

| wild-derived strains | Subspecies       | Pth_rs51104087 | PTH氨酸 acid 28 | Cdkn2a_rs218801173 | p19Arf氨酸 acid 149 |
|----------------------|------------------|----------------|------------------|-------------------|-------------------|
| CAST/EiJ             | castaneus        | G              | Val              | G                 | Val               |
| HM/Ms                | domesticus       | G              | Val              | C                 | Leu               |
| WSB/EiJ              | domesticus       | G              | Val              | C                 | Leu               |
| BFM/2/Ms             | domesticus       | G              | Val              | C                 | Leu               |
| PGN2/Ms              | molossinus       | G              | Val              | G                 | Val               |
| MOLF/EiJ             | molossinus       | A              | Met              | G                 | Val               |
| JF1/Ms               | molossinus       | A              | Met              | G                 | Val               |
| MSM/Ms               | molossinus       | A              | Met              | G                 | Val               |
| CHD/Ms               | muscular         | G              | Val              | G                 | Val               |
| PWK/PhJ              | muscular         | A              | Met              | G                 | Val               |
| KJR/Ms               | muscular         | A              | Met              | G                 | Val               |
| SWN/Ms               | muscular         | A              | Met              | G                 | Val               |
| NJL/Ms               | muscular         | A              | Met              | G                 | Val               |
| BLG2/Ms              | muscular         | A              | Met              | G                 | Val               |

Sequence data from MoG+.
carcinogenesis model, underscore the potential therapeutic effect of using PTH or PTH analogs in skin cancer prevention and treatment.

In our previous report, we demonstrated a genome-wide significant linkage at Stmm3 on chromosome 4. However, this linkage peak at Stmm3 completely disappeared in p53+/− mice. We concluded that corresponding genes for Stmm3 are genetically dependent on p53 [57, 61]. Next, we generated p53+/− and p53+/+ sub-congenic mouse lines for Stmm3. These mice were exposed to two-stages of skin carcinogenesis using DMBA/TPA [62]. The skin carcinogenesis experiments revealed that both p53+/−Stmm3FVB/FVB and p53+/−Stmm3FVB/MSM mice developed late-stage (larger than 6 mm in diameter) papillomas (Fig. 4A). In contrast, when we measured the size of papillomas in p53+/+ sub-congenic mice, only Stmm3FVB/FVB mice developed late-stage papillomas

![Image](https://example.com/figure4.png)

**Fig. 4.** p19ArfMSM more efficiently induces the transcriptional activity of p53 compared with p19ArFVB. (A) Representative dorsal back skin photographs of p53+/+ or p53+/−Stmm3FVB/FVB (Stmm3F/F) and Stmm3FVB/MSM (Stmm3F/M) congenic mice at 20 weeks after 7,12-dimethylbenz(a)anthracene (DMBA)/12-O-tetradecanoylphorbol-13-acetate (TPA) initiation. (B) This figure shows the molecular mechanism behind the suppressive effect of p19ArfMSM in late-stage papillomas. We noted that p19ArfMSM was preferentially localized in the nucleus after TPA treatment. In contrast, p19ArFVB was preferentially localized in the cytoplasm. p19Arf mediates p53 transactivation in the nucleus by stabilizing p53 protein through inactivation of Mdm2 protein. In addition, the nuclear localization of p19Arf is necessary for this transactivation function. Eventually, p19ArfMSM led to an increase in p53 stabilization and activated the p53 pathway. Therefore, we conclude that one of the responsible genes for Stmm3 is p19Arf.
Therefore, the Stmm3 region exerted more potent suppressive effects on papillomas in the presence of two copies of p53, indicating that the effects of Stmm3 are dependent on the p53 gene [62].

Based on congenic mapping results, we focused on p53-related Cdkn2a as a candidate factor for Stmm3 [61]. Two tumor suppressor genes, p19Arf and p16Ink4a, are encoded by Cdkn2a through the sharing of exons. Therefore, each gene knockout mouse was prepared by genome editing with CRISPR/Cas9 using ES cells of MSM/Ms [62]. Furthermore, by mating these mice with FVB/N, MSM/Ms allele knockout mice with an F1 genetic background were produced, and two-stage skin carcinogenesis experiments were performed. The results revealed p19Arf to be the most promising candidate gene for Stmm3, as only MSM/Ms allele KO mice with p19Arf showed significant increases in the number of late-stage benign tumors, the incidence of malignant tumors, and prolongation of mouse survival [62].

No significant difference was observed between the p19Arf RNA expression levels and the p19Arf protein levels between cutaneous tissues of the two mouse strains. On the other hand, the responsible SNP (rs218801173) between MSM/Ms (valine) and FVB/N (leucine) was localized in the C-terminal domain of p19Arf. In addition, this SNP was not found in most inbred and wild-derived mice belonging to M. m. domesticus and was widely conserved in wild-derived mice not belonging to M. m. domesticus subspecies, including MSM/Ms (Table 2).

In cell localization experiments, deletion mutants lacking the C-terminal domain were wholly localized in the cytoplasm without entering the nucleus. Arf has been reported to be degraded, at least in part, by the proteasome in the cytoplasm [82, 83]. Thus, the C-terminal domain of p19Arf may localize proteins in the nucleus to avoid degradation. However, human p14ARF lacks this C-terminal region [84, 85]. Therefore, the biological importance of the C-terminal region may be overlooked. When the SNP was introduced into 3T3 cells by retroviral transfection, it was found that stimulation of TPA resulted in a change in p19Arf localization and that the MSM/MS allele would likely be localized in the nucleoplasm without entering the nucleus. Arf has been reported to be degraded, at least in part, by the proteasome in the cytoplasm without entering the nucleus. Arf has been reported to be degraded, at least in part, by the proteasome in the cytoplasm [82, 83]. Thus, the C-terminal domain of p19Arf may localize proteins in the nucleus to avoid degradation. However, human p14ARF lacks this C-terminal region [84, 85]. Therefore, the biological importance of the C-terminal region may be overlooked. When the SNP was introduced into 3T3 cells by retroviral transfection, it was found that stimulation of TPA resulted in a change in p19Arf localization and that the MSM/MS allele would likely be localized in the nucleoplasm without entering the nucleus [82, 83]. Thus, the C-terminal domain of p19Arf may localize proteins in the nucleus to avoid degradation. However, human p14ARF lacks this C-terminal region [84, 85]. Therefore, the biological importance of the C-terminal region may be overlooked.

When the SNP was introduced into 3T3 cells by retroviral transfection, it was found that stimulation of TPA resulted in a change in p19Arf localization and that the MSM/MS allele would likely be localized in the nucleoplasm without entering the nucleus (Fig. 4B). p19Arf stabilizes p53 protein by inactivating Mdm2 protein and mediating p53 transactivation [86]. In addition, the nuclear localization of p19Arf is necessary for this transactivation function [87, 88] (Fig. 4B). Therefore, we quantified the amount of p53 protein and the expression level of downstream factors (such as p21, Bax, and Noxa) in mouse skin. The results showed that mice carrying the MSM/Ms allele had higher p53 protein levels and elevated levels of downstream factor mRNA expression (Fig. 4B) [62].

We also revealed two SNPs of CDKN2A associated with breast cancer incidence in the Japanese cancer population. An SNP (rs36228836) in linkage disequilibrium with one of the two SNPs, was predicted by in silico analysis to affect transcription factor binding. This result suggests that rs36228836 influences breast cancer risk via transcriptional regulation. In recent years, SNPs near the CDKN2A/B locus on chromosome 9q21.3 have been linked to the risk of various human cancers and metabolic disorders [89].

We demonstrated that genetic polymorphisms in Cdkn2a and CDKN2A affect cancer risk. Therefore, we conclude that one of the genes responsible for Stmm3 is p19Arf. We have shown that linkage analysis using a mouse model can be translated to humans for identification of cancer susceptibility alleles and thus development of targeted therapies.

**Concluding Remarks and Future Perspective**

This review focused on the cancer susceptibility/resistance of a Japanese wild-derived inbred mouse strain, MSM/Ms, and attempted to identify the genetic modifiers. Forward genetics approaches have revealed that multiple genetic factors act on the two-stage process of skin carcinogenesis. In particular, it was shown that Stmm1a/b is involved in the development of early and mid-stage papillomas and that Stmm3 functions in a p53-dependent manner in late-stage papillomas to SCCs.

To the best of our knowledge, this is the first to report that PTH, one of the genes responsible for Stmm1b, is involved in skin carcinogenesis. Additionally, the cSNP responsible for increased iPTH in sera may be derived from the M. m. musculus subspecies (Table 2). In the future, it will be essential to measure iPTH levels in some wild-derived mouse strains to determine their associations with a variety of diseases, including cancer. It is also necessary to clarify why MSM/Ms mice can maintain normal calcium and vitamin D levels despite high serum iPTH. Moreover, we are currently investigating associations with the incidences of several cancers as well as data for SNPs around PTH locus data in human cancer patients. In conjunction with this, we are verifying findings with mouse models other than skin carcinogenesis models using Pth knockout mice.

On the other hand, the SNP in the rodent-specific C-terminal region of p19Arf, the gene we identified as being responsible for Stmm3, is a domain that has not been investigated in detail because it does not exist in humans. However, our research suggests that this domain is essential for the nuclear localization of p19Arf, which may
shed light on the evolutionary properties and unknown functions of p19ARF in other mammals. Further research is needed in the future. We are currently analyzing the gene responsible for Stmm1a, which has the most potent resistance effect, and we believe that identification of it will provide valuable insights into the cancer resistance in MSM/Ms.

In future studies, it will be essential to analyze human cancer susceptibility genes and polymorphisms by utilizing this information and integrating it with information from human GWAS. In addition, wild-derived inbred mice exhibit phenotypes different from those of classical inbred mice for various genetic diseases [16, 90–92]. We are currently screening resistant candidate SNPs from haplotype analysis of wild-derived strain of mice based on Stmm candidate region information using a bioinformatics approach. We plan to introduce these candidate SNPs directly into susceptible inbred mice by CRISPR/Cas9 to perform carcinogenesis experiments and single-cell RNA sequencing. In the future, further technological innovations in areas such as sequencing and genome editing will clarify the cause of genetic diseases at the individual level, and it is thought that wild-derived inbred mouse strains will become an increasingly important and valuable genetic resource.

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