Molecular biology of malignant mesothelioma

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Abstract  Human malignancies develop via a multi-step process that involves the accumulation of several key gene alterations with associated genetic and epigenetic events. Although malignant mesothelioma (MM) has been demonstrated to be clearly correlated with asbestos exposure, it remains poorly understood how asbestos fibers confer key gene alterations and induce cellular transformation in normal mesothelial cells, which results in the acquisition of malignant phenotypes, including deregulated cell proliferation and invasion. Malignant mesothelioma presents with the frequent inactivation of tumor suppressor genes of \( p16^{INK4a}/p14^{ARF} \) on chromosome 9p21 and neurofibromatosis type 2 (\( NF2 \)) on chromosome 22q12, with the latter being responsible for the NF2 familial cancer syndrome. In contrast, MM shows infrequent mutation of the \( p53 \) gene, which is one of the most frequently mutated tumor suppressor genes in human malignancies. Genetic abnormalities of oncogenes have also been studied in MM, but no frequent mutations have been identified, including the epidermal growth factor receptor (\( EGFR \)) and \( K-RAS \) genes. Recent studies have suggested the activation of other receptor tyrosine kinases, including Met, and the deregulations of mitogen-activated protein kinase (MAPK) and phosphatidylinositol-3-kinase (PI3K)-AKT signaling cascades, although the alterations responsible for their activation are still not clear. Thus, further genome-wide studies of genetic and epigenetic alterations as well as detailed analyses of deregulated signaling cascades in MM are necessary to determine the molecular mechanisms of MM, which would also provide some clues for establishing a new molecular target therapy for MM.

Keywords  Asbestos  ·  Malignant mesothelioma  ·  Molecular target therapy  ·  Oncogene  ·  Tumor suppressor gene

Introduction

Malignant mesothelioma (MM), a highly lethal neoplasm arising primarily in the pleural, peritoneal, or pericardial cavity, is thought to develop from surface mesothelial cells [1]. In up to 80% of patients, MM occurs within approximately 30–40 years following exposure to asbestos [2]. Since patients with MM are usually diagnosed at advanced stages and MM is refractory to conventional therapy, the prognosis of patients with MM is very poor. The median survival of patients with malignant pleural mesothelioma (MPM) is 7–11 months after diagnosis, despite the recent advancements in chemotherapeutical modalities combining cisplatin and antifolates such as pemetrexed or raltitrexed [3, 4].

The regulation of asbestos occurred relatively late in Japan, with the result that the number of patients with MM is expected to increase year by year, with the peak incidence predicted around 2030–2040 [5]. To date, in Japan approximately 1000 deaths can be attributed to MM. It has been estimated that 250,000 people will die of MPM in Europe during the next three decades and in the United States, 2500–3000 new cases are diagnosed each year [6, 7]. These numbers are attributable to the wide use of
asbestos in various industrial and building materials, resulting in exposure to asbestos not only among workers in factories and/or at construction and demolition sites but also by ordinary citizens including family members of the workers and local residents near the factories and affected sites.

About 80% of MM develops in the pleura, 20% in the peritoneum, and less than 1% in the pericardium. The pathologically, epithelial type accounts for about 60% of all cases, the sarcomatous type for about 20%, and the biphasic type with both components ranges around 20%. Since MM is a relatively rare human malignancy, both basic and clinical studies have lagged far behind those on other common types of malignancies, such as lung, colon, and breast cancers. However, many researchers are now focusing their efforts on identifying the mechanisms and key genes in MM development with the aim of using this information to develop new diagnostic tools and target molecules of new therapy.

**Asbestos-induced oncogenesis**

Up to 80% of mesothelioma patients have been exposed to asbestos, thus establishing a clear link between asbestos exposure and MM development [1]. Many animal models have also demonstrated the carcinogenicity of asbestos. There are six types of asbestos: amphiboles-amosite (brown asbestos), crocidolite (blue asbestos), anthophyllite, actinolite, and tremolite, the serpentine chrysotile (white asbestos). However, it has not yet been clearly determined whether asbestos fibers act directly on the mesothelial cells or whether they indirectly cause mesothelioma. Several plausible explanations have been put forward on how asbestos fibers can cause MPM [1, 8]. One such suggestion is that asbestos fibers mechanically induce pleural irritation: long and thin asbestos fibers can be inhaled deeply into the lungs, penetrating and repeatedly scratching the mesothelial surface, resulting in prolonged cycles of damage, repair, and local inflammation. Alternatively, asbestos fibers can also mechanically interfere with the mitotic process of the cell cycle by disrupting the mitotic spindle, which may result in chromosomal abnormalities and aneuploidy. A third proposal is that highly reactive oxygen species (ROS) and reactive nitrogen species (RNS) are induced by asbestos, leading to DNA damage and strand breaks. A ramification of the interaction of long fibers with cells is frustrated phagocytosis and a prolonged oxidative burst. Finally, asbestos can induce cytokines and growth factors, such as transforming growth factor-β (TGF-β) and platelet-derived growth factor (PDGF), as well as transcription factors, such as nuclear factor kappa B (NF-κB) and activator protein-1 (AP-1). Additionally, in rat mesothelial cells, crocidolite asbestos was found to cause autophosphorylation of epidermal growth factor receptor (EGFR) [9].

**Inactivation of tumor suppressor genes in MPM**

The most frequently inactivated tumor suppressor genes (TSGs) in human malignancies are p53 and p16^{INK4a}/p14^{ARF}. Most MPMs show frequent inactivation of p16^{INK4a}/p14^{ARF}, although only 20–25% of these show a mutation of p53. Analysis of primary samples of MPM revealed that over 70% of the samples showed downregulation of p16^{INK4a}/p14^{ARF} [10, 11]. Using established MPM cell lines, Taniguchi et al. [12] found that almost all of the cell lines had a homozygous deletion of the p16^{INK4a}/p14^{ARF} gene locus. The p16^{INK4a} gene product controls the cell cycle via the cyclin-dependent kinase 4 (CDK4)/Cyclin-D-RB pathway, while the p14^{ARF} gene product regulates p53 through inactivation of the human homolog of mouse double minute 2 (HDM2), which is an upstream regulator of p53. Thus, the homozygous deletion of p16^{INK4a}/p14^{ARF} indicates the inactivation of two major tumor-suppressing pathways of RB and p53 in the cell.

The loss of chromosome 22q12 is frequently detected in MPM. The neurofibromatosis type 2 (NF2) gene, which had been initially identified as a target gene of NF2 familial cancer syndrome, has also been shown to be the target gene of the 22q12 loss in MPM [13, 14]. Neurofibromatosis type 2 patients frequently develop vestibular schwannoma, meningioma, and other neuronal malignancies, while sporadic tumors of these types of cancer also harbor a NF2 mutation [15]. However, to date, there has been no published report of NF2 patients showing a higher susceptibility to MPM. The NF2 gene is inactivated by homozygous deletion, nonsense mutation, or missense mutation in MPMs. Bianchi et al. [14] and Sekido et al. [13] reported that about 40% of MPMs showed the genetic mutation of NF2; among the 60% of MPMs without this NF2 genetic mutation, about 20% showed a down-regulation of NF2. In total, 50–60% of MPMs showed inactivation of NF2. In an animal model, an established line of Nf2 (±) knockout mice were shown to develop MPMs in the earlier stage and, more frequently, after asbestos exposure [16]. Remarkably, similar to human MM, tumors from Nf2 (±) mice showed frequent homozygous deletions of the p16^{INK4a}/p14^{ARF} locus and adjacent p15^{INK4b} tumor suppressor gene.

The NF2 gene encodes a membrane–cytoskeleton-associated protein, Merlin, an adaptor protein with a FERM (four-point-one, ezrin, radixin, and moesin) domain, which transduces a growth-regulatory signal [17]. Merlin has been shown to interact with several proteins, including...
CD44, ezrin radixin moesin (ERM) proteins, p21-activated kinase 1 (PAK1), and loses its tumor-suppressing activity with phosphorylation at serine 518. This serine 518 site is phosphorylated with RAC/PAK1 and dephosphorylated with myosin phosphatase targeting subunit 1-protein phosphatase 1 (MYPT-1-PP1δ) [18]. The activity of Merlin is thought to be regulated by cell-adhesion (adherence junction), cell-extracellular matrix adhesion, or extracellular growth signals. The downstream signaling of Merlin is mediated by Hippo cascade, which was initially identified via genetic studies in Drosophila [19]. Thus, Merlin is thought to be one of the key molecules in the signaling cascades that determine the properties of invasion, cell growth, and survival of malignant mesothelioma cells (Fig. 1).

**Activation of oncogenes in MPM**

Simian virus 40 (SV40) is a double-stranded DNA polyomavirus of monkey origin, which has been suggested to be associated with MM development [20]. Since the polio vaccine that was used between 1954 and 1963 was widely contaminated with SV40, vaccination was proposed as a plausible vector for the widespread SV40 infection of the human population. However, even people who had not received a polio vaccination were found to be infected with SV40, leaving unanswered the questions as to how SV40 virus infected humans.

Since SV40 expresses large T and small T antigens, and the large T antigen binds and inactivates the p53 and RB tumor suppressors, SV40 infection has been recognized as one of primary mechanisms of mesothelioma development pathogenetically. Intrapleural infection of SV40 in hamsters induces mesothelioma development by 6 months after administration, and SV40 induces Met, Notch-1, and telomerase activity, which also supports the functional roles of SV40.

However, most of the studies showing a strong relationship between SV40 and MM consisted exclusively of PCR-based assays in which the simple amplification of specific segments was considered to indicate a positive result; otherwise, the data were conflicting and reproducibility was limited. The results from several more recent studies suggest that such results were false-positives due to contamination by plasmids that were in general use in many of the laboratories common among the studies [21]. Consequently, the involvement of SV40 in malignant mesothelioma remains controversial.

The activation of receptor tyrosine kinase (RTK) family members has been investigated in MM. Among these, the *Met* oncogene has been shown to be frequently expressed
in primary tumors and cell lines of MM [22]. Hepatocyte growth factor (HGF) is a ligand of Met, and HGF-MET plays a role in the mitogen-activated protein kinase (MAPK) and phosphatidylinositol-3-kinase (PI3K)-AKT cascades. The PI3K-AKT cascade has also been shown to be activated in MM. Furthermore, MMs with positive AKT phosphorylation status show phosphorylated (activated) mTOR (mammalian target of rapamycin), which is one of the downstream molecules of AKT. Thus, the activation of the PI3K-AKT-mTOR signaling cascade in MM is thought to be induced by activation of the upstream HGF-MET (Fig. 1). Additionally, although EGFR, another RTK which is frequently mutated in Asian female adenocarcinoma of the lung, was shown to be overexpressed in 56% of primary tumors, no EGFR mutation was documented [23].

Malignant mesothelioma cell lines produce many other growth factors and cytokines [1]. Platelet-derived growth factor, TGF-β, insulin-like growth factor (IGF) have been studied in depth. In addition, factors involving angiogenic pathways have also been demonstrated to be expressed in MM cells, including interleukin (IL)-6, IL-8, fibroblast growth factors (FGFs), and vascular endothelial growth factors (VEGFs) [24].

**Searching for new key genes in malignant mesothelioma**

Since only a small number of oncogenes or TSGs with genetic alterations has been identified in MM, other, as yet unidentified genes may well be responsible for its development. Traditional allelotyping and karyotype analyses have revealed nonrandom chromosomal abnormalities, including 1p, 3p, 4p15.1-p15.3, 4q25-q26, 4q33-q34, 6q, 9p, 14q11.1-q12, 14q23-q24, and 22q [25, 26]. A comparative genomic hybridization (CGH) technique has recently been introduced to search for additional genes that are potentially involved in MM biology. New alteration regions have been identified, including 1q, 4q, 5p, 6p, 7p, 8p, 8q, 10p13-pter, 13q, 14q, 15q, 17p12-pter, 17q, and 20, in which new cancer-associated genes of MM may be harbored [27, 28]. A recent study of array-based CGH analysis with MMs from a total of 22 individuals identified high-copy gain at 1p32, which includes the JUN protooncogene [12]. JUN is a transcription factor and functions as homo- or hetero-dimerization with FOS to form the transcription factor AP-1, which can bind to the promoter region of intermediate genes involved in cell division and other cell functions. Both crocidolite and chrysotile asbestos reportedly caused increases in the expression of JUN and FOS in rat pleural mesothelial cells [29]. Since the gene amplification of JUN was identified in a subset of MPM tumors, it was suggested that there were some strong and persistent factors for JUN activation during the development of MPM tumor cells. A more recent study reported an activated mutation of N-RAS in three of 38 MMs [30].

Finally, expression profiling using microarray has been also studied to identify specific gene expression changes in MM compared with normal mesothelium [31–34]. Several new candidate oncogenes and TSGs of MM were proposed, and patient prognosis was shown to be predictable with the differences of gene expression profiling.

**Application for molecular target therapy**

Malignant mesothelioma is a highly aggressive tumor, and the patient prognosis with advanced-stage MM is very poor. Combination chemotherapy with cisplatin and antifolate has recently been shown to be superior to cisplatin alone [3, 4]. Although several molecular target therapies have been tested, no satisfactory results have been obtained to date. For example, a phase II study of an EGFR inhibitor, gefitinib, was conducted for 43 patients with previously untreated MM [35]. Although 97% of patients with MM had EGFR overexpression, gefitinib was not active in MM and EGFR expression did not correlate with failure-free survival. Imatinib, another tyrosine kinase inhibitor known to affect both Kit and PDGFα (and β) receptors, has also been shown to have limited efficacy for MPM [36].

A recent microarray analysis on 99 MPM detected advanced-stage, sarcomatous histology and the p16INK4a/p14ARF homozygous deletion to be significant adverse prognostic factors [37]. The same study also found that more aggressive MPM expressed higher levels of Aurora kinases A and B, which are serine/threonine kinases with multiple roles in mitotic progression. Thus, the role of Aurora kinases is of interest due to the recent development of their small-molecule inhibitors. In addition, a small-molecule inhibitor of TGF/β type I receptor has been shown to inhibit murine mesothelioma tumor growth in vivo [38].

**Summary**

The long latency period between asbestos exposure and tumor development implies that multiple—and likely diverse—genetic changes are required for the malignant transformation of mesothelial cells. Many studies have been conducted to determine the underlying key genetic and epigenetic events responsible for the development of MPM, some of which may be directly caused by asbestos fibers. New animal models of MM and human MM cell lines are also being established to present more useful tools for detailed analyses of the carcinogenesis of MM and the
development of new therapeutic modalities [16, 39]. Thus, more in-depth knowledge of key gene alteration, specific expression profiling, and other fundamental abnormalities at the cellular, intercellular, and tissue levels in MM cells will be of great help in developing future strategies for potential molecular targets as well as other therapeutic modalities, such as immunotherapy.

Acknowledgments

This work was supported by a Special Coordination Fund for Promoting Science and Technology from the Ministry of Education, Culture, Sports, Science and Technology (H18-1-3-1). I thank Dr. Hideki Murakami, Dr. Yutaka Kondo, Dr. Hirotaka Osada, and Dr. Tetsuo Taniguchi for their helpful comments. I regret the lack of citations for many important observations in the text, but their omission is made necessary by restrictions on the preparation of review manuscripts.

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