Abstract. Lung cancer is a major cause of cancer-related mortality in both men and women. A 5-year survival of lung cancer patients is only 15% with a negative correlation between progressively advanced lung cancer stage and a 5-year survival period. The only chance for cure is surgical resection if done at the early stage of the disease. Therefore, an early diagnosis and a better prediction of prognosis could decrease mortality. An early diagnosis could provide the opportunity for a therapeutic intervention early in the course of the disease. Genetic alterations in the cancer genome include aneuploidy, deletions and amplifications of chromosomal regions, loss of heterozygosity (LOH), microsatellite alterations, point mutations and aberrant promoter methylation. Of the various types of genetic alterations (i.e. gene amplifications, allele deletions, point mutations or deletions and methylation) reported in different tumor types, aberrant promoter methylation of genes is recent and is the focus of the present review. Specifically, we will briefly review the role of promoter methylation in various malignancies and then focus on lung cancer diagnosis and promoter gene methylation with emphasis on the methylation status of genes of the innate host defense, namely the surfactant proteins A and D.

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1. Lung cancer prevalence and current status

Lung cancer is a major cause of cancer-related mortality in both men and women in industrialized countries and causes more deaths than colorectal, breast and prostate cancer combined (1,2). The overall annual incidence and mortality rate of lung cancer has been estimated to be ~104 million new cases per year and 921,000 deaths in the world, with the highest rates currently observed in Europe and North America (3,4). The incidence and mortality rates are higher for men than for women. However, lung cancer mortality has increased markedly among women since 1960, following an increased prevalence of smoking (5). A high degree of correlation between smoking and lung cancer has been observed. Smoking accounts for 80% of the attributed risk among men and for 45% of the cases among women (6).

The 5-year survival of lung cancer patients is only 15%, which is much lower than the survival rate of colorectal (41%) and breast (67%) cancers. This is largely due to the fact that three quarters of lung cancer patients are diagnosed when their disease has spread regionally or distantly (7). There are studies on negative correlation between a 5-year survival period and lung cancer stage (as per TNM classification, where T is characteristic of primary tumor, N is regional lymph node involvement and M is metastasis). A 5-year survival period has been reported as high as 60-70% following resection of stage I lung cancer, while a 5-year survival rate of lung cancer as low as 8-13% has been observed if treatment starts at stages III-IV (8). At present, the only chance of cure is surgical resection at the early stage of the disease, with better prognosis for small tumors compared to larger ones. Thus, an early diagnosis of lung cancer and prediction of patient’s prognosis on the basis of clinical and genetic...
characteristics of tumor could potentially decrease lung cancer mortality by enabling earlier and more appropriate therapeutic intervention.

2. DNA methylation

DNA methylation mainly refers to methylation at cytosine residues located in dinucleotide CpG sites. The specifics relating to CpG dinucleotide location and methylation status have been reviewed previously (9-15). In brief, the CpG dinucleotide distribution in most of the genome is statistically lower than the expected frequency. However, in some genomic regions that mainly coincide with promoters or regions involved in gene regulation, the frequency of CpG dinucleotides is that of the expected value. These C+G rich regions are termed CpG islands [reviewed in (16)]. In the human genome, the pattern of CpG methylation varies during development [reviewed in (11)]. For example, in the case of germ line-specific genes and certain tissue-specific genes, promoter regions are subject to methylation as part of normal developmental processes [reviewed in (11,17)]. This ensures that specific genes are expressed in specific tissues and at specific times of development. Methylation of regulatory regions is involved in genomic imprinting and X chromosome inactivation in females [reviewed in (18)]. In a healthy individual [reviewed in (11)] most CpG islands are unmethylated and associated genes are expressed. However, if CpG islands are located within genes expressed in a tissue-specific manner, it is possible that the CpG islands remain unmethylated while these genes are silent (11). The lone CpGs found throughout the rest of DNA (i.e. not in CpG islands) are mostly methylated (19).

DNA methylation acting via deregulated gene expression has been recognized as a key component of aging (20,21) and various diseases (22). Although, initially reported in cancer, DNA methylation is now being appreciated as playing a role in neurological, cardiovascular and immunological pathologies as well. For example, there are studies of associations between aberrant promoter methylation of certain genes and hypertension (23), immunodeficiency syndromes (i.e. ICF, ATRX) and autoimmunity diseases (i.e. systemic lupus erythematosus, rheumatoid arthritis) (24,25) and neurological disorders (i.e. Alzheimer’s disease) (22,26).

3. DNA methylation in malignancies other than lung cancer

The cancer genome is characterized by a general decline in methylated cytosine level (genomic hypomethylation) accompanied by a local CpG island hypermethylation (27,28), with both processes leading to dysregulated gene expression. Genomic hypomethylation (29) and perhaps gene-specific CpG island hypomethylation (30-32) may lead to cancer via activation of proto-oncogenes, reactivation of transposable elements, loss of gene imprinting and hyperactivation of other genes (i.e. host defense genes, see below). On the other hand, when CpG islands are hypermethylated, the activity of regulatory proteins that promote transcription is restricted due to tightly packed nucleosomes (33). This often leads to suppression of gene expression of genes that affect tumor suppression, DNA repair and chromatin condensation. Tumor suppressor genes contain unmethylated CpG islands in their promoters that become methylated in various malignancies. Although, both of these processes (i.e. genomic hypomethylation and gene-specific promoter hypermethylation or hypomethylation) occur concurrently in cancer, thus far no relationship has been found between global genomic hypomethylation and local hypermethylation, indicating that these processes may be independent of each other (34).

**Hypermethylation.** DNA hypermethylation is a well established epigenetic alteration seen in various types of malignancies. A growing number of genes are being recognized that harbor dense methylation in normally unmethylated promoter CpG islands, with some of them common across different types of tumors. These include genes mainly involved in functions such as cell-cycle control and apoptosis (p14, p15, p16, Rb, DAPK), DNA repair (MGMT, hMLH1), adhesion and metastasis (CDH1, CDH13), biotransformation (GSTP1) and signal transduction (RARß and APC). For example, epigenetic silencing of glutathione S transferase (GSTP1) is the most common (>90%) genetic alteration reported in prostate cancer. It can be used to accurately distinguish benign and malignant prostate lesions (35). A similar prevalence of methylation changes was shown for RARß gene. The methylation of RARß correlates with more advanced pathological stage and allows discrimination between neoplastic and non-neoplastic prostate tissue (this included high-grade prostatic intraepithelial neoplasias and benign prostate hyperplasia) (36). The combination of methylation analysis with histological tests resulted in detection of 97% of prostate adenocarcinoma cases in comparison with only 64% when a histological test alone was used (37). Furthermore, an analysis of the hypermethylation status of a panel of four genes, GSTP1, RARB, TIG1 and APC, could detect prostate carcinoma with 100% specificity. Similarly, hypermethylation of 3 or more genes from a panel of 5 genes (CDH13, HSPA2, MLH1, RASSF1A, SOCS2) was observed in endometrial cancer, while 91% of the controls showed hypermethylation of none or fewer than 3 genes (38). Hypermethylation of a gene panel consisting of RARB, p16, p14, RASSF1A, DAPK, and GSTP1 was observed in breast cancer tissue compared to non-cancer breast tissue (39).

**Hypomethylation.** Genomic hypomethylation refers to loss of methylation at CpG sites located in intergenic and intronic DNA regions (global hypomethylation), as well as in promoters of specific genes (gene-specific hypomethylation). Global hypomethylation mainly affects repeat sequences and transposable elements. It often leads to cancer via chromosomal instability due to aberrant transcription and recombination, while localized hypomethylation often leads to increased expression of proto-oncogenes, growth factors, and genes, the protein products of which are involved in cancer cell proliferation, invasion and metastasis (40). There is substantial evidence of association of hypomethylation (whether at genome level or individual loci) with cancer. For example, satellite DNAs (juxta centromeric Sat2 and centromeric Sato) were found hypomethylated in ovarian samples with a direct correlation between stage of tumors and...
degree of hypomethylation (41,42). Similarly repeat sequences Sat2, Sato and SatR-1 were found to be hypomethylated in breast cancer (43,44). Hypomethylation at L1, a younger subfamily of autonomous long interspersed nuclear elements, which are a type of non-LTR retrotransposons, was reported in hepatocellular carcinoma (45), tumors of bladder, head and neck (46) and various benign and malignant ovarian carcinomas (47). Evaluation of the methylation status of LINE-1 repetitive sequences in genomic DNA derived from microdissected samples from several human normal and neoplastic tissues revealed a greater percentage of hypomethylation at LINE-1 repetitive sequences in several cancer tissues including breast, colon, lung, head and neck, bladder, esophagus, liver, prostate and stomach than their normal tissue counterparts (46). Similarly, hypomethylation of tumor genes, such as urokinase type plasminogen activator (upA), heparanase and S100A4 led to their increased expression in cancer cells. uPA (48,49) and heparanase (50) belong to a category of proteases produced by tumor cells that degrade extracellular matrix, while S100A4 is a calcium binding protein that regulates production of matrix-degrading enzymes. These enzymes are responsible for remodeling of the extracellular matrix and for an increase in tumor cell proliferation and motility (51). In summary, the available data indicate that a) a derangement of CpG methylation in the promoter of a variety of genes has been shown to associate with numerous cancers, including lung cancer and b) methylation markers of various genes have been shown to be useful in tissue-specific cancer diagnosis.

4. Lung cancer and DNA methylation

Lung cancer is a disease where both, environment and genetics play a role (6,52-56). There are two main types of lung carcinomas: non-small cell lung carcinoma (NSCLC) and small cell lung carcinoma (SCLC) (57). SCLC is most often a centrally located tumor arising from bronchial epithelium. NSCLC constitute the majority of all lung cancers (~85-90%) and are sub-classified into three main histological types: the squamous cell, which is linked to smoking and for the most part is found near the bronchus, the adenocarcinoma found in the outer lungs and the large cell carcinoma that can originate anywhere in the lung. Squamous cell cancer has at least one of three differentiating features that include individual cell keratinization, spherical keratinization, or intercellular cytoplasmic bridges. Adenocarcinomas are malignant tumors with tubular, acinar, or papillary growth patterns, and/or mucus production. Large cell carcinoma is characterized by large nuclei, prominent nucleoli, abundant cytoplasm and well defined cell borders.

Most lung cancers start in the lining of the bronchi. Carcinogenesis, starting from a normal cell to an invasive carcinoma, is a multistep process (58). Lung carcinoma is preceded by a premalignant lesion, such as hyperplasia, metaplasia, or dysplasia of the bronchial epithelium. The transition of a premalignant lesion to lung cancer is often accompanied by several genetic events (some of them are common between SCLC and NSCLC), such as alterations in the expression of proto-oncogenes and tumor suppressor genes (58) and perhaps of other genes (32). These changes include aneuploidy, deletions and amplifications of chromosomal regions, loss of heterozygosity (LOH), microsatellite alterations, point mutations and aberrant promoter methylation. Of interest, several of these genetic changes (i.e. aberrant promoter methylation, chromosome deletions, aneuploidy, altered expression of oncogenes, tumor suppressor genes, and other) occur during the premalignant stage that leads to malignant tumor cells and as such these changes may be useful markers in early detection of lung cancer (59-64). This indicates that several of the molecular, genetic and epigenetic changes that lead to malignant tumor cells occur early in the process, and as such these changes may be useful markers in early diagnosis of cancer. In fact, there is a considerable effort to use these genetic alterations as molecular biomarkers for early cancer diagnosis by applying different approaches. An ideal tumor marker should be highly sensitive, tumor-specific, easy to handle, and inexpensive.

While previous efforts, to identify tumor markers, used screening for mutations, LOH, and microsatellite alterations, more recent strategies concentrate on aberrant promoter methylation (65). This is mainly because of the following reasons: i) Aberrant promoter methylation of specific genes has been shown to be an early event in the pathogenesis of lung cancer. For example, in a rat model of lung carcinogenesis (59), 94% of adenocarcinomas induced by a tobacco-specific carcinogen 4-methylamino-1-(3-pyridyl)-1-butanone were hypermethylated at the p16 gene promoter; most importantly, this methylation change was frequently detected in precursor lesions to the tumors, adenomas and hyperplastic lesions. The timing for p16 methylation was recapitulated in human squamous cell carcinoma (SCC) (59), where the p16 gene was coordinately methylated in 75% of carcinoma in situ lesions adjacent to SCCs harboring this change. Moreover, the frequency of this event increased during disease progression from basal cell hyperplasia (17%) to squamous metaplasia (24%) to carcinoma in situ (50%) lesions. Similarly, aberrant methylation of the p16 and/or O-methyl-guanine-DNA methyltransferase promoter was observed in DNA from spum from 100% of patients with squamous cell lung carcinoma up to 3 years before clinical diagnosis (66). Collectively, these observations indicate that assessment of the prevalence of these methylation markers could be useful for prediction of a lifetime risk of developing lung cancer; ii) The methylation pattern of some genes correlates with clinicopathological characteristics of lung cancer patients. For example, lung cancer patients with methylated RASSF1A had an overall shorter survival than patients who had unmethylated RASSF1A (67). Similarly, a direct correlation has been reported between methylation of the APC and DAPK genes with poor survival rate in NSCLC patients (68,69); iii) it is relatively easy to detect the methylation status quantitatively from a variety of samples ranging from, tumor sample to spum to blood, of lung cancer patients (70). However, to date the penetrance of DNA methylation at any single locus has not been high enough to provide great clinical sensitivity.

A major focus of research in cancer diagnosis is to create a panel of methylation markers (rather than study methylation in a single gene) that could be used to detect cancer with high sensitivity and specificity. The first step in developing a panel of DNA methylation biomarkers is to identify patterns of
tumor methylation signatures (71). One approach is a genome-wide screening (72,73), where, after identifying patterns or signatures of genomic changes in cancer tissues or cancer cell lines, candidate genes and/or regions can be selected to be included in the biomarker panel. Another one is the candidate gene approach, where selected tumor associated genes (such as tumor suppressor genes, proto-oncogenes, genes expressing cell cycle proteins, proteins involved in DNA repair, apoptosis, or host defense and other) are screened in samples of cancer tissue for their aberrant methylation. Using the candidate gene approach several genes have been described to be inactivated by promoter methylation in lung cancer, as previously reviewed (74,75). A recent study exemplified the candidate gene approach along with high throughput technologies (76). Aberrant methylation was studied at 1536 specific CpG sites in 371 genes selected based on their biological relevance in lung cancers and normal lung tissues. The selected genes included tumor suppressor genes and oncogenes, genes that are indirectly involved in cancer development (i.e. DNA repair genes), metastasis-inhibitor genes, genes regulated by various signaling pathways, and/or responsible for altered cell growth, differentiation and apoptosis, genes considered to be targets for oncogenic transformation, genes of innate host defense including lung surfactant protein genes, imprinted genes and previously reported differentially methylated genes (29,75,77).

Among the genes identified with this high throughput approach that showed an altered methylation pattern in lung cancer compared to normal lung tissue were the surfactant protein (SP) genes, SP-A and SP-D. These genes, in addition to their role in surfactant-related functions, have been shown to play important roles in the innate host defense of the lung and/or regulation of inflammatory processes. Since infections and ensuing inflammatory processes have been implicated in cancer pathogenesis (78-80), the surfactant proteins, given their physiological relevance in lung health, are good candidate genes for study in lung cancer and their methylation signatures may serve as valuable markers in lung cancer detection.

5. SP-A and SP-D in lung cancer

An essential for life function is carried out by the lung, namely the oxygen and carbon dioxide exchange, where the delicate gas-exchanging alveolar epithelium that lines the terminal lung airspaces allows for efficient diffusion of oxygen and carbon dioxide between inspired air and the pulmonary circulation. The lung via inspired air is virtually in direct contact with the external environment and therefore it is exposed daily to thousands of insults (i.e. bacteria, viruses, allergens, particulate matter, ozone and other) and yet under normal circumstances remains well functioning. A delicate network comprising of various immune cells (i.e. macrophages, neutrophils and other) and molecules of innate host defense together maintain a normal functioning lung via their ability to mount an appropriate inflammatory response to help get rid of foreign insulting agents. Inflammation under normal circumstances is self-limiting and it subsides as soon as foreign material is removed. However, an inappropriate response (whether less than optimum or more than optimum) can have a negative impact on lung structure and function. A suboptimal inflammatory response may lead to persistence or continuous presence of an infection or foreign matter in the lung, while an overzealous inflammation may take the form of chronic inflammation. Either scenario of deranged host defense results in cellular and/or tissue damage that may ultimately lead to lung cancer or other pulmonary diseases. The relationship between inflammation and lung cancer is not new. It was first proposed by Virchow in 1868, when he first observed leukocytes in neoplastic tissue (81). Since then several cancer types have been shown to associate with inflammation. Also, there is a growing body of evidence that many malignancies are initiated by infections (78-80). According to an estimate >15% of malignancies can be attributed to infections (79).

Molecules of innate immunity and host defense in the lung include the surfactant proteins (SP-) A or SP-A and SP-D. SP-A is a major protein component (by weight) of the lipoprotein complex called surfactant, which is essential in preventing, at low lung volumes, alveolar collapse via its ability to lower surface tension at the air-liquid interface in the terminal airspaces or alveoli. SP-D co-purifies with surfactant. SP-A and SP-D are hydrophilic proteins [reviewed in (82-85)] and have been shown to play a role in surfactant-related functions of the lung (i.e. surfactant homeostasis, structure, inhibition of surfactant secretion and other) as well as in host defense and/or the regulation of inflammatory processes in the lung. Both have been implicated in diverse activities of pulmonary host defense and regulation of inflammation via their effects on immune cells in the presence or absence of a pathogenic insult (86-93), production of reactive oxygen and nitrogen intermediates (94,95), clearance of apoptotic cells (96-98) and others. Furthermore, both SP-A and SP-D have been shown to provide a link between innate and adaptive immunity (99,100), via their effects on dendritic cells.

Study of the SP-A-/- mice has provided in vivo support of the role of SP-A in innate host defense. These mice lack tubular myelin, a structural form of surfactant, and although they are able to survive with no apparent pathology in a sterile environment (101) and respond similarly to wild-type mice following exercise or hyperoxic conditions (102), their pulmonary immune responses are insufficient during immune challenge (103,104). SP-D-/- mice on the other hand although early in life appear healthy with no differences in the biophysical function of surfactant, later in life exhibit quite a complex phenotype. Even in the absence of any apparent insult, the SP-D-/- mice display phenotypic abnormalities in alveolar macrophages and type II epithelial cells, and increased lipid pools, indicating that SP-D has an important role in surfactant homeostasis (101,105,106). Metalloproteinases are also elevated in their lungs. SP-D-/- lungs develop an emphysema-like phenotype (105). Therefore, collectively, given their role in infection and inflammation, alterations in the level or activity of lung collectins (SP-A and SP-D) may lead to pathogenesis of various diseases, including lung cancer.

In human, SP-D is a single gene locus, whereas the SP-A locus consists of two functional genes, SP-A1 and SP-A2 with several intragenic haplotypes or variants identified for each SP-A gene (107,108). The in vitro expressed products of the SP-A1 and SP-A2 genes have been shown to exhibit different
levels of activity in terms of their ability to modulate cytokine production by a macrophage-like cell line (109-111), enhance phagocytosis of bacteria by alveolar macrophages (112-114), inhibit secretion of surfactant by epithelial type II cells (115) and bind to carbohydrates (116). Given the observed differences in activity between SP-A1 and SP-A2, it is possible that the overall functional activity of SP-A in the lung depends on the relative functional content of each SP-A1 and SP-A2 product and that derangement in SP-A1 and/or SP-A2 expression or functional ability compromises lung health. In a recent study, we showed that the SP-A1 to total SP-A ratio, in samples of bronchoalveolar lavage from different individuals, differs as a function of lung health status and age (117). Moreover, both SP-A1 and SP-A2 genes have been identified with extensive genetic variability (107) that affects function and/or structure (110,112-116) or regulation (118-121). This extensive genetic variability may serve as 'tag' to identify individuals at disease risk and also provide insight into the underlying mechanisms of a given lung disease.

6. Methylation signatures in SPs

Although the study of methylation signatures of SPs in lung cancer is recent, the role of SPs in lung cancer has been under investigation for >10 years. Several studies based on different approaches including, reverse transcriptase PCR, immunoblot analysis, ELISA and immunohistochemical analysis have been carried out to study alterations of SP-A in lung cancer (122-128), while there is a single study (129) that has reported usefulness of SP-D as a diagnostic marker for lung cancer. Immunohistochemical detection of SP-A in lungs (130) and pleural effusions (131) has been reported to be useful for differential diagnosis of lung cancer from metastatic carcinomas to lung and pleural mesotheliomas.

Various genetic variants and/or epigenetic changes of SPs may under certain micro-environmental conditions, alter surfactant protein expression and/or function and these changes in turn may modulate susceptibility to lung cancer. Studies carried out by our group and others support this postulate. We have shown an association of lung cancer susceptibility with different SP-A variants (132) and a high resolution comparative genomic hybridization (CGH) analysis of a cDNA microarray showed deletion of the SP-A gene to be one of the most common genomic changes in primary lung cancer (133). The latter genomic aberrations were found to be associated with tumor progression and a history of smoking and were proposed as useful biomarkers in the identification of poor prognoses in patients with NSCLC (134). Recently, using a high-throughput technology platform, consisting of miniaturized bead-based array, we studied the methylation level of 1536 specific CpG sites in a panel of 371 genes (76). The panel included the lung surfactant protein genes, SP-A1, and SP-D, as well as the two hydrophobic surfactant protein genes, SP-B and SP-C, along with other genes documented to be involved in cancer and shown to be differentially methylated (29,75,77) in cell lines, normal tissues, or lung biopsy cancer samples. This study led to identification of a panel of 55 methylation markers that included surfactant protein (SP-) methylation markers. With these markers, lung adenocarcinomas could be distinguished from normal lung tissues with high specificity (76).

With regard to SP gene methylation, a total of 4 groups were obtained, when the collective DNA methylation profile was assessed, by clustering analysis, for 11 CpG sites of lung surfactant proteins (SP-A1, SP-B, SP-C and SP-D) in 23 adenocarcinoma tissues and 23 matched non-cancerous lung tissues obtained from the same patient (32). The percentage of cancer samples increased from group 1 to group 4, while the overall level of methylation for the SP markers tested decreased as one moved from group 1 to group 4. This indicated an inverse association of the collective level of methylation of the 11 SP CpG sites with lung adenocarcinoma. A similar pattern was observed in the case of squamous cell carcinoma compared to matched control tissues obtained from the same patient, although the clustering was not as tightly defined. A further comparison of the methylation levels of each CpG site in cancerous and non-cancerous samples revealed that from the 11 CpG markers analyzed, only 4 CpG sites (namely SP-A1_1370, SP-A1_1080, SP-D_1170, and SP-D_1370) differed significantly between cancerous and non-cancerous samples in both adenocarcinoma and squamous cell cancer cohorts, indicating that these 4 CpG sites can serve as potential biomarkers for lung cancer diagnosis. Of interest, a better group separation was obtained when the methylation content of all 11 SP CpG sites were included in the analysis compared to that obtained with only the four CpG sites that showed significant differences between cancer and non-cancer tissues. Therefore, at this point, we can not exclude the possibility that inclusion of small changes in methylation of certain genes may help improve the specificity of methylation panels of genes identified with significant CpG changes.

A preliminary analysis of the impact of individual CpG methylation on gene expression revealed an inverse correlation (32) between the methylation content and mRNA expression for one of the SP-D CpGs (SP-D_1170). This CpG showed significant changes in methylation between cancer and normal lung tissue. An association between the methylation content of an SP-A1 CpG (SP-A1_1468 that did not show significant differences between cancer and normal tissue) and mRNA expression was also shown. These preliminary observations indicate that methylation of a given CpG site may be one of the mechanisms that regulates SP-A1 and SP-D expression. However, the details of the mechanisms via which an overexpression of SP-A and/or SP-D (presumably a result of promoter hypomethylation) may contribute to lung cancer development are currently unknown.

It is possible that SP-A and SP-D contribute to deranged immunity in cancer via their role on dendritic cell maturation (99,100). Dendritic cells in lung tumor infiltrates appear to be largely defective regarding their ability to stimulate T-cells (135). The effect of SP-A and SP-D on dendritic cell maturation and/or activity appears to be in opposite direction (99,100) indicating that together SP-A and SP-D provide under normal conditions the necessary balance required for a healthy immune lung response. Although hypomethylation at both SP-A1 and SP-D CpG sites was observed in lung cancer (32) and an association was shown between hypomethylation
of certain CpG sites and SP expression, there is no information available to indicate whether the changes in expression are similar between SP-A1 and SP-D. Also information is lacking with regards to the role of SP-A2 CpG methylation in lung cancer. It is currently unknown whether any SP-A2 CpG sites are hyper- or hypomethylated and whether these associate with lung cancer. Moreover, as discussed above, SP-A1 and SP-A2 exhibit differences in their ability to modulate macrophase function with regards to proinflammatory cytokine production (92,109-111) and bacterial phagocytosis (112-114). Since the level of SP-A1 and SP-A2 differences on macrophase function depends on SP-A protein concentration and functional integrity, an altered SP-A expression or SP-A dysfunction, as it occurs after SP-A is exposed to ozone, may adversely affect macrophage host defense function. This may lead to an overall deranged lung host defense and perhaps lung cancer. Thus, although deranged SP-A and SP-D expression may adversely affect lung immunity via their effects on dendritic cells, macrophages, or other cells, our current knowledge is limited to further comment on this.

7. Summary and expert opinion

The available data are consistent with a potential contribution of SP-A in lung cancer. We postulate that a hyperactivity of SP-A via increased expression as it may occur via hypomethylation (32), or a decrease of SP-A activity as it may occur via SP-A dysfunction due, for example, to oxidation (109,111,112), or lack of SP-A as it may occur by SP-A gene deletion (133) may contribute to lung cancer pathogenesis. An excessive SP-A activity may lead to an exuberant and sustained inflammatory reaction that may have dire consequences on lung tissue integrity. On the other hand, reduced SP-A activity or lack of SP-A activity may compromise the innate host defense ability of the lung to maintain a healthy lung status. In either case, where SP-A (and/or SP-D) over-functions or under-functions, the host defense ability of the lung is compromised and with time and/or in the presence of various macro- or micro-environmental insults, this may contribute to lung carcinogenesis.

Although the role of innate host defense molecules, such as SP-A and SP-D, in cancer is still in its early stages, the available literature argues for an inclusion of SP-A and SP-D in future studies, where panels of methylation markers for early lung cancer diagnosis are being considered. The SP-A (and/or SP-D) panel may be more useful in the early diagnosis of lung cancer than the SP-A1 and SP-D panel. Since environmental insults have been shown to contribute to lung cancer development, inclusion of genes responsive to such insults may also be warranted. Moreover, as we may move towards ‘individualized medicine’ selection of a panel of markers for diagnostic use, may require prior understanding of the individual’s life style. Such a knowledge may help determine which gene-environment interactions are more relevant to that individual and consequently choose a panel most appropriate for that individual. Finally, we expect that as our knowledge on the subject increases many more factors and considerations are likely to enter the design and selection of diagnostic methylation marker panels.

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