Genetic variants in the plasminogen activator inhibitor-1 gene are associated with an increased risk of radiation pneumonitis in lung cancer patients

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Introduction

Radiotherapy is a common modality for treating lung cancer [1]. Radiation pneumonitis (RP) is a major side effect associated with radiotherapy which limits the therapeutic ratios of tumor treatment and reduces the living quality in patients who are irradiated for lung cancer. Approximately 16–30% of lung cancer patients experience severe RP after thoracic irradiation [2]. Therefore, the exploration and application of RP biomarkers may help maximize efficacy and minimize adverse effects of radiotherapy. Previous studies have investigated and identified multiple therapeutic and patient-related factors that are associated with the incidence of RP including chemotherapy, smoking status, chronic lung disease, dosimetric parameters, and transforming growth factor (TGF)β plasma concentrations [3–8]. However, only a small proportion of patients exposed to similar doses and volumes of irradiation develop RP, indicating that genetic factors perform a crucial role in the RP process. Our previous studies found that single-nucleotide polymorphisms (SNPs) in the TGFβ pathway were associated with RP risk [9, 10]. Here, we expand upon our previous work by analyzing SNPs in plasminogen activator inhibitor-1 (PAI-1), an important target gene of TGFβ with RP risk [11].

Abstract

Plasminogen activator inhibitor-1 (PAI-1) plays a crucial role in the process of lung injury, although its association with radiation pneumonitis (RP) is unclear. We hypothesized that genetic variants in PAI-1 may influence the risk of RP. In this study, 169 lung cancer patients were genotyped for six single-nucleotide polymorphisms in PAI-1 using the Sequenom MassARRAY system. The risk of RP was evaluated by Cox proportional hazards analyses. The cumulative RP probabilities by genotype were assessed using Kaplan–Meier analyses. Univariate and multivariate analyses revealed that PAI-1:rs7242 GT/GG was correlated with an increased occurrence of grade ≥3 RP (crude hazard ratio = 3.331; 95% confidence interval, 1.168–9.497; P = 0.024). Our results indicated that PAI-1:rs7242 in the 3′-untranslated region of PAI-1 can be a predictor of grade ≥3 RP before radiotherapy.
recent study indicated that a truncated PAI-1 protein (rPAI-123) protects against radiation-induced lung injury in a murine model [16]. Overall, these results implied that PAI-1 could be involved in the RP process.

Human PAI-1 is located on chromosome 7q21.3–q22 and consists of eight introns and nine exons. SNPs in PAI-1 may affect the transcriptional activation and plasma concentrations of PAI-1 [17]. Previous studies demonstrated that PAI-1 polymorphisms were associated with keloids, susceptibility to idiopathic interstitial pneumonia, myocardial infarction, and lung cancer prognosis [18–21]. However, no studies have examined how PAI-1 polymorphisms influence the risk of RP. Here, we investigated the association of SNPs in PAI-1 with RP risk in lung cancer patients treated with radiotherapy.

Materials and Methods

Study populations

This prospective study (NCT02490319) included 169 lung cancer patients. All patients received radiotherapy between September 2008 and June 2014 at Tongji Hospital (Wuhan, China). The enrolled patients had an expected survival > 6 months, Karnofsky Performance Status > 60, and received a radiation dose more than 45 Gy. Exclusion criteria included respiratory infection or lung fibrosis, pulmonary emboli, cardiac disease, drug toxicity, and previous thoracic irradiation. The Tongji Hospital Review Board approved our study. All patients enrolled in the study signed written informed consents for DNA and clinical information.

All enrolled patients underwent radiotherapy with a 6-MV linear accelerator (Elekta, Stockholm, Sweden). The total radiation dose was reached by administering 1.5–2 Gy per treatment. Dose–volume histogram data were shown in Table S1. Seventy-nine patients received intensity-modulated radiation therapy. One hundred sixty patients received induction chemotherapy followed by radiation or concurrent chemotherapy and radiation, with 32.9% receiving a gemcitabine/cisplatin regimen, 19% a CPT-11/cisplatin regimen, 19.6% a docetaxel/cisplatin regimen, and 15.8% an etoposide/cisplatin regimen. We used a three-dimensional planning system (Pinnacle software, version 9.2; Philips Healthcare, Cleveland, OH) to delineate critical normal organs and target volumes.

Details of the follow-up schedule and the RP scoring criteria have been described previously [10]. Briefly, RP was diagnosed by two radiation oncologists after reviewing chest X-ray or computed tomography scans, pulmonary function tests, and clinical information, including symptoms, at each follow-up visit. The patients were followed during and 1 month after therapy, then every 3 months. RP was scored according to the Common Terminology Criteria for Adverse Events 4.0. Symptomatic RP interfering with daily activities, or a requirement for oxygen, were defined as grade 3.

Genotyping methods

Genomic DNA from all patients was extracted from peripheral blood via a blood DNA Kit (K1820-01; Invitrogen, Carlsbad, CA). Based on the public HapMap SNP database and HaploView 4.2 software, we searched for SNPs in PAI-1 that had minor allele frequencies greater than 10%, positioned within the 15-kb region or in its upstream or downstream regulatory regions. We found that all eligible SNPs could be captured with r2 > 0.8 by five tagged SNPs: rs2227631, rs2227667, rs2227672, rs2227692, and rs7242. Together with the well-studied functional SNP rs1799768 (or 4G5G) [20], six SNPs in PAI-1 were selected (Table 1). The SNPs were genotyped by the Sequenom MassARRAY system (Agena Bioscience, San Diego, CA) as described previously [10].

Statistical analyses

The time for developing grade ≥3 RP was the endpoint used for this analysis. Data from patients were censored if they did not develop grade ≥3 RP within 1 year. SPSS version 19.0 (IBM, Chicago, IL) was used for statistical analyses. The Cox proportional hazards model was applied to estimate hazard ratios with 95% confidence intervals of different genotypes. Multivariate Cox regression analysis

Table 1. Characteristics of six SNPs selected for analysis.

| SNP ID   | Chromosome | Position  | Allele | Function class |
|----------|------------|-----------|--------|---------------|
| rs2227631 | 7          | 101126257 | G>A    | promoter      |
| rs1799768 | 7          | 101126425 | –>G    | promoter      |
| rs2227667 | 7          | 101131468 | A>G    | intron 3      |
| rs2227672 | 7          | 101132405 | G>T    | Intron 4      |
| rs2227692 | 7          | 101135963 | C>T    | Intron 7      |
| rs7242    | 7          | 101138164 | G>T    | 3′-UTR        |

SNP, single-nucleotide polymorphisms; UTR, untranslated region.
was used to adjust other covariates. Kaplan–Meier analyses were used to evaluate influences of the genotypes on RP between groups by log-rank tests. \( P < 0.05 \) was considered statistically significant in all tests.

**Results**

**Patient characteristics and association with RP**

Table 2 lists characteristics of the 169 (125 male and 44 female) lung cancer patients (114 non-small-cell lung carcinoma and 55 small-cell lung carcinoma). The median age of patients was 57 years (28–78 years). One hundred six (62.0%) of the patients were smokers. Among the 169 patients, 145 (85.8%) had stage III–IV disease, 160 (94.7%) were treated with chemotherapy, and 86 (50.9%) underwent surgery before radiotherapy.

The median follow-up time in this study was 22 months (6–52 months). After treatment with radiotherapy, 32 patients (18.9%) had grade \( \geq 3 \) RP (grades 3, 4, and 5 were found in 29, 1, and 2 patients, respectively). We evaluated the association between clinicopathologic characteristics and grade \( \geq 3 \) RP risk. According to multivariate analysis, \( V_5 \geq 48\% \), \( V_{10} \geq 38\% \), \( V_{20} \geq 24\% \) and a mean lung dose (MLD) \( \geq 15 \) Gy were associated with increased grade \( \geq 3 \) RP risk (\( P = 0.009 \), \( P = 0.019 \), \( P = 0.034 \), and \( P = 0.014 \), respectively). None of the other clinicopathologic characteristics were associated with a risk of RP in this study (Table S1 and Table 3).

**RP and PAI-1 polymorphisms**

The associations between genetic polymorphisms and the risk of grade \( \geq 3 \) are shown in Table 4 using the Cox proportional hazards model. A significant association was found between rs7242 and the risk of grade \( \geq 3 \) RP. Compared with the rs7242 TT genotype, the GT/GG genotypes had increased hazards of grade \( \geq 3 \) RP (\( P = 0.024 \)). We found a similar result after multivariate analyses with adjustment for potential confounding factors of RP. The RP-free survival for grade \( \geq 3 \) RP, according to rs7242 is plotted in Figure 1A. Development of grade \( \geq 3 \) RP was prolonged in the rs7242 GG/GT genotypes, while no associations with grade \( \geq 3 \) RP were found for the other SNPs.

**PAI-1:rs7242 and dosimetric factors**

The cumulative probability of grade \( \geq 3 \) RP on the basis of genotype and \( V_{20} \) as a function of time is shown in Figure 1B. The incidence of RP in patients receiving \( V_{20} \geq 24\% \) and GT/GG genotypes in rs7242 were higher than patients who received \( V_{20} \geq 24\% \) with the TT genotype in rs7242 (\( P = 0.013 \)). We also analyzed the cumulative RP incidence on the basis of MLD and genotypes as a function of time (Fig. 1C). Patients with a
MLD $\geq$ 15 Gy and GT/GG genotypes in rs7242 displayed a higher RP hazard than patients with the TT genotype and a MLD $\geq$ 15 Gy ($P = 0.010$). However, we did not observe this difference in patients who received $V_{20} < 24\%$ or a MLD $< 15.0$ Gy. These results suggest the independent role of rs7242 genotypes in grade $\geq 3$ RP.

**Discussion**

This study examined whether genetic polymorphisms in PAI-1 gene might be associated with an increased risk of RP in lung cancer patients receiving radiotherapy. We believe this is the first finding of an association between...
the presence of rs7242 in the 3′- untranslated region (UTR) of PAI-1 and the risk of grade ≥3 RP. We found that patients with the rs7242 GG or GT genotypes exhibited an increased risk of RP following radiotherapy. Our results also indicated that the association between rs7242 and grade ≥3 RP risk was independent of V20 and MLD. Moreover, a group of patients (GT/GG genotypes in rs7242 and V20 ≥24% or MLD ≥15 Gy) were found with the highest occurrence of grade ≥3 RP.

RP is a common complication following radiotherapy and is characterized by diffuse alveolar damage and subsequent fibrosis with excessive ECM deposition in the lung [22]. PAI-1 is the main inhibitor of the plasmin system and has a crucial role in ECM accumulation by inhibiting fibrinolysis [12]. Although little is known about the association between PAI-1 polymorphisms and RP risk, several facts indicate that this association is biologically plausible. First, genetic variants in PAI-1 influence the plasma levels of PAI-1 and are associated with other inflammatory or fibrotic diseases including keloids, myocardial infarction, and idiopathic interstitial pneumonia [18–20]. Second, PAI-1 is implicated in the development of other radiation injury diseases. For example, there is a high level of PAI-1 in radiation-induced nephrosclerosis and the process of radiation enteritis [23, 24]. PAI-1 knockout mice have better survival and intestinal function compared with wild-type mice in radiation-induced intestinal injury [25]. Finally, PAI-1 is closely regulated by TGF-β1, the cytokine that has a critical role in the RP process [22, 26]. TGF-β1 can regulate PAI-1 expression via SMAD-dependent and -independent pathways in numerous fibrotic diseases [11, 27–31]. Moreover, TGF-β1 increases PAI-1 plasma levels and promotes the epithelial-mesenchymal transition (EMT), while PAI-1 small interfering RNA prevents the TGF-β1-induced EMT in mouse lung epithelial cells [12].

In this study, rs7242 was associated significantly with grade ≥3 RP. The rs7242 polymorphism is located in the 3′-UTR of PAI-1 and is characterized by the substitution of a guanine with thymine. Studies have examined the

### Table 4. Association between PAI-1 genotypes and grade ≥3 RP.

| Polymorphism and Genotype | No. of event | No. of total | Univariate analysis | Multivariate analysis |
|---------------------------|--------------|--------------|---------------------|----------------------|
|                           |              |              | HR  | 95% CL   | P        | HR  | 95% CL   | P        |
| **PAI-1:rs2227631**       |              |              |     |         |          | 1   |         |          |
| GG                        | 10           | 64           | 1.208 | 0.543–2.688 | 0.644 | 1.564 | 0.695–3.519 | 0.279 |
| AG                        | 15           | 81           | 2.389 | 0.909–6.28  | 0.077 | 2.636 | 0.982–7.076 | 0.054 |
| AA                        | 7            | 21           | 1.433 | 0.679–3.027 | 0.345 | 1.974 | 0.840–3.829 | 0.131 |
| AA+AG                     | 22           | 102          | 1.161 | 0.563–2.391 | 0.686 | 1.399 | 0.633–2.834 | 0.445 |
| **PAI-1:rs1799768**       |              |              |     |         |          | 1   |         |          |
| 4G/4G                     | 12           | 71           | 1.247 | 0.590–2.636 | 0.563 | 1.403 | 0.651–3.022 | 0.387 |
| 4G/5G                     | 16           | 76           | 0.848 | 0.239–3.004 | 0.798 | 1.111 | 0.278–3.875 | 0.956 |
| 5G/5G                     | 3            | 21           | 1.161 | 0.563–2.391 | 0.686 | 1.339 | 0.633–2.834 | 0.445 |
| **PAI-1:rs2227667**       |              |              |     |         |          | 1   |         |          |
| AA                        | 13           | 54           | 0.867 | 0.425–1.770 | 0.695 | 0.816 | 0.390–1.710 | 0.590 |
| AG                        | 18           | 85           | 0.130 | 0.017–0.992 | 0.049 | 0.115 | 0.014–0.924 | 0.042 |
| GG                        | 1            | 29           | 0.668 | 0.330–1.352 | 0.262 | 0.658 | 0.313–1.385 | 0.270 |
| GG+AG                     | 19           | 114          | 0.742 | 0.286–1.927 | 0.540 | 0.777 | 0.249–1.842 | 0.445 |
| **PAI-1:rs2227672**       |              |              |     |         |          | 1   |         |          |
| GG                        | 27           | 135          | 0.777 | 0.387–1.563 | 0.479 | 0.829 | 0.396–1.735 | 0.619 |
| GT                        | 5            | 34           | 0.620 | 0.308–1.246 | 0.179 | 0.673 | 0.316–1.433 | 0.304 |
| **PAI-1:rs2227692**       |              |              |     |         |          | 1   |         |          |
| CC                        | 18           | 76           | 0.777 | 0.387–1.563 | 0.479 | 0.829 | 0.396–1.735 | 0.619 |
| CT                        | 14           | 74           | 0.777 | 0.387–1.563 | 0.479 | 0.829 | 0.396–1.735 | 0.619 |
| TT                        | 0            | 17           | NC    | NC        | 0.971 | NC    | NC        | 0.971 |
| CT+TT                     | 14           | 91           | 0.777 | 0.387–1.563 | 0.479 | 0.829 | 0.396–1.735 | 0.619 |
| **PAI-1:rs7242**          |              |              |     |         |          | 1   |         |          |
| TT                        | 4            | 51           | 2.710 | 0.899–8.165 | 0.077 | 3.558 | 1.127–11.23 | 0.030 |
| GT                        | 15           | 76           | 4.532 | 1.477–13.90 | 0.008 | 5.200 | 1.623–16.66 | 0.006 |
| GG                        | 13           | 42           | 3.331 | 1.168–9.497 | 0.024 | 4.188 | 1.404–12.50 | 0.010 |
| GG+GT                     | 28           | 118          | 3.331 | 1.168–9.497 | 0.024 | 4.188 | 1.404–12.50 | 0.010 |

Multiple analyses in this table were adjusted for sex, age, smoking, surgery, chemotherapy, and V20.

PAI-1, Plasminogen activator inhibitor-1; HR, hazard ratio.

P < 0.05 are presented in bold.

NC not calculated.
relationships between this polymorphism and the risk of myocardial infarction, diffuse-type gastric cancer susceptibility, and primary ovarian insufficiency [19, 32, 33]. In addition, previous studies found that haplotypes of this polymorphism may affect the plasma level of PAI-1 [32, 34]. Other research reported that the rs7242 polymorphism may affect blood insulin concentrations [19]. Because insulin levels play particular roles in lung diseases [35], rs7242 may also modulate the risk of RP by influencing insulin levels in cancer patients.

In summary, these facts suggest that the influence of rs7242 on RP is biologically plausible. However, in this study, we did not observe that other PAI-1 polymorphisms affected the risk of RP. This included rs1799768 that can influence PAI-1 plasma levels and confer an increased risk of several inflammatory or fibrotic diseases such as myocardial infarction, asthma, nephropathy, and idiopathic interstitial pneumonia [20, 36–38]. This finding may have been due to the different nature of the diseases and the small size of the study population.

Our study suggested that the rs7242 polymorphism can be used as a predictor of RP. In combination with our previous findings concerning RP susceptibility and SNPs in TGF\(\beta\)1, ITGB6, PI3CA, AKT2, and MMP1 [9, 10, 39, 40], we can establish a more accurate model using these variants, enabling the prediction of RP by genotyping patients prior to radiotherapy. This would enable patients lacking RP susceptibility genotypes to receive appropriately elevated radiation doses to enhance tumor-related therapies.

In spite of these positive findings, some limitations of our study should be addressed. First, the population of this study was relatively small and thus the results need to be confirmed by further validation. Moreover, we were unable to explore the exact mechanism by which PAI-1 polymorphisms led to RP in lung cancer patients. Finally, as the power in this exploratory study was limited, the \(P\) values in this study were not adjusted using Bonferroni corrections. Therefore, our findings are considered preliminary.

In conclusion, this study identified that rs7242 GT/GG genotypes located in the 3'UTR of PAI-1 were significantly associated with an increased risk of RP in lung cancer patients treated with radiotherapy. Our findings suggested that this polymorphism could be used to predict RP in
lung cancer patients prior to initiating radiotherapy. However, further studies are essential to confirm our findings.

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**Conflict of Interest**

The authors have no conflict of interests.

**References**

1. Killock, D. 2014. Lung cancer: thoracic radiotherapy improves survival in small-cell lung cancer. Nat. Rev. Clin. Oncol. 11:623.
2. Kong, F. M., and S. Wang. 2015. Nondosimetric risk factors for radiation-induced lung toxicity. Semin. Radiat. Oncol. 25:100–109.
3. Palma, D. A., S. Senan, K. Tsujino, R. B. Barriger, R. Rengan, M. Moreno, et al. 2013. Predicting radiation pneumonitis after chemoradiation therapy for lung cancer: an international individual patient data meta-analysis. Int. J. Radiat. Oncol. Biol. Phys. 85:444–450.
4. Stenmark, M. H., X. W. Cai, K. Shedden, J. A. Hayman, S. Yuan, T. Ritter, et al. 2012. Combining physical and biologic parameters to predict radiation-induced lung toxicity in patients with non-small-cell lung cancer treated with definitive radiation therapy. Int. J. Radiat. Oncol. Biol. Phys. 84:e217–e222.
5. Asakura, H., T. Hashimoto, S. Zenda, H. Harada, K. Hirakawa, M. Mizumoto, et al. 2010. Analysis of dose-volume histogram parameters for radiation pneumonitis after definitive concurrent chemoradiotherapy for esophageal cancer. Radiother. Oncol. 95:240–244.
6. Zhang, J., B. Li, X. Ding, M. Sun, H. Li, M. Yang, et al. 2014. Genetic variants in inducible nitric oxide synthase gene are associated with the risk of radiation-induced lung injury in lung cancer patients receiving definitive thoracic radiation. Radiatio. Oncol. 111:194–198.
7. Palmer, J. D., N. G. Zaorsky, M. Witek, and B. Lu. 2014. Molecular markers to predict clinical outcome and radiation induced toxicity in lung cancer. J. Thorac. Dis. 6:387–398.
8. Anscher, M. S., F. M. Kong, K. Andrews, R. Clough, L. B. Marks, G. Bentel, et al. 1998. Plasma transforming growth factor beta1 as a predictor of radiation pneumonitis. Int. J. Radiat. Oncol. Biol. Phys. 41:1029–1035.
9. Yuan, X., Z. Liao, Z. Liu, L. E. Wang, S. L. Tucker, L. Mao, et al. 2009. Single nucleotide polymorphism at rs1982073:T869C of the TGFbeta 1 gene is associated with the risk of radiation pneumonitis in patients with non-small-cell lung cancer treated with definitive radiotherapy. J. Clin. Oncol. 27:3370–3378.
10. Tang, Y., B. Liu, J. Li, H. Wu, J. Yang, X. Zhou, et al. 2016. Genetic variants in PI3K/AKT pathway are associated with severe radiation pneumonitis in lung cancer patients treated with radiation therapy. Cancer Med 5:24–32.
11. Samarakoon, R., J. M. Overstreet, and P. J. Higgins. 2013. TGF-beta signaling in tissue fibrosis: redox controls, target genes and therapeutic opportunities. Cell. Signal. 25:264–268.
12. Ghosh, A. K., and D. E. Vaughan. 2012. PAI-1 in tissue fibrosis. J. Cell. Physiol. 227:493–507.
13. Prabhakaran, P., L. B. Ware, K. E. White, M. T. Cross, M. A. Matthay, and M. A. Olman. 2003. Elevated levels of plasminogen activator inhibitor-1 in pulmonary edema fluid are associated with mortality in acute lung injury. Am. J. Physiol. Lung Cell. Mol. Physiol. 285:L20–L28.
14. Eitzman, D. T., R. D. McCoy, X. Zheng, W. P. Fay, T. Shen, D. Ginsburg, et al. 1996. Bleomycin-induced pulmonary fibrosis in transgenic mice that either lack or overexpress the murine plasminogen activator inhibitor-1 gene. J. Clin. Invest. 97:232–237.
15. Senoo, T., N. Hattori, T. Tanimoto, M. Furonaka, N. Ishikawa, K. Fujitaka, et al. 2010. Suppression of plasminogen activator inhibitor-1 by RNA interference attenuates pulmonary fibrosis. Thorax 65:334–340.
16. Chung, E. J., G. McKay-Corkum, S. Chung, A. White, B. T. Scroggins, J. B. Mitchell, et al. 2016. Truncated Plasminogen Activator Inhibitor-1 Protein Protects From Pulmonary Fibrosis Mediated by Irradiation in a Murine Model. Int. J. Radiat. Oncol. Biol. Phys. 94:1163–1172.
17. Diamanti-Kandarakis, E., G. Palioniko, K. Alexandraki, A. Bergiele, T. Koutsouba, and M. Bartzis. 2004. The prevalence of 4G5G polymorphism of plasminogen activator-1 (SERPINE1) Gene -675 4G/5G and -844 A/G promoter polymorphism with risk of keloid in a Chinese Han population. Med. Sci. Monit. 20:2069–2073.
18. Wang, Y., J. Long, X. Wang, and Y. Sun. 2014. Association of the plasminogen activator inhibitor-1 (PAI-1) Gene -675 4G/5G and -844 A/G promoter polymorphism with risk of keloid in a Chinese Han population. Med. Sci. Monit. 20:2069–2073.
19. Morange, P. E., N. Saut, M. C. Alessi, J. S. Yudkin, M. Margaglione, G. Di Minno, et al. 2007. Association of plasminogen activator inhibitor (PAI)-1 (SERPINE1) SNPs with myocardial infarction, plasma PAI-1, and metabolic parameters: the HIFMECH study. Arterioscler. Thromb. Vasc. Biol. 27:2250–2257.
20. Kim, K. K., K. R. Flaherty, Q. Long, N. Hattori, T. H. Sisson, T. V. Colby, et al. 2003. A plasminogen...
activator-inhibitor-1 promoter polymorphism and idiopathic interstitial pneumonia. Mol. Med. 9:52–56.
21. Di Bernardo, M. C., A. Matakidou, T. Eisen, and R. S. Houlston. 2009. Plasminogen activator inhibitor variants PAI-1 A15T and PAI-2 S413C influence lung cancer prognosis. Lung Cancer 65:237–241.
22. Flechsig, P., M. Dadrich, S. Bickelhaupt, J. Jenne, K. Hauser, C. Timke, et al. 2012. LY2109761 attenuates radiation-induced pulmonary murine fibrosis via reversal of TGF-beta and BMP-associated proinflammatory and proangiogenic signals. Clin. Cancer Res. 18:3616–3627.
23. Brown, N. J., S. Nakamura, L. Ma, I. Nakamura, E. Donnert, M. Freeman, et al. 2000. Aldosterone modulates plasminogen activator inhibitor-1 and glomerulosclerosis in vivo. Kidney Int. 58:1219–1227.
24. Vozenin-Brotton, M. C., F. Milliat, C. Linard, C. Strup, A. Francois, J. C. Sabourin, et al. 2004. Gene expression profile in human late radiation enteritis obtained by high-density cDNA array hybridization. Radiat. Res. 161:299–311.
25. Abderrahmani, R., A. Francois, V. Buard, M. Benderitter, J. C. Sabourin, D. L. Crandall, et al. 2009. Effects of pharmacological inhibition and genetic deficiency of plasminogen activator inhibitor-1 in radiation-induced intestinal injury. Int. J. Radiat. Oncol. Biol. Phys. 74:942–948.
26. Park, J. H., S. H. Ryu, E. K. Choi, S. D. Ahn, E. Park, K. C. Choi, et al. 2015. SKI2162, an inhibitor of the TGF-beta type I receptor (ALK5), inhibits radiation-induced fibrosis in mice. Oncotarget. 6:4171–4179.
27. Liu, R. M., J. Choi, J. H. Wu, K. A. Gaston Pravia, K. M. Lewis, J. D. Brand, et al. 2010. Oxidative modification of nuclear mitogen-activated protein kinase phosphatase 1 is involved in transforming growth factor beta1-induced expression of plasminogen activator inhibitor 1 in fibroblasts. J. Biol. Chem. 285:16239–16247.
28. Liu, R. M. 2008. Oxidative stress, plasminogen activator inhibitor 1, and lung fibrosis. Antioxid. Redox Signal. 10:303–319.
29. He, W., R. Tan, C. Dai, Y. Li, D. Wang, S. Hao, et al. 2010. Plasminogen activator inhibitor-1 is a transcriptional target of the canonical pathway of Wnt/beta-catenin signaling. J. Biol. Chem. 285:24665–24675.
30. Das, F., N. Ghosh-Choudhury, B. Venkatesan, X. Li, L. Mahimainathan, and G. G. Choudhury. 2008. Akt kinase targets association of CBP with SMAD 3 to regulate TGFbeta-induced expression of plasminogen activator inhibitor-1. J. Cell. Physiol. 214:513–527.
31. Zirlik, A., S. Ernst, A. Leugers, F. Willecke, B. E. Sobel, C. Bode, et al. 2009. Inhibition by fibrates of plasminogen activator inhibitor type-1 expression in human adipocytes and preadipocytes. Thromb. Haemost. 101:1060–1069.
32. Jeon, Y. J., Y. R. Kim, B. E. Lee, S. H. Cha, M. J. Moon, D. Oh, et al. 2014. Association of five common polymorphisms in the plasminogen activator inhibitor-1 gene with primary ovarian insufficiency. Fertil. Steril. 101:825–832.
33. Ju, H., B. Lim, M. Kim, S. M. Noh, W. H. Kim, C. Ihm, et al. 2010. SERPINE1 intron polymorphisms affecting gene expression are associated with diffuse-type gastric cancer susceptibility. Cancer 116: 4248–4255.
34. Jeon, Y. J., Y. R. Kim, B. E. Lee, Y. S. Choi, J. H. Kim, J. E. Shin, et al. 2013. Genetic association of five plasminogen activator inhibitor-1 (PAI-1) polymorphisms and idiopathic recurrent pregnancy loss in Korean women. Thromb. Haemost. 110:742–750.
35. Honiden, S., and M. N. Gong. 2009. Diabetes, insulin, and development of acute lung injury. Crit. Care Med. 37:2455–2464.
36. Onalan, O., G. Balta, A. Oto, G. Kabakci, L. Tokgozoglu, K. Aytemir, et al. 2008. Plasminogen activator inhibitor-1 4G/4G genotype is associated with myocardial infarction but not with stable coronary artery disease. J. Thromb. Thromboly. 26:211–217.
37. Cho, S. H., I. P. Hall, A. Wheatley, J. Dewar, D. Abrahaj, J. Del Mundo, et al. 2001. Possible role of the 4G/5G polymorphism of the plasminogen activator inhibitor 1 gene in the development of asthma. J. Allergy Clin. Immunol. 108:212–214.
38. Wong, T. Y., P. Poon, C. C. Szeto, J. C. Chan, and P. K. Li. 2000. Association of plasminogen activator inhibitor-1 4G/4G genotype and type 2 diabetic nephropathy in Chinese patients. Kidney Int. 57: 632–638.
39. Yi, M., Y. Tang, B. Liu, Q. Li, X. Zhou, S. Yu, et al. 2016. Genetic variants in the ITGB6 gene is associated with the risk of radiation pneumonitis in lung cancer patients treated with thoracic radiation therapy. Tumour Biol. 37:3469–3477.
40. Liu, B., M. Yi, Y. Tang, Q. Liu, H. Qiu, Y. Zou, et al. 2016. MMP-1 promoter polymorphism is associated with risk of radiation-induced lung injury in lung cancer patients treated with radiotherapy. Oncotarget 7:7075–70184.

Supporting Information

Additional supporting information may be found in the online version of this article:
Table S1. Association between Dose–volume histogram data and grade ≥ 3 RP.
Table S2. Association between PAI-1 genotypes and grade ≥ 2 RP.