Short Communication

Suppression of Pulmonary CYP2A13 Expression by Carcinogen-Induced Lung Tumorigenesis in a CYP2A13-Humanized Mouse Model

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ABSTRACT

CYP2A13 is a human cytochrome P450 (P450) enzyme important in the bioactivation of the tobacco-specific lung procarcinogen 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK). CYP2A13 expression levels vary dramatically among lung biopsy samples from patients, presumably owing in part to a suppression of CYP2A13 expression by disease-associated inflammation. Here, we determined whether CYP2A13 expression in the lungs of CYP2A13-humanized mice is suppressed by the presence of lung tumors. Tissues from an NNK lung tumor bioassay were examined. CYP2A13-humanized mice (95–100%) had multiple lung tumors at 16 weeks after NNK (30 or 50 mg/kg) treatment; whereas only ~9% of saline-treated CYP2A13-humanized mice had lung tumor (~1/lung). Mice with lung tumors, from the NNK-treated groups, were used for dissecting adjacent tumor-free lung tissues; whereas mice without visible lung tumors, from the saline-treated group, were used as controls. Compared with the controls, the levels of CYP2A13 protein and mRNA were both reduced significantly (by ≥50%) in the NNK-treated groups. The levels of mouse CYP2B10 and CYP2F2 mRNAs were also significantly lower in the dissected normal lung tissues from tumor-bearing mice than in lungs from the control mice. Pulmonary tissue levels of three proinflammatory cytokines, tumor necrosis factor alpha, interferon gamma, and interleukin-6, were significantly higher in the tumor-bearing mice than in the controls, indicating occurrence of low-grade lung inflammation at the time of necropsy. Taken together, these findings support the hypothesis that CYP2A13 levels in human lungs can be suppressed by disease-associated inflammation in tissue donors, a scenario causing underestimation of CYP2A13 levels in healthy lungs.

Introduction

CYP2A13, a functional member of the human CYP2A gene subfamily, is selectively expressed in the respiratory tract (Koskela et al., 1999; Su et al., 2000; Zhu et al., 2006) and is the most efficient cytochrome P450 (P450) enzyme in the metabolic activation of the tobacco-specific lung procarcinogen 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK) (Su et al., 2000; He et al., 2004; Jalas et al., 2005). CYP2A13 protein has been detected in human nasal mucosa and lung (Wong et al., 2005; Zhang et al., 2007). In human lungs, levels of CYP2A13 protein expression were correlated with rates of lung microsomal NNK metabolic activation (Zhang et al., 2007). The CYP2A13*2 allele, which has a decreased level of gene expression, and encodes a variant CYP2A13 protein with reduced activity toward NNK (Zhang et al., 2007; D’Agostino et al., 2008), is associated with a reduced risk of smoking-induced lung adenocarcinoma (Dong et al., 2003). More recently, CYP2A13 was found to mediate NNK-induced lung tumorigenesis in a CYP2A13-humanized mouse model (Megaraj et al., 2014). These findings strongly suggest that CYP2A13 plays an important role in the metabolic activation of NNK in the respiratory tract of human smokers.

A large interindividual variation in the detected levels of CYP2A13 expression (<2–20 fmol/mg of microsomal protein) in human lung biopsy samples was previously reported (Zhang et al., 2007). Given the potential impact of differences in lung CYP2A13 expression levels on the susceptibility to tobacco smoke–induced lung cancer, we have been searching for factors that dictate the apparently large interindividual variations in CYP2A13 expression. Whereas some CYP2A13 genetic variants (e.g., *2 and 7520C>G) are associated with decreased allelic expression in human lung (Zhang et al., 2004; D’Agostino et al., 2008; Wu et al., 2009), we also obtained evidence supporting the hypothesis that CYP2A13 levels in human lung can be suppressed by inflammation associated with disease status in tissue donors (Wu et al., 2013). In the latter study, we demonstrated that the bacterial endotoxin lipopolysaccharide (LPS) can suppress CYP2A13 expression in vitro, in the NCI-H441 human lung cell line, and in vivo, in a CYP2A13-humanized (CYP2A13-transgenic/Cyp2a5-null) mouse model, via a mechanism that involves regulatory sequences in the CYP2A13 promoter region and the nuclear factor NF-κB.

The human lung biopsy samples examined in our previous study were mainly resected, histologically normal lung tissues from patients with lung tumors (Zhang et al., 2007). The aim of this study was to test the hypothesis that CYP2A13 levels in resected nontumor lung tissues can be suppressed by the inflammation induced by the presence of lung tumors, in the CYP2A13-humanized mouse model. Lung tumors were induced in CYP2A13-humanized mice (on A/J background) by a single, intraperitoneal NNK injection. Tumor-bearing mice and control saline-treated, tumor-free mice were examined 16 weeks later for levels of CYP2A13 expression in lung tissues, and for signs of pulmonary or

ABBREVIATIONS: GAPDH, glyceraldehyde-3-phosphate dehydrogenase; IFN, interferon; IL, interleukin; LPS, lipopolysaccharide; NNK, 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone; P450, cytochrome P450; PCR, polymerase chain reaction; SAP, serum amyloid P component; TNF, tumor necrosis factor.

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systemic inflammation. Our results provide further experimental proof to support the idea that the levels of CYP2A13 (and possibly other P450s) detected in patient-derived lung tissues are lower than the levels present in normal, healthy human lung, a notion that has significant implications for efforts to delineate the roles of human lung P450 enzymes in xenobiotic metabolism, pulmonary pathogenesis, and chemical carcinogenesis.

Materials and Methods

Animal Treatments. All procedures involving animals were approved by the Institutional Animal Care and Use Committee of the Wadsworth Center. The conditions and protocols for NNK lung tumor bioassay for the CYP2A13-humanized mouse model (CYP2A13/2B6/2F1-transgenic/Cyp2a5-null; A/J-NS) have been described (Megaraj et al., 2014). Tissues from that project were used for this study. For tumor induction, 8-week-old female mice were treated with a single ip injection of either saline or NNK (either 30 or 50 mg/kg body weight).

Blood Collection, Tissue Dissection, Lung Homogenate Preparation, and Enzyme-Linked Immunosorbent Assay for Cytokine Detection. Post sacrifice, blood was collected from left ventricle and serum was stored at −80°C until use for cytokine determination. Mouse lungs were dissected on ice to remove all tumors visible from the surface; the remaining tumor-free, adjacent lung tissues were quickly frozen on dry ice and stored at −80°C until use. In pilot studies, the dissection procedure (≤5 minutes/mouse) was found to not affect CYP2A13 protein or mRNA levels. The frozen lung tissues were weighed, thawed, and placed into tubes containing 1 ml of T-PER on ice (Tissue Protein Extraction Reagent; Thermo Scientific, Rockford, IL), containing freshly added Halt protease inhibitor single-use cocktail (1×), and homogenized on ice with a POLYTRON (PT 10-35; Kinematica, Bonheim, NY). Lung homogenates were centrifuged at 10,000g for 5 minutes at 4°C; the supernatant was transferred to a new tube and stored at −80°C until use. Interleukin (IL)-6, tumor necrosis factor (TNF)-α, and interferon (IFN)-γ were determined in sera and lung homogenates using the mouse cytokine Duoset ELISA kit (R&D System, Minneapolis, MN). Lung tissue homogenates were prediluted with 1% bovine serum albumin (BSA) in phosphate-buffered saline (PBS) to a final concentration of 1 mg of lung protein/ml.

RNA Isolation and Polymerase Chain Reaction Analysis. Total RNA was isolated from mouse lung and liver using Trizol reagent (Invitrogen, Carlsbad, CA). First-strand cDNA was prepared using the SuperScriptIII first-strand synthesis system (Invitrogen). Briefly, 2.5 μg of total lung RNA or 5 μg of total liver RNA was treated with DNase I (Invitrogen) at room temperature for 15 minutes, and then was mixed with standard amounts of oligo(dT) and other components of the system in final volume of 20 μl. Each reaction mixture was incubated at 50°C for 50 minutes. Polymerase chain reaction (PCR) reactions were carried out essentially as described (Zhang et al., 2007). Levels of various mRNAs were normalized to those of glyceraldehyde-3-phosphate dehydrogenase (GAPDH). Primers used were: GAPDH-F (5′-ggtagcaaggttaggcgca-3′) and GAPDH-R (5′-tgctgtagaggtgatgaggc-3′) (Wei et al., 2012); mSAP-F (5′-tcagcttcacatgattttcag-3′) and mSAP-R (5′-tcagcttcacatgattttcag-3′) (Charles et al., 2006); CYP2B10-F (5′-ccaggtgactctgcaacc-3′) and CYP2B10-R (5′-tgctgtagaggtgatgaggc-3′) (Wei et al., 2012). CYP2A11-F (5′-ggtagcaaggttaggcgca-3′) and CYP2A11-R (5′-tgctgtagaggtgatgaggc-3′) (Pan et al., 2000); CYP2F2-F (5′-ggtagcaaggttaggcgca-3′) and CYP2F2-R (5′-tgctgtagaggtgatgaggc-3′) (Braunning et al., 2009); CYP2A13E6F (5′-tcagcttcacatgattttcag-3′) and CYP2A13E7R (5′-tcagcttcacatgattttcag-3′) (Zhang et al., 2004).

Microsome Preparation and Immunoblot Analysis. Microsome preparation and immunoblot analysis were performed essentially as described previously (Ding and Coon, 1990). Microsomal proteins were separated on NuPAGE Bis-Tris Mini gels (10%) (Invitrogen). Rabbit antibodies against CYP2A5 (Gu et al., 1998) or calnexin (Genscript, Piscataway, NJ) and a goat anti-rabbit secondary antibody (Sigma-Aldrich, St. Louis, MO) were used.

Histopathological Analysis of Dissected Adjacent Lung Tissues. Frozen lung tissues that had been dissected to remove all visible tumors were thawed in Bis-Tris Mini gels (10%) (Invitrogen). Rabbit antibodies against CYP2A5 and immunoblot analysis were performed essentially as described previously (Megaraj et al., 2014). Tissues from that project were used for this study. For tumor induction, 8-week-old female mice were treated with a single ip injection of either saline or NNK (either 30 or 50 mg/kg body weight).

Data Analysis and Statistics. Quantitative data are expressed as means ± S.E. in all groups. For comparison of gene and protein expression in control and tumor-bearing mice, either Student’s t test or one-way analysis of variance (ANOVA) with Dunnett’s post-hoc test was performed, as indicated.

Results

Repression of CYP2A13 Expression in Lungs of NNK-Treated, Tumor-Bearing Mice. Four different NNK doses (30, 50, 100, and 200 mg/kg) were used in the original tumor bioassay (Megaraj et al., 2014), but tissues from only the two lower doses and the saline control group are analyzed here, as the lungs from the two high dose groups had too many tumors, which made it difficult to dissect adjacent “tumor-free” tissues. As reported in the original study (Megaraj et al., 2014), essentially all NNK-treated CYP2A13-humanized mice had observable lung tumors at 16 weeks after the NNK treatment (tumor frequency: 100% and 95%; tumor multiplicity: 13.1 and 4.4 tumors per lung; for the 50 mg/kg- and 30 mg/kg-NNK dose groups, respectively). Approximately 9% of saline-treated CYP2A13-humanized mice also had tumors (each mouse had ~1 tumor per lung). Ten mice with lung tumors, from each of the two NNK-treated groups, were used for dissecting adjacent lung tissues, and three mice without lung tumors, from the saline treated group, were used as controls. The lung tumors were visible from the surface. The identification of lung tumors was confirmed by histopathological analysis (Megaraj et al., 2014). The boundaries of the tumors, which distort the alveolar architecture, were usually well demarcated, allowing easy removal of the tumors during dissection.

The “tumor-free,” adjacent lung tissues are defined as tissues without visible tumors. In a pilot study, we assessed the potential presence of smaller tumors, which were too small to be seen, by performing histologic analysis of dissected lung tissues from four NNK-treated mice (one lung lobe each). By examining serial sections through the entire lung lobes from which visible tumors had been removed, we found two to five additional (small) tumors lesions per lung lobe. However, these small tumors lesions were found to correspond to, on average, no more than ~2% (on the basis of space occupied) to ~6% (on the basis of cells) of the lung lobes analyzed (Supplemental Table 1).

CYP2A13 protein expression was examined by immunoblot analysis in microsomes prepared from tumor-free lung tissues. CYP2A13 protein was detected in both NNK-treated groups and the saline control group (Fig. 1A). As reported recently, the anti-CYP2A5 polyclonal antibody detects a nonspecific band in the lungs of both Cyp2a5-null mice and the CYP2A13-humanized mice (Wei et al., 2013). As expected, the CYP2A13 band was not detected in lung microsomes from the Cyp2a5-null mice, which served as a negative control. Compared with the saline-treated group, the level of CYP2A13 protein was reduced significantly (~60%) in both NNK-treated groups (Fig. 1B).

CYP2A13 mRNA expression was also examined, in total RNA prepared from tumor-free lung tissues. Compared with the saline control group, CYP2A13 mRNA levels were decreased by ~50% in the 30-mg/kg group (though statistical significance was not reached) and by ~80% in the 50-mg/kg group (P < 0.01) (Fig. 1C). Notably, there was no difference in CYP2A13 mRNA level between saline and NNK (30- or 50-mg/kg) groups at 1 week after dosing (Fig. 1D), a result indicating that NNK injection per se did not suppress lung CYP2A13 expression.
The effects of lung tumorigenesis on the expression of mouse P450s were also examined. The levels of murine CYP2B10 and CYP2F2 mRNAs were significantly lower \((P < 0.05)\) in the dissected adjacent lung tissues from tumor-bearing mice than in lungs from saline-treated control mice (Supplemental Fig. 1, A and B). In contrast, the levels of hepatic CYP2B10 and CYP3A11 mRNAs in the tumor-bearing mice were similar to those in the control group (Supplemental Fig. 1, C and D). These results suggest that the lung tumorigenesis had a broad but tissue-specific effect on lung P450 expression.

### Tumor-Related Inflammatory Response in NNK-Treated, Tumor-Bearing Mice.
To evaluate whether systemic inflammation was induced in the tumor-bearing mice, we measured serum levels of three proinflammatory cytokines, IL-6, TNF-\(\alpha\), and IFN-\(\gamma\), by ELISA, as well as mouse hepatic serum amyloid P component (SAP) mRNA levels [known to respond to induction by proinflammatory cytokines (Sunman et al., 2004)] by RNA-PCR. We have recently reported the large effects of systemic inflammation, induced by LPS injection, on serum levels of IL-6 and hepatic levels of SAP in the CYP2A13-humanized mouse (Wu et al., 2013). However, none of the three cytokines were detected, albeit at relatively low levels, in lung tissue homogenates, and their levels were significantly higher \((P < 0.05)\) in tumor-bearing mice than in control mice (Fig. 2, B–D). These data suggest occurrence of a low grade of lung inflammation in the tumor-bearing mice that were treated with NNK at 30 or 50 mg/kg for tumor induction. In other studies not presented, more severe lung inflammation was observed in tumor-bearing mice from the 200-mg/kg NNK group, including tissue injury and mononuclear cell infiltration.

### Discussion
We found no evidence of systemic inflammation in our mouse lung tumor model at the time when tissues were collected for analysis of CYP2A13 expression. This finding was not surprising, as systemic inflammation usually occurs only in cases of advanced cancer (Martin et al., 1999). On the other hand, local inflammation in the lungs of the tumor-bearing mice was evident, as shown by significant elevation in levels of IL-6, TNF-\(\alpha\), and IFN-\(\gamma\) in lung homogenates. Thus, it is highly probable that the suppression of CYP2A13 expression observed in the present lung tumor model was mediated through tumor-derived local inflammation in the lung. In that regard, we have already shown that systemic inflammation, induced by LPS treatment, can suppress lung CYP2A13 expression in the same CYP2A13-humanized mouse model (Wu et al., 2013). Additionally, data in Fig. 1D confirmed that a single-dose...
we have demonstrated for the first time that the expression of human in adjacent, histologically normal lung tissues, rather than a possible loss tumor-bearing mice mainly reflect suppression of CYP2A13 expression observed decreases in CYP2A13 levels in the lungs of NNK-treated, expression level that was actually observed. Thus, we conclude that the was on average only no more than 6% of the total cells, an abundance that total number of tumor cells in the dissected lungs of NNK-treated mice (Ling et al., 2007), we suspect that CYP2A13 expression within the tumor cells was also suppressed. This reduction would contribute to the overall lung tumorigenesis.

NNK injection per se would not repress CYP2A13 expression, prior to lung tumorigenesis.

The absence of systemic inflammation in our lung tumor model also explains the tissue-specific suppression of P450 expression in the lung, relative to the liver, of NNK-treated, tumor-bearing mice. Our results contrast with a previous report that tumor-bearing mice had reduced hepatic P450 levels and altered P450 enzyme activity. By using an explant sarcoma in a transgenic mouse model of human CYP3A4 regulation, Charles et al. (2006) demonstrated an association between the reduction in CYP3A4 expression and occurrence of tumor-derived inflammation. However, in the xenograft model of solid tumors used, the tumor mass reached ~3 g or 10% of total body weight over 17–21 days, and acute-phase response was induced in the tumor-bearing mice. In contrast, the tumor-bearing mice in our study had normal weight gain (at the two doses examined) and the volume of tumor lesion was very small, which explains the lack of a systemic inflammation.

In this study, the dissected “tumor-free” lung tissues might have contained some additional tumors that were too small to be seen. Given that CYP2A13 is normally not expressed in human lung cancer cell lines (Ling et al., 2007), we suspect that CYP2A13 expression within the tumor cells was also suppressed. This reduction would contribute to the overall decrease in CYP2A13 levels detected in the dissected “tumor-free” lung tissues from the tumor-bearing mice. Nonetheless, we estimated that the total number of tumor cells in the dissected lungs of NNK-treated mice was on average only no more than 6% of the total cells, an abundance that would not explain the more than 50% decreases in overall CYP2A13 expression level that was actually observed. Thus, we conclude that the observed decreases in CYP2A13 levels in the lungs of NNK-treated, tumor-bearing mice mainly reflect suppression of CYP2A13 expression in adjacent, histologically normal lung tissues, rather than a possible loss of CYP2A13 expression in the tumor itself.

In summary, by studying a CYP2A13-humanized mouse model, we have demonstrated for the first time that the expression of human CYP2A13 transgene was repressed in the presence of lung tumors. Accompanying the decrease in CYP2A13 expression, three proinflammatory cytokines (IL-6, TNF-α, and IFN-γ) were upregulated in the lungs of tumor-bearing mice. These findings further support our hypothesis (Wu et al., 2013) that the low expression of CYP2A13 observed in the resected, histologically normal lung biopsy tissues from patients with lung cancers (Zhang et al., 2007) was caused at least in part by the tumor-induced chronic inflammation in the lung. In that regard, most studies on gene expression in human lungs use tissue samples that were available from patients suffering from lung diseases or from organ donors who had been subjected to surgical conditions that could induce lung inflammation. The results of these studies are commonly taken to represent tissue levels of gene expression in people (healthy or not). However, our findings imply that, for CYP2A13, the detected levels in available human lung tissue samples are much lower than the levels in healthy, intact human lungs.

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References


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Suppression of Pulmonary CYP2A13 Expression by Carcinogen-induced Lung Tumorigenesis in a CYP2A13-humanized Mouse Model

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Supplemental Table I

Extent of contamination of resected, “tumor-free” lung tissues by tumorous lesions

Lung lobes from NNK-treated CYP2A13-humanized mice (one each at 30, 50, 100, or 200 mg/kg NNK), already dissected to remove all visible tumors, were subjected to histopathological analysis. Ten serial sections, evenly spaced to cover an entire lung lobe, were obtained from each lung and stained with H&E. Areas occupied by residual tumorous lung lesions, the total area of each section, the number of cells within lung tumors, and the total number of cells on each section, were measured. The portion of the section that are occupied by tumorous lesions and the fraction of all cells that are within residual lung tumors are shown.

<table>
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<tr>
<th>Sample No.</th>
<th>NNK dose (mg/kg)</th>
<th>No. of residual tumors</th>
<th>Area occupied by tumors (% of total area)</th>
<th>Fraction of cells within tumors (% of all cells)</th>
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<td>2</td>
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<td>200</td>
<td>5</td>
<td>1.7</td>
<td>2.8</td>
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Supplemental Fig. 1

Effects of lung tumorigenesis on expression of mouse CYPs in the lungs and livers of CYP2A13-humanized mice. Mice (2-month old females) were treated with saline (control) or NNK (either 30 or 50 mg/kg, combined as one group) to induce lung tumorigenesis, and tissues were obtained 16 weeks later. Dissected lung tissues with no visible tumors (from individual mice) and livers from the same groups of mice were used for RNA preparation. Quantitative RNA-PCR analysis was performed, for determination of relative levels of CYP2B10 (A) and CYP2F2 (B) in the lung and CYP2B10 (C) and CYP3A11 (D) in the liver. Data represent means ± S.E. (n=3 for saline/lung, n=14 for NNK/lung, n=4 for saline/liver, n=10 for NNK/liver), normalized by the level of murine GAPDH. *, p < 0.05; compared to the saline group; Student’s t-test.
Supplemental Fig. 1

A

Lung CYP2B10 mRNA (arbitrary units)

Saline
NNK

B

Lung CYP2F2 mRNA (arbitrary units)

C

Liver CYP2B10 mRNA (arbitrary units)

D

Liver CYP3A11 mRNA (arbitrary units)