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Yongjun Yin
Washington University School of Medicine in St. Louis

Xiaodi Ren
Eli Lilly and Company

Craig Smith
Washington University School of Medicine in St. Louis

Qianxu Guo
Eli Lilly and Company

Maria Malabunga
Eli Lilly and Company

See next page for additional authors

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Inhibition of fibroblast growth factor receptor 3-dependent lung adenocarcinoma with a human monoclonal antibody

Yongjun Yin1, Xiaodi Ren2, Craig Smith1, Qianxu Guo3, Maria Malabunga4, Ilhem Guernah4, Yiwei Zhang5, Juqun Shen5, Haijun Sun6,*, Nabil Chehab4,‡, Nick Loizos4, Dale L. Ludwig6 and David M. Ornitz1,§

ABSTRACT

Activating mutations in fibroblast growth factor receptor 3 (FGFR3) have been identified in multiple types of human cancer and in congenital birth defects. In human lung cancer, fibroblast growth factor 9 (FGF9), a high-affinity ligand for FGFR3, is overexpressed in 10% of primary resected non-small cell lung cancer (NSCLC) specimens. Furthermore, in a mouse model where FGF9 can be induced in lung epithelial cells, epithelial proliferation and ensuing tumorigenesis is dependent on FGFR3. To develop new customized therapies for cancers that are dependent on FGFR3 activation, we have used this mouse model to evaluate a human monoclonal antibody (D11) with specificity for the extracellular ligand-binding domain of FGFR3, that recognizes both human and mouse forms of the receptor. Here, we show that D11 effectively inhibits signaling through FGFR3 in vitro, inhibits the growth of FGFR3-dependent FGF9-induced lung adenocarcinoma in mice, and reduces tumor-associated morbidity. Given the potency of FGF9 in this mouse model and the absolute requirement for signaling through FGFR3, this study validates the D11 antibody as a potentially useful and effective reagent for treating human cancers or other pathologies that are dependent on activation of FGFR3.

KEY WORDS: Lung cancer, NSCLC, Fibroblast growth factor receptor 3, FGFR3, Adenocarcinoma, Inhibitory monoclonal antibody

INTRODUCTION

The fibroblast growth factor (FGF) signaling pathway is essential for organogenesis, tissue homeostasis, repair, and injury-induced angiogenesis. Abrupt activation of the FGF signaling pathway is linked to genetic disease and cancer (Dieci et al., 2013; Ellman et al., 2013; Lemieux and Hadden, 2013; Oladipupo et al., 2014; Ornitz and Itoh, 2015; Turner and Grose, 2010). The interactions of FGF receptors (FGFRs) with eighteen signaling FGF ligands regulates cell proliferation, differentiation, shape, and movement (Ornitz and Itoh, 2015). The precise cellular response to FGFR signaling, however, depends on the cell type, developmental stage and physiologic status of the organism.

Activation of the FGF signaling pathway has been implicated in animal and human cancers, where it is of considerable interest as a potential target for therapeutic intervention (Dieci et al., 2013; Liang et al., 2013; Turner and Grose, 2010). In human lung adenocarcinoma, whole genome sequencing ranked FGFR3 among the top 25 significantly mutated genes (Imaiinski et al., 2012). Additionally, increased FGFR3 expression, mutations and gene fusions were observed in primary human NSCLC, and in cell lines derived from these tumors (Liao et al., 2013; Majewski et al., 2013); however, it has not been proven that FGFR3 activation is a driver event in these cancers. Additionally, increased expression of FGFR1, FGFR2 and FGFR3 have been implicated in the acquisition of resistance to activating mutations in the epidermal growth factor receptor (EGFR) family (Kono et al., 2009; Marek et al., 2009; Olveras-Ferraros et al., 2012; Terai et al., 2013; Ware et al., 2013, 2010). Overexpression, activating mutations and activating gene fusions in FGFR3 have also been identified in multiple myeloma, glioblastoma multiforme, bladder, cervical, gastric, colorectal, head and neck squamous, and germ cell-derived cancers (Dieci et al., 2013; Ornitz and Itoh, 2015; Turner and Grose, 2010). Mutations in FGFR3 have also been identified as an escape pathway for inhibitors of B-Raf in melanoma (Yadav et al., 2012).

FGF9 is a potent ligand for FGFR3 (Hecht et al., 1995; Ornitz et al., 1996). Like FGFR3, FGF9 expression has also been identified in several tumor types, including breast, prostate, endometrioid and lung (Hendrix et al., 2006; Li et al., 2008; Marek et al., 2009; Ohgino et al., 2014), suggesting an important role in tumorigenesis. Additionally, expression of FGF9 in lung cancer was associated with poorer prognosis (Ohgino et al., 2014). To model potential oncogenic roles for FGF9, an inducible transgenic system was designed to express FGF9 in adult lung epithelium (White et al., 2006; Yin et al., 2013). Induction of FGF9 expression in adult mice resulted in the rapid conversion of cells in the bronchioalveolar duct junction into proliferative cells thought to have progenitor properties that co-express surfactant protein C (Sftpc), club cell antigen 10 (CC10, Sgb1a1) and Sca-1. Further, rapidly expanding epithelial tumors could be identified within 24-48 h of FGF9 induction. Analysis of these tumors indicated a papillary adenocarcinoma histology and expression of Sftpc, but not CC10. Moreover, genetic studies showed that the formation of these tumors was absolutely dependent on FGFR3 (Yin et al., 2013). The rapid formation of tumors and specificity for FGFR3 indicated that this model could serve as a highly stringent system to test therapeutic agents that target FGFR3 or FGF9.

In this study, we characterize a human monoclonal antibody (D11) that targets the extracellular domain of FGFR3, where it blocks ligand binding and ligand-induced signaling of both major...
splice variants of FGFR3. Using the FGF9-inducible mouse model, we show that treatment with the D11 monoclonal antibody can be used to prevent the initiation of tumors and slow the progression of tumors after induction of FGF9. Furthermore, treatment with D11, improved tumor-associated weight loss, reduced macrophage infiltration into lung tissue and reduced cell proliferation in the bronchioalveolar duct junction.

RESULTS
Characterization of a ligand-blocking anti-FGFR3 human monoclonal antibody
To further evaluate the role of FGFR3 in tumorigenesis and to explore the therapeutic potential of targeting this receptor, we screened a human Fab phage display library and selected an anti-hFGFR3 fully human monoclonal antibody (IMC-D11). D11 bound to human FGFR3 major splice variants FGFR3b and FGFR3c extracellular domain-Fc fusion proteins with an EC50 of ~0.1 nM, and showed minimal binding to FGFR1, FGFR2, or FGFR4 extracellular domains (Fig. 1A). Additionally, D11 bound to human and mouse FGFR3b (Fig. 1B) or FGFR3c (Fig. 1C) with similar affinities. Finally, surface plasmon resonance analysis, a method for measuring protein interactions (Patching, 2014), indicated that D11 had similar binding affinity to murine, rat, cynomolgus monkey and human FGFR3b or FGFR3c (data not shown).

To determine whether D11 could recognize FGFR3 splice variants expressed on the cell surface, we examined antibody interactions with BaF3 cells that stably express FGFR3b or FGFR3c. Flow cytometry analysis showed that D11 bound equally well to BaF3 cells that expressed either the FGFR3b or FGFR3c splice variants on their cell surface, but not to parental BaF3 cells that lack FGFR expression (Fig. 1D). Importantly, the binding of D11 to FGFR3 also blocked the subsequent binding of FGF1 and FGF9 to both FGFR3b and FGFR3c (Fig. 1E-H) with an IC50 of ~1-2 nM. To determine whether the inhibition of ligand binding was sufficient to inhibit receptor activation, FGFR3-expressing BaF3 cells were assayed for survival and growth in response to FGF1 or FGF9 in the presence of increasing concentrations of D11. Consistent with inhibition of ligand binding, D11 also effectively inhibited the mitogenic response of FGFR3-expressing BaF3 cells to FGF1 or FGF9 with an IC50.
ranging from 1.6 to 41.3 nM (Fig. 2A-D). D11 had no detectable agonist activity under tested conditions (data not shown).

We next examined the growth of UMUC-14 cells (Liebert et al., 1989; Miyake et al., 2007), a bladder carcinoma cell line that contains the constitutive activating point mutation, FGFR3$^{S249C}$, in the presence of increasing concentrations of D11. D11 treatment significantly inhibited UMUC-14 cell growth in vitro in a dose-dependent manner compared with human IgG control (Fig. 2E). Importantly, D11 also caused FGFR3 receptor loss, possibly through internalization and degradation, in a dose-dependent manner in UMUC-14 cells (Fig. 2F,G). FGFR3 receptor loss mediated by D11 was also observed on multiple myeloma cell lines, OPM-2 and KMS-11 (data not shown). Thus, D11 was able to inhibit FGFR3 pathway-dependent cell proliferation through blocking ligand binding to receptors, and possibly downregulating cell surface receptors by antibody-induced receptor internalization and degradation.

**D11 inhibits FGF9-dependent lung adenocarcinoma**

To determine whether the D11 antibody can inhibit signaling through FGFR3 in vivo, we employed a mouse model in which lung epithelial hyperplasia and adenomatous tumor formation is dependent on FGFR3 (Arai et al., 2015; Yin et al., 2013). The surfactant protein C–reverse tetracycline activator (Sftpc-rtTA) transgenic allele was used to activate the doxycycline (Dox)-responsive Tre-Fgf9-Ires-eGfp transgene (Tichelaar et al., 2000; White et al., 2006; Yin et al., 2013). In the absence of doxycycline, adult Sftpc-rtTA, Tre-Fgf9-Ires-eGfp double transgenic mice did not express GFP and their lungs were histologically normal (Fig. 3B). In response to doxycycline (provided in chow), Fgf9 and eGFP expression were robustly induced (Fig. 3C). Within one day following doxycycline administration, small eGFP-positive nodules became visible on the lung surface. After four days of doxycycline induction, the lungs fluoresced green under UV illumination. Lung histology showed epithelial nodules with a papillary adenoma-like architecture. The receptor responsible for transducing the FGF9 signal has previously been identified as FGFR3 because induction of FGF9 on an Fgfr3$^{-/-}$ background did not cause epithelial hyperplasia, tumors, or an increase in epithelial proliferation (Yin et al., 2013).

To evaluate the ability of D11 to functionally block the FGFR3 response to induced FGF9, two experimental conditions were
In the lower stringency condition, the D11 antibody was injected intraperitoneally into Sftpc-rtTA, Tre-Fgf9-Ires-eGfp mice (40 mg/kg) beginning two days prior to induction of FGF9 with doxycycline chow (solid line). D11 injections and doxycycline induction continued for an additional 14 days. Mice were euthanized 24 h after the last antibody injection. (B-D) Representative images of whole lungs, corresponding eGFP fluorescence, and histology, from Sftpc-rtTA, Tre-Fgf9-Ires-eGfp mice. (B) Uninduced control mice; (C) control mice induced with doxycycline for 14 days (1 of 9 shown); (D) experimental mice induced with doxycycline for 14 days and injected with D11 antibody (3 of 14 shown). (E) Body weight change of doxycycline-induced control (n=9) and D11 antibody-treated (n=14) mice showing reduced weight loss in D11-treated mice (P<0.0001). (F) Pathology score of doxycycline-induced control (n=9) and D11 antibody-treated (n=14) mice showing improved lung histology in D11 antibody-treated mice (P<0.0001). Scale bars: B-D whole lungs and eGFP panels, 2.0 mm; histology panels, 200 µm.

In the higher stringency condition, the D11 antibody was injected intraperitoneally into Sftpc-rtTA, Tre-Fgf9-Ires-eGfp mice for an additional 14 days and the D11 antibody was repeatedly injected every two days. On day 17 of the experiment (24 h after the last antibody injection), mice were weighed and then euthanized. Plasma was collected to assay for circulating antibody levels and lungs were evaluated for green fluorescence and histopathology.

Control Sftpc-rtTA, Tre-Fgf9-Ires-eGfp mice that were not placed on doxycycline chow showed no eGFP fluorescence, and had normal lung histology (Fig. 3B) consistent with the previously observed very low background expression of the Tre-Fgf9-Ires-eGfp transgene. Sftpc-rtTA, Tre-Fgf9-Ires-eGfp mice that were
given doxycycline chow for 16 days and not injected with antibody or injected with saline showed brightly fluorescent lungs with large bright green nodules (Fig. 3C). These mice showed a significant weight loss \([-3.2±0.3 \text{ g (mean±s.d.)}, n=9]\) over the 17-day experiment. Lung histology showed large tumor nodules, in some cases occupying nearly an entire lobe. In contrast, Sftp/c-rtTA, Tre-Fgf9-Ires-eGfp mice pre-treated with D11 antibody and injected every two days thereafter, showed improved lung histology (Fig. 3D). Importantly, these mice gained weight (0.5±0.5 g, \(n=14\)) over the experimental period (Fig. 3E). Plasma levels of D11 antibody were not detectable in un.injected or saline-injected mice. In D11-injected mice, the mean plasma levels, measured 24 h after the last injection, were 1124 μg/ml (range 660 to 1620 μg/ml).

To semi-quantitatively evaluate the efficacy of D11 treatment, we developed a pathology scoring system (see materials and methods and Fig. S1). In this system, a score of 0 represents normal lung and a score of 5 represents lung tissue that is filled with tumor (no normal lung). Sftp/c-rtTA, Tre-Fgf9-Ires-eGfp mice that were induced with doxycycline and injected with saline showed an average pathology score of 3.8±0.1, \(n=9\) (Fig. 3F). In contrast, mice that were treated with D11 antibody showed a significantly reduced average pathology score of 1.8±0.2, \(n=14\) (\(P<0.0001\) compared with controls).

The efficacy of D11 antibody treatment was next evaluated using a more stringent protocol, in which Sftp/c-rtTA, Tre-Fgf9-Ires-eGfp mice were induced with doxycycline for two days before beginning antibody injections. After two days of doxycycline induction Sftp/rtTA, Tre-Fgf9-Ires-eGfp mice showed small epithelial tumor nodules and widespread epithelial hyperplasia (Fig. S2). After a total of 17 days induction with doxycycline, control mice developed widespread adenomatous tumors (Fig. 4B) and lost significant weight (−6.3±1.1 g, \(n=4\)) (Fig. 4D). These mice had a mean pathology score of 3.9±0.1, \(n=4\). In contrast, mice treated with D11 antibody showed improved pathology (Fig. 4C), average weight gain of 0.4±0.6 g, \(n=10\) (\(P<0.0001\) compared with controls) (Fig. 4D), and a significantly lower average pathology score of 2.3±0.3, \(n=10\) (\(P<0.008\) compared with controls) (Fig. 4E). In D11-injected mice, the plasma levels of D11 antibody, measured 24 h after the last injection, were 1271 μg/ml (range 937 to 1665 μg/ml).

A third experiment was performed to examine tumor regression. Previous studies demonstrated that after doxycycline induction for up to two weeks, followed by doxycycline withdrawal for up to 18 weeks, the established tumor mass remained stable and became independent of FGFR3–FGFR3 signaling (Arai et al., 2015; Yin et al., 2013). To determine whether treatment with the D11 antibody could cause tumor regression, we induced Sftp/c-rtTA, Tre-Fgf9-Ires-eGfp mice with doxycycline for 7 days. Doxycycline chow was then replaced with normal chow and D11 antibody was injected every 2 days for 14 days. Pathology scores of control mice not injected with D11 antibody, or experimental mice injected with D11 antibody, were not different (mean pathology score: without antibody 2.5±1.4, with antibody injection 2.5±1.4), indicating that once tumors were established, inhibition of FGFR3 was not sufficient to accelerate tumor regression of these FGFR3–FGFR3-independent tumors (data not shown).

D11 inhibits the proliferation of Sftp/c-CC10+ bronchioalveolar duct junction cells

Previous studies using the Sftp/c-rtTA, Tre-Fgf9-Ires-eGfp mouse model demonstrated that short-term (16 h) induction with doxycycline resulted in increased proliferation of Sftp/c-CC10+ cells within 25 cells of the bronchioalveolar duct junction (BADV) (Yin et al., 2013). To determine whether D11 antibody treatment could affect proliferation or the number of Sftp/c-CC10+ cells in the bronchioalveolar duct junction region, Sftp/c-rtTA, Tre-Fgf9-Ires-eGfp mice were injected with two doses of D11 antibody 24 h and 9 h before being placed on doxycycline chow for 16 h (Fig. 5A). After 16 h of induction, control Sftp/c-rtTA, Tre-Fgf9-Ires-eGfp mice showed conversion of nearly all epithelial cells in the BADJ region to Sftp/c-CC10+ cells in response to induction of FGFR9 (Fig. 5F-J). However, D11 antibody treatment was associated with a significant (\(P<0.002\)) decrease in PCNA-positive cells in the BADJ region in these mice (Fig. 5K).

Deincreased infiltration of multinucleated macrophages in D11-treated mice

Tumor induction in Sftp/c-rtTA, Tre-Fgf9-Ires-eGfp mice is often associated with the infiltration of large multinucleated macrophages (Fig. 6A,B; Fig. S4) (Arai et al., 2015). Macrophage infiltration is evident by 24 h of induction and often resolves after several weeks,
To determine whether D11 antibody treatment affects immune cell infiltration we developed an immune score scale to evaluate the degree of immune cell infiltration (Fig. S3). Interestingly, treatment with D11 antibody significantly reduced multinucleated macrophage infiltration (Fig. 6E) but had little effect on lymphocyte infiltration (Fig. 6F; Fig. S4). This suggests that multinucleated macrophage infiltration is very sensitive to lung pathology mediated by signaling through FGFR3 whereas lymphocyte infiltration might be secondary to signals mediated by other FGF receptors.

DISCUSSION
The future for treatment of human cancers will require redundant therapies targeted to tumor-specific pathways and the tumor microenvironment. Thus, the identification of signaling pathways required by specific tumor types and the development of therapeutic agents that effectively target these pathways will be essential to the next generation of cancer treatments. Mutation, gene rearrangements and gene amplifications that effectively activate the FGFR3 tyrosine kinase domain are currently associated with several human tumors, including adenocarcinoma of the lung, multiple myeloma, glioblastoma multiforme, nasopharyngeal, bladder, cervical, gastric, colorectal and germ cell-derived cancers (Cappellen et al., 1999; Cheng et al., 2013; Chesi et al., 2001; Ewen et al., 2013; Jang et al., 2001; Kompijer et al., 2010; Ornitz and Itoh, 2015; Singh et al., 2012; Turo et al., 2015; Wang et al., 2014; Williams et al., 2013; Wu et al., 2013). Furthermore, FGF9, a high-affinity ligand for FGFR3 is expressed in a large percentage of lung, prostate, and colon cancers (Leushacke et al., 2011; Ohgino et al., 2014; Teishima et al., 2014).

In a mouse model for adenocarcinoma of the lung, activation of epithelial-expressed FGFR3 by induced expression of FGF9 results in rapid changes in the differentiation state of BADJ cells, increased proliferation of Sftpc+/CC10+ BADJ cells and type II pneumocytes, and the rapid formation of adenomatous tumor nodules (Yin et al., 2013). The rapid development of lung pathology and its complete dependence on signaling through FGFR3 makes this model ideal for testing or screening for drugs that can inhibit the FGFR3 signaling pathway in vivo, such as a blocking antibody to FGFR3.

D11 is a human monoclonal antibody that targets the extracellular domain of FGFR3. We show that the D11 antibody specifically recognizes human and mouse FGFR3 and has very low affinity for other FGFRs. We further show that D11 inhibits a mutant form of FGFR3 harboring a gain-of-function missense mutation, and additionally, inhibits ligand-induced activation of FGFR3. Because D11 also readily cross-reacts with mouse FGF3, to demonstrate potential therapeutic efficacy we showed that mice given serial intraperitoneal injections of D11 over two weeks attained average plasma levels of >1000 µg/ml and showed normal weight gain and no apparent adverse effects of treatment. In a short-term treatment protocol in mice, adverse effects would most likely result from consequences of on-target inhibition of FGFR3, possible off-target interference with other physiologically active molecules, or an acute immune response. The lack of adverse effects of D11 treatment is consistent with on-target activity against FGFR3 and the knowledge that mice that congenitally lack FGFR3 are viable (Colvin et al., 1996). These studies further suggest that D11 has minimal off-target interference with other FGFRs or other non-related proteins.

The Sftpc-rtTA, Terc-Fgf9-Ires-eGfp mouse model is highly stringent test for an anti-FGFR3 drug, in that FGF9 is induced throughout the lung (type II pneumocytes and distal airway epithelial cells), and the resulting lung pathology has a rapid onset characterized by acutely increased epithelial proliferation in distal airways, formation of adenomatous nodules, and inflammatory infiltration. The observation that D11, even when administered two days following FGF9 induction, has the ability to suppress FGFR3-dependent pathology in vivo without adverse side effects suggests that this antibody has significant therapeutic potential.

A previously developed monoclonal antibody, R3Mab, directed towards the FGFR3 extracellular domain (Qing et al., 2009) effectively blocks ligand binding and ligand-induced activation of the wild-type receptor and inhibits ligand-independent FGFR3 with activating missense mutations in the extracellular domain. R3Mab was also effective in inhibiting multiple myeloma cells containing the t(4;14)(p16.3;q32) chromosomal translocation that results in the
expression of an FGFR3-MMSET fusion protein (Chesi et al., 1998; Qing et al., 2009). Other researchers have demonstrated that a single-chain anti-FGFR3 Fv fragment fused to a toxin gene product, rGel, had the ability to inhibit the growth of a xenograft bladder cancer cell line (Martinez-Torrecuadrada et al., 2008). The D11 antibody also effectively inhibits ligand binding, ligand-induced receptor activation, and activation of FGFR3 containing an activating point mutation. Furthermore, D11 causes internalization and degradation of the FGFR3 protein.

In addition to the use of antibodies to target FGFR3, small-molecule inhibitors of the FGFR tyrosine kinase domain have demonstrated efficacy with several xenograft tumor models. For example, the pan-FGFR selective tyrosine kinase inhibitors, SU5402 and PD173074, effectively blocked the growth of t(4;14) (p16.3;q32) multiple myeloma cells (Grand et al., 2004). However, no small-molecule inhibitors have been identified with specificity for FGFR3 over other FGFRs. Thus, the use of antibodies, potentially coupled to toxins or other biologically functional molecules, has the potential to provide more customized and disease-specific therapy while minimizing adverse side effects.

In addition to the role of activating FGFR3 mutations in a variety of cancers, germline-activating mutations in FGFR3 are the etiology of achondroplasia, the most common form of skeletal dwarfism in humans, and somatic activating mutations in FGFR3 cause seborrheic keratosis, a benign skin tumor, and epidermal nevi, a benign hyperplastic skin lesion (Hafner et al., 2006; Logie et al., 2005; Naski et al., 1996). Activation of the C-type natriuretic peptide (CNP) signaling pathway, which antagonizes FGFR3 signaling in chondrocytes, is being evaluated as a therapeutic for achondroplasia (Lorget et al., 2012). Antibodies that effectively inhibit FGFR3 activity in vivo would also be potential candidates to treat skeletal dwarfism and other pathologies associated with achondroplasia. Synergism with other activators of chondrocyte proliferation, such as CNP, might provide a more effective treatment with fewer side effects. This is important, given that treatment of achondroplasia will be required throughout much of the prepubertal growth years of affected children.

MATERIALS AND METHODS
Identification of a human anti-FGFR3 antibody from a phage display library
A human Fab phage display library was panned (Eli Lilly and Company, New York, NY) for anti-FGFR3IIIc antibodies using human FGFR3IIIc-Fc (R&D Systems, Minneapolis, MN; #766-FR-050) as the bait. Individual phage clones recovered after the second and third rounds of selections were examined for binding to immobilized hFGFR3IIIc/Fc and for blocking hFGF3/FGF2 interaction by ELISA. The DNA sequences encoding the heavy- and light-chain variable genes for the selected antibody, IMC-D11 (D11; Eli Lilly and Company), were amplified by PCR and cloned into an expression vector containing human λ light-chain constant region and human γ1 heavy-chain constant region. D11 monoclonal antibody was produced from individual clones of stably transfected CHO cells.

ELISA binding and MSD blocking assays
Various amounts of phage, Fab, or monoclonal antibody were serially diluted in 0.2% Tween 20/PBS containing 1% BSA, and added to hFGFR3IIIc-coated plates (50 µl at 1 µg/ml) and incubated at room temperature for 2 h. The bound antibodies were detected with an anti-human Fab antibody conjugated with HRP (Jackson ImmunoResearch, West Grove, PA; #109-035-097) in blocking assays, or continuously probed with SULFO-TAG-labeled FGF1s (Meso Scale Discovery, Rockville, MD; # R91AN-1) in blocking assays according to the supplier’s instructions. 250 nM NaCl was present in ELISA binding and washing steps, and 10 µg/ml hparin was present in ligand-blocking assays. Human FGF1 (Genway Biotech Inc., San Diego, CA; #GW-B-54AE0), Human FGF9 (R&D Systems; #273-F9).

Cell lines and proliferation assays
Murine pro-B cell line BaF3 parental cells (Mathey-Prevot et al., 1986; Palacios and Steinmetz, 1985) were maintained in RPMI-1640 (Invitrogen, Carlsbad, CA) with 10% FBS (heat inactivated, HyClone, Logan, UT), 5 ng/ml murine IL-3 (R&D Systems); BaF3 cells stably transfected with FGFR3 were maintained in the above medium supplemented with 2 µg/ml puromycin (Sigma-Aldrich, St. Louis, MO). Cell lines were confirmed to be IL3-dependent and puromycin-resistant. No further authentication was carried out on these cells. For flow cytometry, BaF3 cells were incubated with 10 µg/ml D11 and 1:200 dilution of secondary anti-hIgG-PE (Jackson Immunoresearch: #109-116-088). After washing, cells were analyzed on a Guava EasyCyte Plus flow cytometry system (Millipore, Billerica, MA). For IMC-D11-blocking BaF3/FGFR3 cell proliferation assays, 25,000 cells/well were seeded in 96-well plates. After incubation with serially diluted D11 for 1 h, FGF ligands were added to 3.7 nM final concentration for 72 h. Cell viability was determined using the CellTiter-Glo Luminescent Cell Viability Assay (Promega, Madison, WI).

UMUC-14 cells (Liebert et al., 1989), obtained from MD Anderson Cancer Center, were maintained in DMEM medium (ThermoFisher, Invitrogen, Grand Island, NY) supplemented with 10% FBS (HyClone) under conditions of 5% CO2 at 37°C. The mutation in FGFR3 was confirmed by PCR amplification and sequencing. No further authentication was carried out on these cells. For D11-blocking UMUC-14 cell proliferation assay, cells were incubated with 1:3 serial-diluted D11 from 200 nM and/or 200 nM human IgG control antibody (Equatech-Bio, Kerrville, TX; #SLH56) for 72 h. Cell viability was determined using the CellTiter-Glo Luminescent Cell Viability Assay (Promega, Madison, WI).

Western blot
Cell lysates (15 µg) from UMUC-14 cells were subjected to SDS-PAGE followed by western blot. Blots were probed with anti-FGFR3 antibody (1:2,000; Sigma-Aldrich; #F0425) or anti-tubulin antibody (1:2,000; Cell Signaling Technology, Danvers, MA; #2125), and secondary anti-Rabbit IgG HRP antibody (1:10,000; Jackson ImmunoResearch; #111-035-144). Bands were detected and quantified using a Fujifilm LAS-4000 Luminescence Image Analyzer (GE Healthcare Life Sciences, Pittsburg, PA).

Detecting mouse plasma IMC-D11 antibody
Serially diluted plasma samples or purified D11 antibodies were added to goat anti-human IgG-coated plates, and detected with an HRP-conjugated goat anti-human IgG Fc γ-specific antibody (1:10,000; Jackson ImmunoResearch; #109-035-098).

Mice
The Sftpc-rtTA, Tre-Fgf9-ires-eGfp mouse strain has been previously described (Perl et al., 2002; Tichelaar et al., 2000; White et al., 2006; Yin et al., 2013, 2008). Sftpc-rtTA, Tre-Fgf9-ires-eGfp mice were maintained on an inbred FVB genetic background. All mice were housed in a pathogen-free animal facility under the veterinary care of the Department of Comparative Medicine at Washington University School of Medicine, and used at the age of six-to-twelve weeks. All protocols were approved by the Washington University Animal Studies Committee and were performed in accordance with the Animal Welfare Act and the Guide for the Care and Use of Laboratory Animals.

Transgene induction and D11 antibody injection
Doxycycline (Dox) diet was purchased from Bio-Serv Inc. (200 mg/kg green pellets; #S388). Mice were fasted for 6 h prior to providing doxycycline chow at 6 pm on the first day of induction. Doxycycline chow was provided ad libitum throughout the experiment. The D11 antibody was diluted with 1× PBS and injected intraperitoneally at a dose of 40 mg/kg (~0.5 ml/mouse). Control mice were either not injected or injected with 0.5 ml of 0.9% saline. Body weight was measured before injection and on the day of necropsy.
Lung pathology, histology, and immunohistochemistry

Mice were anesthetized with KXA (31 mg/kg ketamine, 6 mg/kg xylazine, 1 mg/kg acepromazine) and transcardially perfused with a vascular rinse of 0.9% NaCl followed by 10% neutral buffered formalin (VWR International; #9370-094). Lungs were dissected and photographed under bright-light illumination and under UV light using an Olympus SZX12-ILLD100 dissecting microscope (Olympus Optical Co. Ltd). Tissues were postfixed in 10% phosphate buffered formalin overnight at 4°C. All five lobes were separated, embedded in paraffin and sectioned using standard procedures. For histology, slides were stained with hematoxylin and cosin (H&E). For immunofluorescence staining, sections were rehydrated. Antigen retrieval was achieved by boiling at 121°C for 15 min in 10 mM citrate buffer followed by gradual cooling to room temperature. Sections were incubated overnight at 4°C with the primary antibodies, and after incubation with the secondary antibodies, appropriate Alexa Fluor-coupled secondary antibodies (Thermo Fisher; #A21206, #A21203, #A21447) were applied and incubated at a 1:200 dilution. Sections were photographed on an ApoTome fluorescence microscope (Zeiss Inc). The following primary antibodies were used for staining: CC10 (Scgb1a1, sc-9772; Santa Cruz Biotechnology Inc.; 1:200); pro-SP-C (SftpC, AB3786; Millipore; 1:2000); CD45 (ab10558; Abcam, Cambridge, MA; 1:200). For the CD45 antibody, immunostaining was detected using the Histostain-SP Broad Spectrum (DAB) kit (Thermo Fisher; #95-9643).

For quantification of cell number, multiple optical sections were scored manually to distinguish cell boundaries and identify the bronchioalveolar duct junction. SftpC- and CC10-positive cells were counted within 25 cells of the bronchioalveolar duct junction as described (Yin et al., 2013). Three different whole-lung longitudinal sections containing the main axial bronchi were scored for each mouse.

Pathology and immunology scores

We developed a histopathological scoring system (Fig. S1 and S3) to provide semi-quantitative analysis of the extent of tumor growth and inflammatory infiltrates (Gibson-Corley et al., 2013). For lung tumorigenesis, we based this scoring system in part on the Ashcroft scoring system, which is commonly used to evaluate the extent of lung fibrosis (Ashcroft et al., 1988; Guzy et al., 2015). For each mouse sample, 2-3 H&E slides were selected from different depths of the tissue, separated by at least 50 μm. Three to five lobes were viewed for each mouse. Two persons scored the slides and were blind to the identity of the mouse or the treatment. The pathology score was defined as:

- 0, normal lung tissue; 1, alveolar or ductal hyperplasia; 2, 1-3 tumor nodules in at least one lobe; 3, many tumor nodules in multiple lobes; 4, solid tumor in at least one lobe; 5, solid tumor, no normal lung (Fig. S1).

The immunology score was defined as:

- 0, no inflammation; 1-L, diffuse lymphocyte infiltration, no nodules; 2-L, peribronchiolar lymphocyte clusters, ≤4 clusters in one lobe with >10 cells per cluster; 3-L, peribronchiolar lymphocyte clusters, ≥4 clusters in one lobe with >10 cells per cluster; 4-M, diffuse multinucleated macrophage infiltration; 2-M, dense multinucleated macrophage infiltration (Fig. S3).

Statistical analysis

All data are expressed as mean±standard derivation (s.d.). Student’s t-test values indicate statistical significance for comparable tissues, P values were obtained using GraphPad Prism and Microsoft Excel software.

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Competing interests

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Supplementary information

Supplementary information available online at http://dmm.biologists.org/lookup/suppl/doi:10.1242/dmm.024760/-/DC1

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