Bronchiolar Adenoma Transforming to Invasive Mucinous Adenocarcinoma: A Case Report

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Abstract: Bronchiolar adenoma (BA) is recognized as a neoplasm with benign clinical course. Histologically, BA is characterized by nodular proliferation of the bilayered bronchiolar-type epithelium, including multipartite epithelial cells and a continuous layer of basal cells. Recent reports have revealed the frequent presence of driver gene mutations in BA, suggesting its neoplastic nature. However, it is still debatable whether BA has malignant potential. Herein, we report the first case of BA harboring the same KRAS mutation with the adjacent invasive mucinous adenocarcinoma (IMA). Additionally, the loss of continuity of the basal cell layer in the junctional zone between BA and IMA indicated a malignant transformation from BA to IMA in this particular case.

Keywords: bronchiolar adenoma, invasive mucinous adenocarcinoma, KRAS, transformation

Introduction
Bronchiolar adenoma (BA) is a newly designated rare entity by Chang et al in 2018, including the currently established ciliated mucinous papillary tumor (CMPT) and the so-called non-classic CMPT. Commonly, BAs present as discrete, sharply circumscribed lesions with a median size of 0.5 cm (0.2 to 2 cm). The most prominent histological feature of BA is the bilayered cell structures composed of the continuous basal cell layer (p40 and CK5/6-positive), and the surface cell layer consists of different proportions of mucinous cells, ciliated cells, Clara cells and type II pneumocytes. The composition of the surface epithelium renders BA with subtypes, including proximal-type and distal-type. The primary differential diagnosis for proximal-type BA with prominent mucinous feature is invasive mucinous adenocarcinoma (IMA). To date, no convincing evidence has been found on whether BA can transform to IMA. Herein, we report the pathological observation and molecular study of a special BA, which may shed new light on this point.

Case Presentation
A 70-year-old man with thirty years of smoking history presented with complaints of shortness of breath. Computed Tomography (CT) revealed a 1.5 cm × 1.4 cm nodule in the middle lobe of the right lung. The lesion was lobulated, with spicule formation, associated with distal airway tracts (Figure 1A, red arrow). After excluding extrapulmonary metastases by imageology, lobectomy was performed. Gross examination revealed a demarcated, gray-white mucoid mass in the peripheral lung. No necrosis, hemorrhage or pleural indentation was noticed.

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Microscopically, the tumor was mainly composed of two patterns: glands lined with proliferative mucinous cells and tall columnar mucinous cells growing along the alveolar wall (Figure 1B). Papillary, micropapillary structures and intra-alveolar mucus can be seen. Based on the skipping growth pattern seen in certain areas (Figure 1C), the intraoperative pathological diagnosis favored IMA.

Postoperative pathological examination showed that, in some regions of glands, especially the center of the tumor, the presence of basal layers could not be ruled out. The luminal layers of the glands were lined predominantly by cuboidal cells, interpreted by ciliated cells and columnar cells (Figure 1D). Intraluminal papillaries composed of mucinous cells could be noted in certain glands (Figure 1E). No distinct boundary was found between the glandular and the lepidic areas (Figure 1F). Hence, immunohistochemistry was performed with a panel including CK7, TTF-1, Napsin A, CK5/6, p63, p40, and Ki-67. The expression of p40, p63, and CK5/6 demonstrated an area composed of continuous basal layer measuring 5×3 mm² (Figure 2A and B). TTF-1 was positive in the columnar and cuboidal cells in the luminal layer of the glands (Figure 2C, red arrows).
Interestingly, in the adjunct lepidic areas, basal cells were absent, as demonstrated by the negative staining of basal markers (Figure 2D–F, black arrows). In the junctional zone, p63, p40, and CK5/6 showed only sporadic expression (Figure 2D–F, purple arrows). The Ki-67 index was similar among the two areas (1%).

Furthermore, these two distinct areas were micro-dissected and analyzed by a panel covering EGFR, ALK, ROS1, KRAS, NRAS, PIK3, HER2, and BRAF (Supplementary Figure 1). The same KRAS mutations (G12V) in both portions were identified (Figure 3), supporting the identical cellular origin of both parts. Therefore, the post-operative diagnosis was revised to BA transforming to IMA.

**Discussion**

The hallmark feature of BA is the continuous basal cell layer,¹ this unique feature sets pitfalls in the differentiation between BAs and lung malignancies, especially in the rapid frozen pathological diagnosis. In proximal-type BA with prominent mucinous feature, epithelial cells can grow in lepidic and skipping patterns over alveolar walls, with abundant mucinous cells and secreted mucin. Because of

![Figure 2 Hematoxylin-eosin (HE) and P40 stain of the glandular area with continuous basal layer, an area of 5×3 mm² was demonstrated (A and B, 40×). The cuboidal and columnar cells in the luminal layer were TTF-1 positive (red arrows; C, 100×). Loss of continuity of the basal cell layers at the BA to IMA junctional zone: red arrows indicate the continuous basal cell layer; purple arrows indicate the sporadic staining of basal cell marker in the junctional area; black arrows indicate the absence of basal cell layer (D–F, 200×).](https://www.dovepress.com/press-center/oncotargets-and-therapy/volume-14)
the histological commonalities, it is difficult to discriminate between the proximal-type BA and IMA without the assistance of immunohistochemistry (IHC). Recognition of a continuous basal cell layer is the key to distinguish these lesions, which can be confirmed by IHC with basal cell markers (p40, p63, and CK5/6) following morphological suspicion. Additionally, the presence of ciliated cells (Figure 1D) would also support the diagnosis of BA in this case.

According to Chang et al, BAs exhibit a morphologic spectrum reflecting either proximal or distal bronchiolar differentiation, with cases showing overlaps. Zheng et al found similar immunophenotypes among classic and non-classic ciliated muco-adenomatous papillary tumor (CMPT), including tripartite differentiation of the epithelium. In this case, the epithelium is lined predominantly by terminal respiratory epithelial cells (TTF-1 expression shown in Figure 2C; diffuse CK7 and focal Napsin A expression are not shown), along with the presence of ciliated cells and abundant mucinous cells. Several markers for mucinous cells are usually positive in CMPTs, including HNF4α, MUC1, and MUC5AC. HNF4α is highly sensitive for identifying neoplastic mucinous epithelial cells. However, its reliability and specificity are questioned because divergent percentages of expression were reported in different papers (ranging from 11% to 92%). Besides, Udo et al found HNF4α positivity in four cases of CMPTs. Additional studies are still needed to validate the positivity of diverse immunomarkers in CMPTs.

Peribronchiolar metaplasia (PBM) is a reactive extension of bronchiolar-type epithelium along the peribronchiolar alveolar walls. Usually, PBM demonstrates as small (diameter < 1mm) and multifocal lesions. Histologically, PBM can resemble atypical adenomatous hyperplasia (AAH) or CMPT, but immunohistochemically, the proliferating epithelium in PBM are usually TTF-1 negative. In the current case, the larger size of the glandular area (5×3 mm²), the positive TTF-1 staining and the mixed cellularity in the luminal layer favored the diagnosis of bronchiolar adenoma.

The genomic profile of BA is considered unique and distinct from lung adenocarcinomas. However, several BA cases have been reported for harboring mutations of common driver genes for lung adenocarcinoma, such as KRAS, BRAF, and EGFR. A relatively high percentage (50%) of BRAF mutation was found in BA despite its benign clinical course. Hence, BRAF mutation in BA merely represents evidence for a neoplastic process and is not synonymous with malignancy.

Figure 3. Quantitative reverse-transcript polymerase chain reaction (qRT-PCR) revealed the same KRAS mutations (G12V) in both BA (A) and IMA (B).
Data Sharing Statement
The datasets supporting the conclusions of this case are included within the article.

Ethical Approval and Consent to Participate
Ethical approval for this study was obtained from the First Affiliated Hospital of China Medical University’s institutional ethics review boards. Writing consent to participate was provided by the patients for the present research.

Patient Consent for Publication
Informed consent was obtained from the patients for the publication of cases and any accompanying images. Copies of the written consent are available for review by the Editor-in-Chief of this journal.

Author Contributions
All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure
The authors declare that they have no competing interests.

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