Gatekeeper Mutations and Intratumoral Heterogeneity in FGFR2-Translocated Cholangiocarcinoma

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Summary: FGFR2 genetic translocations are frequent in cholangiocarcinoma, yet despite initial sensitivity to FGFR inhibitors in clinic, patients quickly become resistant to targeted therapies. The work published by Goyal and colleagues demonstrates that acquisition of gatekeeper mutations in FGFR2 and intratumoral heterogeneity drive resistance in patients with FGFR2-translocated intrahepatic cholangiocarcinoma, which will have important implications for management of the disease in clinic. Cancer Discov; 7(3): 248–9. ©2017 AACR.

See related article by Goyal et al., p. 252 (6).

Oncogenic chromosomal rearrangements in the FGFRs present a potential therapeutic target, with translocations in FGFR2 found most frequently in cholangiocarcinoma (1, 2) and translocations in FGFR3 prevalent in bladder cancer and glioblastoma (2). Preclinical models of these fusion proteins have demonstrated sensitivity to FGFR inhibitors, and recent data from early-phase clinical trials substantiated these findings in the clinic (3–5). However, the extent of these translocations as true oncogenic drivers and the potential mechanisms of resistance to therapy have previously been investigated only preclinically. Goyal and colleagues (6) demonstrate that FGFR2 translocations in intrahepatic cholangiocarcinoma are classic oncogene drivers, with response to FGFR inhibition in the clinical follow-up of common gatekeeper mutations in FGFR2 resulting in clinical resistance.

Cholangiocarcinoma has historically been dichotomized by anatomic location into intrahepatic and extrahepatic subtypes. Recent developments in genomic and transcriptomic technologies have revealed fundamental molecular differences between these anatomic subtypes, reflecting distinct etiologies (7). Intrahepatic cholangiocarcinoma, a disease with growing incidence for yet unknown reasons, harbors FGFR2 translocations in up to 16% of cases (1, 2). Fusion proteins generated by genetic translocations generally fuse a variety of protein fragments to the cytoplasmic tail of FGFR2, thereby deleting the C-terminus. The fusion partners usually contain domain inactivation contributed substantially to acquired resistance in at least 2 of the 3 patients, although FGFR2 p.V564F was the
only selected mutation detected in plasma in the patient with the longest duration of response. Overt geographical intratumoral heterogeneity was further demonstrated from multiregion sequencing of a rapid autopsy, with lesion-specific acquisition of distinct resistance mutations. The study suggests that intrahepatic cholangiocarcinoma has substantial intratumoral genetic heterogeneity, which presents a major challenge to targeted therapy for these cancers. Among the lesion-specific mutations identified, acquired genetic loss of PTEN was selected in multiple lesions, confirming prior preclinical observation that silencing PTEN reduced sensitivity to FGFR inhibitors in FGFR2-amplified cancers (10).

The study provides a rationale for developing third-generation inhibitors that target FGFR2V564F and other common mutations to treat resistant tumors or cut off one of the mechanisms to achieved resistance. Goyal and colleagues also profiled currently available first- and second-generation FGFR inhibitors to identify which may be more active against gatekeeper mutations. Here, the authors switched to using TEL–FGFR3 fusions in BaF3 cell lines for inhibitor screening. Although the kinase domains of FGFR2 and FGFR3 are highly similar, the use of FGFR3 as a model is a weakness in this article. Nevertheless, the results are interesting. Polyclonal gatekeeper mutations develop in TEL–FGFR3-expressing cells with acquired resistance at low doses of BGJ398, but at higher doses, only V555M mutation develops (equivalent to V564 in FGFR2). The authors proceeded to evaluate the differential sensitivity of BaF3 cells expressing an FGFR3 resistance mutation to several other FGFR inhibitors, of which the pan-FGFR inhibitor LY2874455 displayed the most activity against cells harboring the TEL–FGFR3V555M mutant, and also against FGFR2V564F. This finding may have direct clinical relevance in terms of treatment sequencing for patients with cholangiocarcinoma in the future.

The contribution of Goyal and colleagues to the field of FGFR2-translocated cholangiocarcinoma is important. Although this article is limited by small sample size, shared gatekeeper mutations were characterized. Identification of similar gatekeeper mutations in EGFR and ALK-positive lung cancer has led to the development of superior second- and third-generation EGFR and ALK tyrosine kinase inhibitors and improved survival for patients with non–small cell lung cancer. It will be important to extend these observations into more patients, and into other pan-FGFR tyrosine kinase inhibitors. Identification of gatekeeper mutations confirms beyond doubt that cholangiocarcinomas are addicted to translocated FGFR2, emphasizing the pressing clinical need to move inhibitors forward to pivotal studies. A more fundamental question that the current study raises is the utility of post-progression biopsies to evaluate mechanisms of resistance in the face of gross tumor heterogeneity. As the sensitivity of circulating DNA sequencing increases, this tool may become the primary method through which the genomic evolution of tumors is evaluated, as the limitations imposed by the spatial boundaries of biopsies may needlessly limit our capacity to fully comprehend the complexity of heterogeneous tumors.

**Disclosure of Potential Conflicts of Interest**

N. Turner reports receiving a commercial research grant from AstraZeneca and is a consultant/advisory board member for AstraZeneca, Lilly, and Novartis. E. Smyth has received honoraria for advisory board participation from Five Prime Therapeutics and for an advisory role from Bristol-Meyers Squibb. No potential conflicts of interest were disclosed by the other author.

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