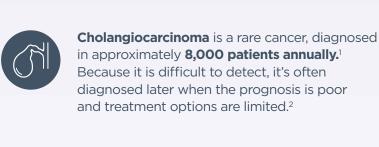
## CDx Announcement

FoundationOne®CDx is now FDA-approved as a companion diagnostic for PEMAZYRE™ (pemigatinib)







FGFR2 fusions or rearrangements occur in

10-16%

of intrahepatic cholangiocarcinoma (iCCA) patients.<sup>3,4,5</sup>





Foundation Medicine testing can help identify *FGFR2* fusions or select rearrangements in cholangiocarcinoma patients who may be eligible for treatment with **PEMAZYRE**™ (pemigatinib).



FoundationOne"CDx is a next-generation sequencing based *in vitro* test intended for use by healthcare professionals for advanced cancer patients with solid tumors. The test analyzes 324 genes as well as genomic signatures including microsatellite instability (MSI) and tumor mutational burden (TMB) and is FDA-approved as a companion diagnostic to identify patients who may benefit from treatment with a specific list of therapies (listed in Table I in the full intended use) in accordance with the approved therapeutic product labeling. Additional genomic findings, other than those listed in Table I, may be reported and are not prescriptive or conclusive for labeled use of any specific therapeutic product. Use of the test does not guarantee a patient will be matched to a treatment or clinical trial option, or that all relevant alterations will be detected. Some patients may require a biopsy. For the complete label, including important risk information, please visit www.foundationmedicine.com/flcdx



## Order FoundationOne®CDx

Web: home.foundationmedicine.com

Email: client.services@foundationmedicine.com

Call: **888.988.3639** 

PEMAZYRE™ is a trademark of Incyte Corporation.

## References

<sup>1</sup>American Cancer Society, cancer.org, accessed 4/27/20

<sup>2</sup> Uhlig J, et al. Ann Surg Oncol. 2019;26:1993-2000.

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<sup>4</sup> Farshidfar F, et al. Cell Rep. 2017;18(11):2780-2794.

<sup>5</sup>Ross JS et al. The Oncologist. 2014;19:235-242.

