

### Highlights from PBC DAY - Analyst Events, February 2021

FRENCH EVENT REPLAY

ENGLISH EVENT REPLAY

### 360° view of PBC

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PBC is a severe chronic, cholestatic, autoimmune liver disease causing injury to the intrahepatic bile ducts, resulting in liver injury and cirrhosis<sup>1</sup>.

#### There is no known cure for PBC<sup>2</sup>

and at present, only two approved treatment options for first line or second line treatment.

Pruritus and fatigue are not addressed by existing therapies<sup>3</sup> and ~40% of patients are non or partial responders to first line therapy<sup>4</sup>, resulting in a highly underserved patient population.

## Patient with PBC Interview

Itch bothers a lot. It really affects you when you go out socially, when you are scratching your face **it's very embarrassing**.

> Dry mouth is pretty bad, especially upon waking up or early in the morning , it's like cotton dry.



Interview conducted by Dr. Kris Kowdley, with one of his patients with PBC, on February 5, 2021

# 3 PBC commercial opportunity



Julien Perrier, Vice President, Global Account Management

Elafibranor is expected to achieve **\$515 million in peak year revenue,** as second line treatment for patients with PBC that cannot benefit from the first line therapy<sup>5</sup>.

#### The PBC market is expected to reach \$1billion in 2025<sup>6</sup>



New therapies required to address the high unmet medical needs in PBC<sup>2</sup> Probability of success of the ongoing Ph3 ELATIVE™: based on positive Ph2 data<sup>7</sup>

Competitive profile of elafibranor, a PPAR α/δ: a promising drug candidate Existing ~\$315MM PBC market<sup>8</sup>, with a double-digit growth, and \$1Bn potential by 2025<sup>6</sup>

### **RELIVE THE EVENT:**

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PRESS RELEASE

1. Kuiper, E.M., et al., Relatively high risk for hepatocellular carcinoma in patients with primary biliary cirrhosis not responding to ursodeoxycholic acid. Eur J Gastroenterol Hepatol, 2010. 22(12): p. 1495-502.; Kumagi, T. and E.J. Heathcote, Primary biliary cirrhosis. Orphanet J Rare Dis, 2008. 3: p. 1.; 2. Hirschfield, G.M.et al, The immunobiology and pathophysiology of primary biliary cirrhosis. Annu Rev Pathol, 2013. 8: p. 303-30.; 3. Lindor et al. Hepatol. 2019; 69 (1): 394-419.; 4. Ali, A., et al., Orphan drugs in development for primary biliary cirrhosis: challenges and progress. 2015. 5: p. 83-97.; 5. IQVIA analysis, Primary research comprised of qualitative interviews with KOLs (28) & payers (15) + quantitative survey with 240 heps.; 6. Iqvia Commercial Opportunity Presentation, 2020 - Resarch on File, November 2019.; 7. Schattenberg et al. J of Hepatology 2021 DOI:https://doi.org/10.1016/j.jhep.2021.01.013.; 8. Intercept Corporate Press Release, November 9, 2020.

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