

Highlights from PBC DAY - Analysts Events, February 2021

FRENCH EVENT REPLAY

ENGLISH EVENT REPLAY

1 360° 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 cirrhosis¹.

There is no known cure for PBC²

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

Pruritus and fatigue are not addressed by existing therapies³ and **~40% of patients are non or partial responders** to first line therapy⁴, resulting in a highly underserved patient population.

2 Patient with PBC Interview

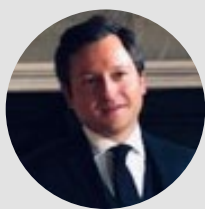
*Itch bothers a lot. It really affects you when you go out socially, when you are scratching your face **it's very embarrassing.***

*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



The total PBC market is estimated to reach **\$1,5bn annually in 2035⁵,** and elafibranor could achieve **\$515 million in peak year revenue,** as potential second line treatment if approved for patients with PBC that cannot benefit from the first line therapy⁶.

4 Conclusion

New therapies required to address the high unmet medical needs in PBC²

Probability of success of the ongoing Ph3 ELATIVE™ trial: based on positive Ph2 data⁷

Competitive profile of elafibranor, a PPAR α/δ: a promising drug candidate

Existing ~\$315MM PBC market⁸, with a double-digit growth, and \$1,5Bn potential by 2035⁵

RELIVE THE EVENT:

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PRESS RELEASE

Note: Elafibranor is an investigational compound and has not been approved by any regulatory authority in any indication. 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 Commercial Opportunity Presentation, 2020 - Resarch on File, November 2019.; 6. IQVIA analysis, Primary research comprised of qualitative interviews with KOLs (28) & payers (15) + quantitative survey with 240 heps.; 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.