

Smart, targeted chemotherapy for advanced liver cancer (HCC)

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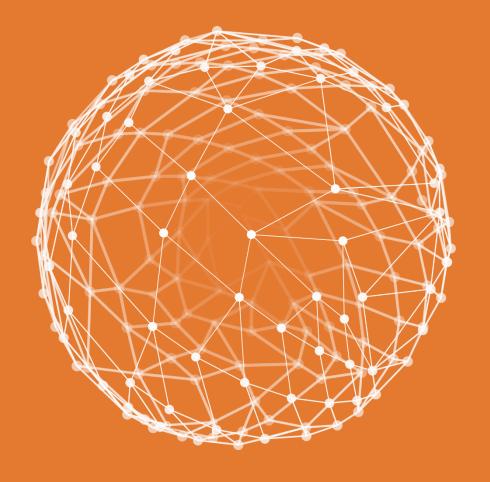
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Medivir is a pharmaceutical company developing innovative drugs with a focus on cancer where the unmet medical needs are high





Fostrox
Selectively killing cancer in the liver



Targeted, smart chemotherapy for liver cancer



Wholly owned phase II asset with strong patent protection

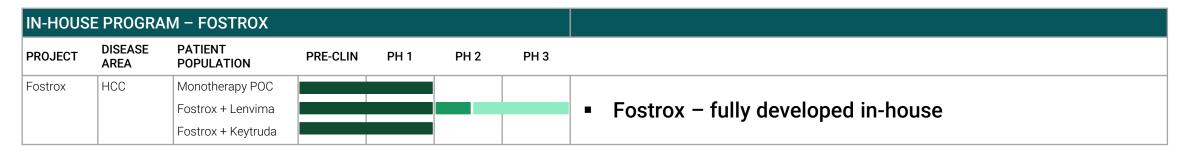


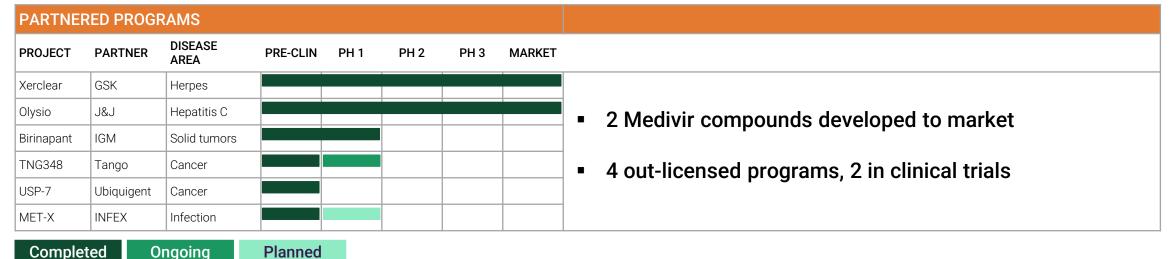
Focus on growing, high-value market segment with large unmet need



Potential for accelerated path to approval

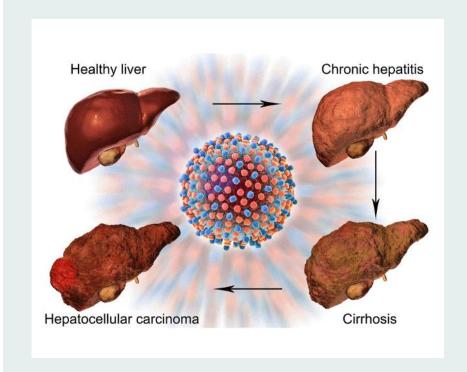
Medivir – experience developing drugs to market with a strong track-record in out-licensing





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Targeted treatment approach critical in liver cancer



- ~80% have underlying liver disease, negatively impacting ability to tolerate systemic anti-cancer treatments^{1,2}
- Tumor growth in HCC is unique as it primarily occurs locally in the liver¹



Only 10% of second line patients respond to current therapies

First line advanced HCC

~1 in 3 responds

- ~90% of patients treated with immunotherapy
- Majority of patients will not respond to current Standard of Care

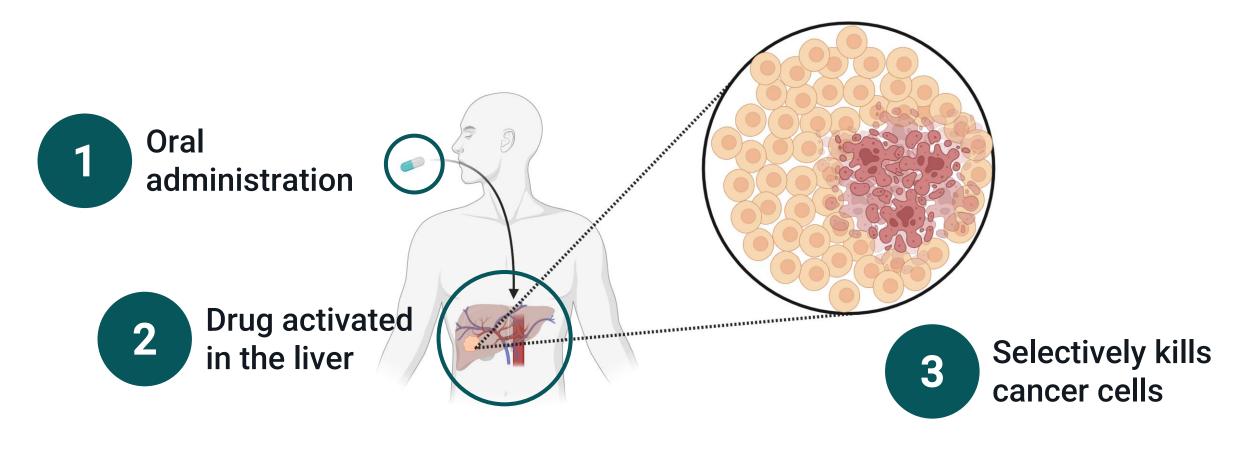
Second line advanced HCC

- No approved treatments after 1st line SoC
- Only ~10% respond to best available therapy¹





Fostrox – a smart chemotherapy targeting cancer in the liver



Global phase 1b/2a study with fostrox + Lenvima (TKI)

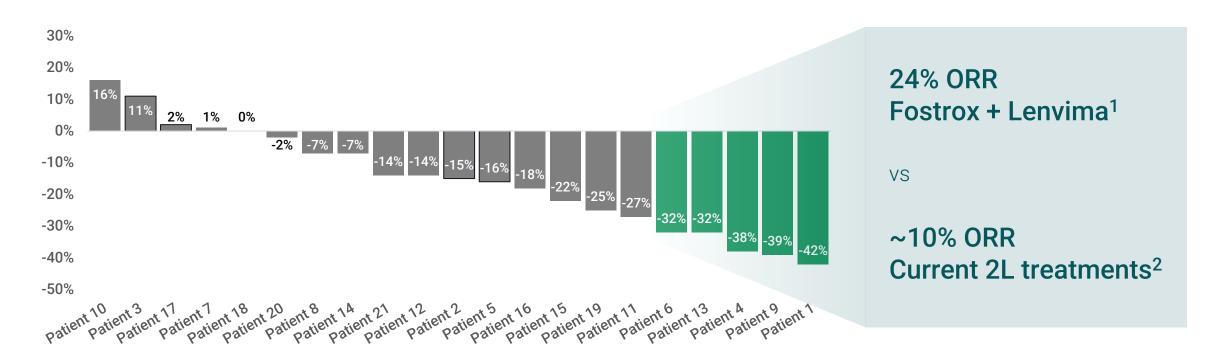


Key study features

- Advanced HCC with generous inclusion criteria, including 2L & 3L patients
- Evaluates potential for synergy between fostrox and Lenvima
- Open-label, single arm, 21 pts
- Final read-out anticipated H2 2024

Significantly higher response rate than current 2L treatments^{1,2}

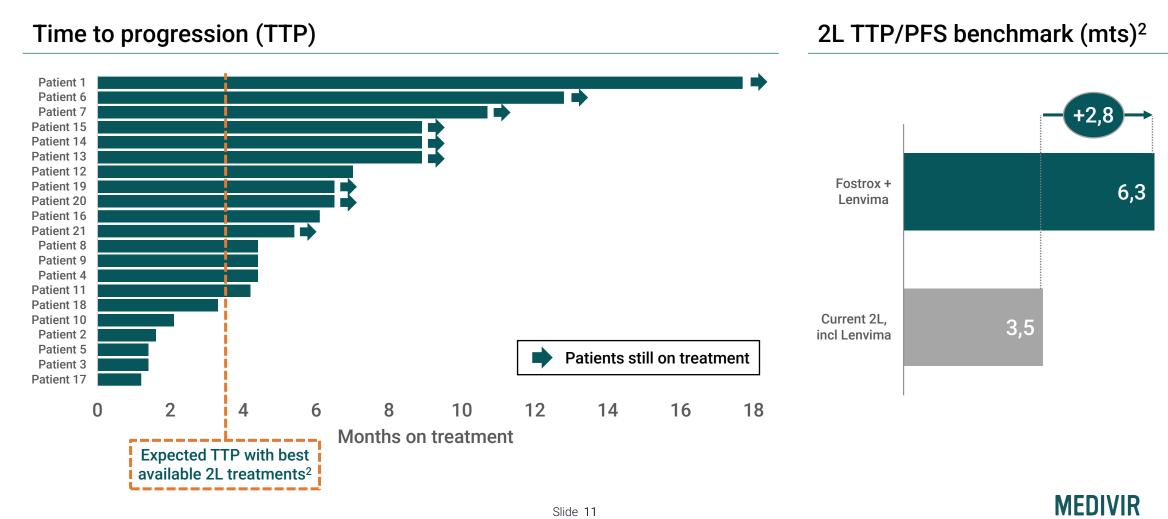
Best percentage change in target lesion size



Patients with partial response

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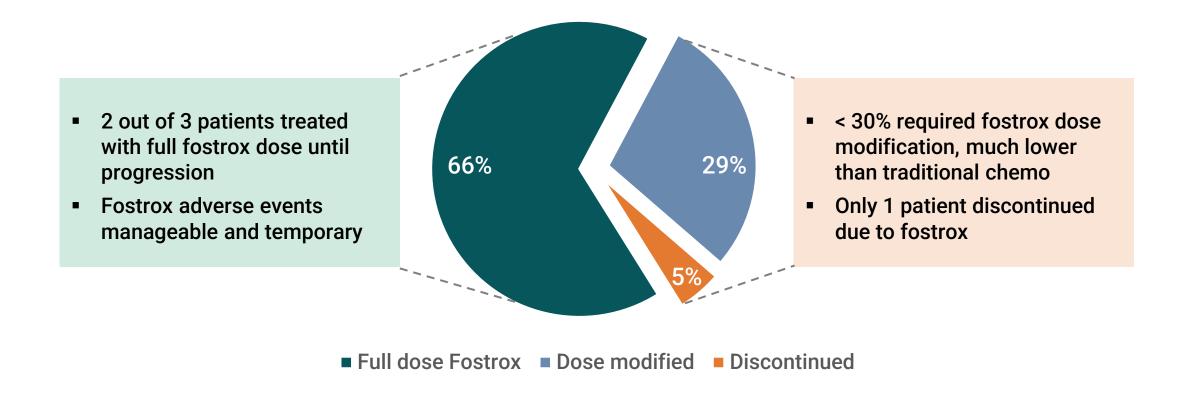
Fostrox extends TTP compared to current 2L treatments¹



¹Local review (All 21 patients data cut-off February 14, 2024), RECIST 1,1

²Based on data from previous 2L phase 3 HCC studies with Stivarga, Cyramza & Cabometyx and investigator initiated prospective & retrospective 2L studies with lenvatinib

A majority of patients tolerate full dose of fostrox long-term¹





Second line HCC market worth over USD 2.5 billion by 2030

Large unmet need in fast growing population

3rd

leading cause of cancer death worldwide¹

+122%

HCC expected to increase +122% in the US and +82% in China² by 2030, caused by fatty liver disease

No

approved treatments in second line post IO-combo

Total market potential > USD 2.5bn by 2030 & growing³

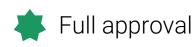




Next step – scaled up global phase 2b to provide opportunity for accelerated approval

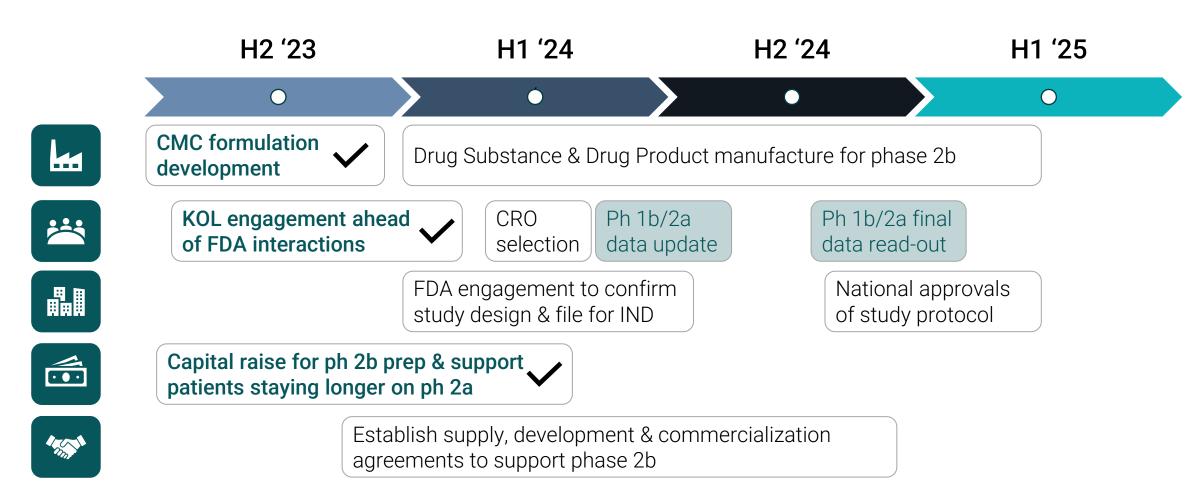
2025 2026 2027 2028 2029 2030 2031 Phase 2b: Phase 3: **Traditional** Randomized but smaller to Randomized with OS as primary endpoint inform phase 3 approach ■ $N \approx 600$ ■ N ≈ 100 Phase 2b: **Confirmatory phase 3: Accelerated** Randomized, scaled up for stats & Randomized with OS as primary endpoint safety database approach ■ N ≈ 600 ■ $N \approx 200-250$ Larger investment upfront





Potential to shorten time-to-market by ~3 years & become first approved treatment in 2L

Fostrox – making good progress preparing for phase 2b





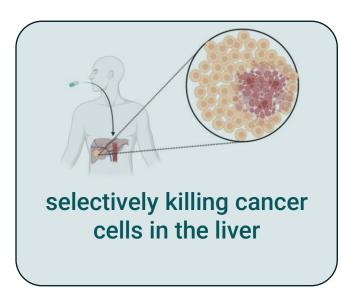


Fostrox – potential to improve second line HCC therapy

Smart chemotherapy

Improving outcomes

First-to-market opportunity



2x

response rate & time to progression

\$2.5bn

market in patients with no approved treatments

Thank You! **MEDIVIR** Slide 17