MEDIVIR Q1 REPORT 2024



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Fostrox + Lenvima efficacy continues to strengthen



Acceleration of fostrox development according to plan



MIV-711 granted RPDD & MIV-701 positive ph. 1 in dogs



Today's presenters



CEO Jens Lindberg



CMO Pia Baumann



CFO Magnus Christensen



CSO Fredrik Öberg



Continued acceleration of fostrox development, strengthened by improved TTP to 7 months

Fostrox – a smart chemotherapy targeting cancer in the liver





Only 10% of second line patients respond to current therapies



~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

~1 in 10 responds



Slide 7

¹Based on data from previous 2L phase 3 HCC studies with Stivarga, Cyramza & Cabometyx and investigator initiated prospective & retrospective 2L studies with lenvatinib

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



Key study features

- Advanced HCC with generous inclusion criteria, including 2L & 3L patients
- Open-label, single arm, 21 pts
- Final read-out anticipated H2 2024
- Next data presentation at ESMO GI, June 26-29, in Munich



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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¹Local review (All 21 patients data cut-off April 8, 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

Slide 9

Fostrox extends TTP compared to current 2L treatments¹

Time to progression (TTP)

2L TTP/PFS benchmark (mts)²



¹Local review (All 21 patients data cut-off April 8, 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



Full dose Fostrox Dose modified Discontinued

Randomized phase 2b; enhanced study design similar in size & timing post FDA feedback

Phase 2b: randomized, double-blind study design



FDA feedback on study design

- ✓ Support for overall phase 2b study design
- ✓ Lenvima rational combination partner & control arm in 2L
- ✓ Two dose arm run in (new fostrox formulation & project OPTIMUS)
- ✓ ORR used in prior accelerated approvals, OS primary endpoint for final approval



Fostrox – well under way preparing for phase 2b



Activity delivered

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



Fostrox – potential to improve second line HCC therapy



Pipeline progress with MIV-711 granted RPDD & MIV-701 positive phase 1 in dogs



Legg-Calvé-Perthes Disease (LCPD) Affects younger children, no approved medicines

Affects younger children

- Incidence varies greatly and seems to increase with latitude
- Highest risk in Caucasian population, studies indicating incidence of ~9/100,000 children <15 years of age in US.
- LCPD usually occurs between 3–12 years of age with the highest rate of occurrence being 5–7 years of age

Pathology clear but not genetic



- Temporary disruption of blood supply to the femoral head
- Pathological changes include osteonecrosis & collapse of the femoral head as well as insufficient normal bone formation
- No single gene with major impact, low concordance in twin studies



Scientific rationale for MIV-711 in LCPD

Cathepsin K inhibition could be key to address LCPD pathology

- Necrosis and inflammation increases levels of cathepsin K which causes bone degradation and femoral head deformity
- Cathepsin K inhibition reduces bone degradation while maintaining osteoclast/osteoblast coupling and ability to stimulate bone formation
- Has shown similar disease-modifying efficacy in OA

LCPD animal model data supports benefit as MIV-711:

- Prevents femoral head deformity
- Improves bone structure
- Does not impact normal bone growth negatively







MIV-711 – Moving forward



Partnership / collaboration approach



Outreach following RPDD and ODD approval



Evaluate additional disease opportunities for cathepsin K



Positive phase 1 for MIV-701 in animal population with high unmet medical need & significant commercial potential

PARTNEF		AMS									
PROJECT	PARTNER	DISEASE AREA	PRE-CLIN	PH 1	PH 2	PH 3	MARKET				
Xerclear	GSK	Herpes									
Olysio	J&J	Hepatitis C						 Positive phase 1 PoC in dogs with periodontal disease (PD) for 			
Birinapant	IGM	Solid tumors						MIV-701, phase 1 in cats ongoing, planned Q3-24 read-out			
TNG348	Tango	Cancer						 80% of dogs & cats >3 years suffer from PD, no disease-modifying treatments approved Vetbiolix estimating annual sales of ~150m USD by 2030 in period 			
USP-7	Ubiquigent	Cancer									
MET-X	INFEX	Infection						oral care market estimated at ~3bn USD			
MIV-701	Vetbiolix	Periodontal									



Financial highlights Q1



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Financial summary Q1, 2024

Consolidated Income Statement, summary	Q	Full year	
(SEK m)	2024	2023	2023
Net turnover	0.5	0.4	7.6
Other operating income	0.1	0.4	1.4
Total income	0.6	0.8	9.0
Other external expenses	-20.7	-13.1	-68.9
Personnel costs	-6.5	-6.2	-27.4
Depreciations and write-downs	-0.7	-0.7	-2.7
Other operating expenses	-0.1	-0.3	-1.4
Operating profit/loss	-27.4	-19.6	-91.4
Net financial items	1.3	0.7	2.1
Profit/loss after financial items	-26.1	-18.9	-89.3
Тах	-	-	-
Net profit/loss for the period	-26.1	-18.9	-89.3

- Net turnover for Q1 was SEK 0.5 million
- Operating loss for Q1 was SEK -27.4 million
- Cash flow from operating activities for Q1 was SEK -35 million
- Cash balance end of Q1 was SEK 153.4 million

Fostrox – potential to improve second line HCC therapy





Fostrox – potential to improve second line HCC therapy



Thank You!

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