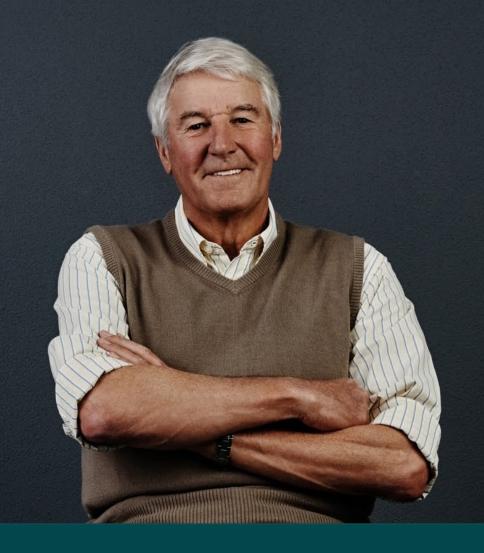
Q4 & YEAREND REPORT 2023

Webcast at 10:00 am CET – February 28, 2024

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Agenda

SCO-101 Dual Mode of Action

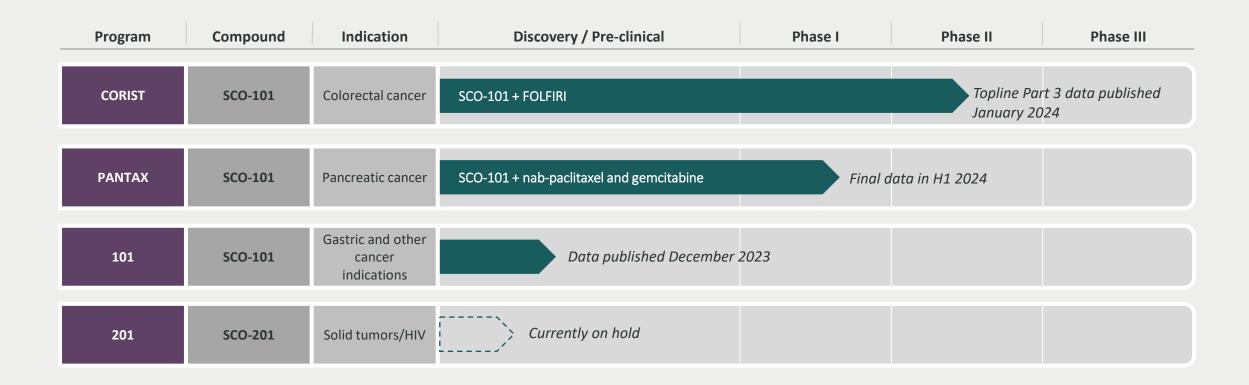
CORIST overview

Financial performance



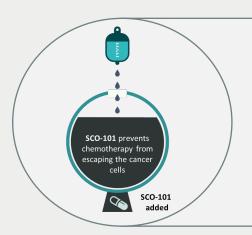


Scandion Oncology has a strong pipeline of innovative assets



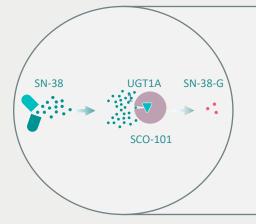


SCO-101 is a dual-acting molecule targeting ABCG2 and UGT1A1



Tumor Effect

Inhibition of ABCG2 prevents the drug efflux pump from removing the chemotherapy from the cell, thereby increasing the chemotherapy concentration in the tumor



Plasma Effect

Inhibition of UGT1A1 prevents the conversion of the active chemotherapeutic metabolite (SN-38) into its inactive form (SN-38-G), resulting in an increased concentration of the chemotherapy in the plasma



CORIST: Lessons learned and Next Steps



Impressive median Overall Survival (OS) of 10.4 months in CORIST part 2

PFS in CORIST part 2 was 2 months

PFS in CORIST part 3 is 4.6 months



Several dosing schedules have been explored in CORIST part 1 and 3 with an intermediate dose in CORIST part 2

Maximal tolerated dose has been established in CORIST 1 and for the 6-day schedule in CORIST part 3

Maximal administrated dose has been reported for CORIST part 3 in the 4-day schedule



Exciting Clinical Benefit Rate (CBR) at week 8

In CORIST part 2 the CBR was 42%

In CORIST part 3 the CBR was 76%



Successful tumor reduction

In CORIST part 2 we saw several patients with a decrease in tumor burden, but no patient had a 30% or more reduction (threshold for a partial remission)

In CORIST part 3 we saw 1 patients with a PR and another patient who after 24 weeks continue to have tumor reduction but not yet at 30%



Next steps: Enabling study to optimize the dose of irinotecan in combination with SCO-101. Goal is to Initiate a randomize proof-of-concept phase IIb study

In CORIST part 3 have one or more additional cohorts in the 4-day schedule, where the dose of irinotecan will be increased to optimize the combination in efficacy but not toxicity

Continue the evaluation to address the observation that bilirubin could be used in the future as a potential biomarker



Financial Overview

- Result in line with expectations
- Strong cost focus and control again in Q4 and throughout the full year

Q4 & Yearend report				
Financial summary	Q4		Full year	
DKK '000	2023	2022	2023	2022
Other income	-48	1,968	446	2,057
Research & development	-6,240	-13,166	-31,851	-65,065
General & administration	-4,868	-4,189	-13,952	-17,158
Financial items	167	-300	653	-2,034
Tax	0	0	5,500	5,500
Net result	-10,989	-15,687	-39,204	-76,700
Earnings per share (DKK)	-0.27	-0.62	-0.96	-1.88
Cash position	26,520	77,605	26,520	77,605

Funding of the Company's activities into 2025



Cash Position

Quarterly cash spend Q4 2023

9.8 MDKK

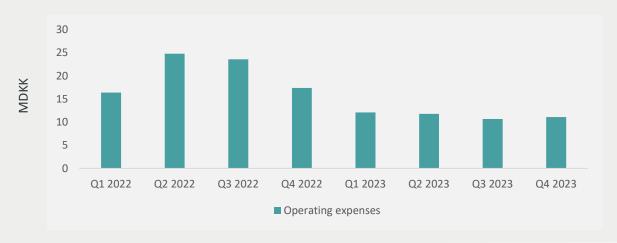
Cash position
September 30, 2023

26.5 MDKK



Financing of the Company's operations into 2025

Operating Expenses 2022 - 2023



Progression of expenses

Strong cost focus and control throughout the year extending cash runway



A series of news flows expected during 2024

2024 H1



CORIST

Part 3 Topline data



Part 3 PK and Safety data

PANTAX

Phase 1b final results

2024 H2



CORIST

Part 3 Final data

Financing secured into **2025**



Scandion Oncology core strengths

One of the first movers in cancer drug resistance

 Potentially first-in-class, targeting a huge market

High medical need in a mature and competitive market

- 10M cancer-related deaths annually
- SCO-101 has broad potential

Financial position

 Current cash funds operations into 2025

Strongly focused pipeline and clinical development

- Focused early-stage pipeline for value creation
- Opportunities to broaden into other cancer indications

Run by seasoned leadership team

- Leadership team with a clear track record
- Best in class Clinical Advisory Board

Multiple value inflection points in 2024

- CORIST part 3 topline results
- Final PANTAX phase 1b results
- Final CORIST part 3 results

