

# CORIST topline results

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**SCANDION**  
ONCOLOGY

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# The Global and European Burden of Cancer

19 million new cancer cases every year in the world



10 million deaths every year in the world



2 million deaths every year in Europe



## Leading causes of cancer death

Lung



1.800.000

Colorectal

916.000

Liver



830.000

Pancreatic

466.000



Colorectal cancer:  
2nd most common cause of cancer death



Pancreatic cancer:  
7th most common cause of cancer death



90% of cancer deaths are due to **resistance** against current treatment options

No drugs are yet available to counteract drug resistance and increase patient survival

# Pipeline

*Developing first-in-class medicines for personalized therapy targeting cancer drug resistance*

Program	Compound	Indication	Discovery / Pre-clinical	Phase I	Phase II	Phase III
CORIST	SCO-101	Colorectal cancer	SCO-101 + FOLFIRI		Part 2: Topline data in Q3, 2022 Part 3: Topline data in Q3, 2023	
PANTAX	SCO-101	Pancreatic cancer	SCO-101 + nab-paclitaxel and gemcitabine		Topline data in H1, 2023	
Immuno-oncology	SCO-101	Multiple cancers				
201	SCO-201	Solid tumors				

The background is a solid teal color. On the left side, there are four white paper airplanes flying upwards and to the right, each following a dashed white line path. On the right side, there is a single yellow paper airplane flying horizontally to the right, also following a dashed white line path. The text 'CORIST topline results' is centered in the middle of the image.

# CORIST topline results

# CORIST Design and Endpoints

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- CORIST is a phase II study, with primary endpoint overall response rate (ORR) according to RECIST 1.1
- As originally designed, it also contains a dose escalation part (part 1) to identify the most suitable dose of SCO-101 in combination with FOLFIRI chemotherapy in a schedule over 7 days. After a suitable dose was found, an expansion of 25 patients (part 2) would assess the efficacy
- Based on part 1 and part 2 pharmacokinetics and pharmacodynamics evidence, the study has been recently amended to explore the potential of a different schedule over 6 days, expected to be possibly the best way to combine SCO-101 and FOLFIRI chemotherapy (part 3 & 4)
- The new part 3 will try to escalate the dose of SCO-101 and FOLFIRI above those employed in part 2
- Part 4 will confirm in up to 24 patients the efficacy of this combination, again based on ORR. The usual secondary endpoints of PFS and OS will also be assessed



# Topline Results of CORIST part 2

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- The dose identified in part 1 was explored in 25 Ras WT patients, all of whom had at least a CT scan after 8 weeks of treatment as per protocol
- The feasibility and safety of combining SCO-101 and FOLFIRI in a schedule over 7 days was confirmed
- Tumor reduction has been observed in some patients, however below the +30% threshold defined as the trial's primary endpoint
- Thus, a proof of concept for efficacy was not reached as no RECIST responses were observed at the planned timepoint of 8 weeks from treatment start
- Evidence of prolonged progression free survival and stable disease (secondary endpoints) was observed in some patients
- Overall, the evidence gathered so far from part 2 of the study, including safety, preliminary activity and pharmacokinetics, support the expansion of the study to investigate a potentially improved modality for combining SCO-101 and FOLFIRI chemotherapy





# Expansion of CORIST (part 3 and 4)

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- The CORIST trial was expanded by adding a new schedule for combining SCO-101 and chemotherapy over 6 days, which will be evaluated in patients with both RAS wild-type (WT) and RAS mutated mCRC
- CORIST part 3 will evaluate the safety and tolerability of SCO-101 in combination with FOLFIRI when dosed according to the different schedule
- CORIST part 3 is planned to include up to 36 mCRC patients with RAS WT and RAS mutated tumors (up to 6 escalation cohorts with a 3+3 design)
- Topline results from CORIST part 3 are expected most likely within Q3, 2023
- In CORIST part 4, up to 24 mCRC patients will be enrolled to assess the preliminary activity of SCO-101 combined with FOLFIRI, administered at the optimal dose found in part 3






# Next Planned Communication

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- In Q1 2023 we will update on the expected timeline of part 3 completion (if based on the number of patients recruited and toxicity observed we are confident the Q3 2023 timeline can be met)
- When CORIST part 3 is completed we will inform about the dose reached with topline results about the safety and tolerability of the new schedule and any activity observed in the part 3 patients
- At this time point there will also be an update about part 2 patients, with a focus on those who are continuing treatment as of today
- Topline results of part 4 will be communicated after all patients have undergone at least the first CT scan on study at 8 weeks, similarly to present communication about part 2
- This may be in the second half of 2023 or first half of 2024, mainly depending on the number of patients recruited in part 3
- The final CORIST study results can be expected approximately 6 months later





**Our vision is to overcome  
cancer drug resistance  
and improve lives for  
cancer patients and their  
families**

**Our aim is to make  
existing cancer  
treatments work  
better and longer**