

Today's presenters



Fredrik Alpsten CEO



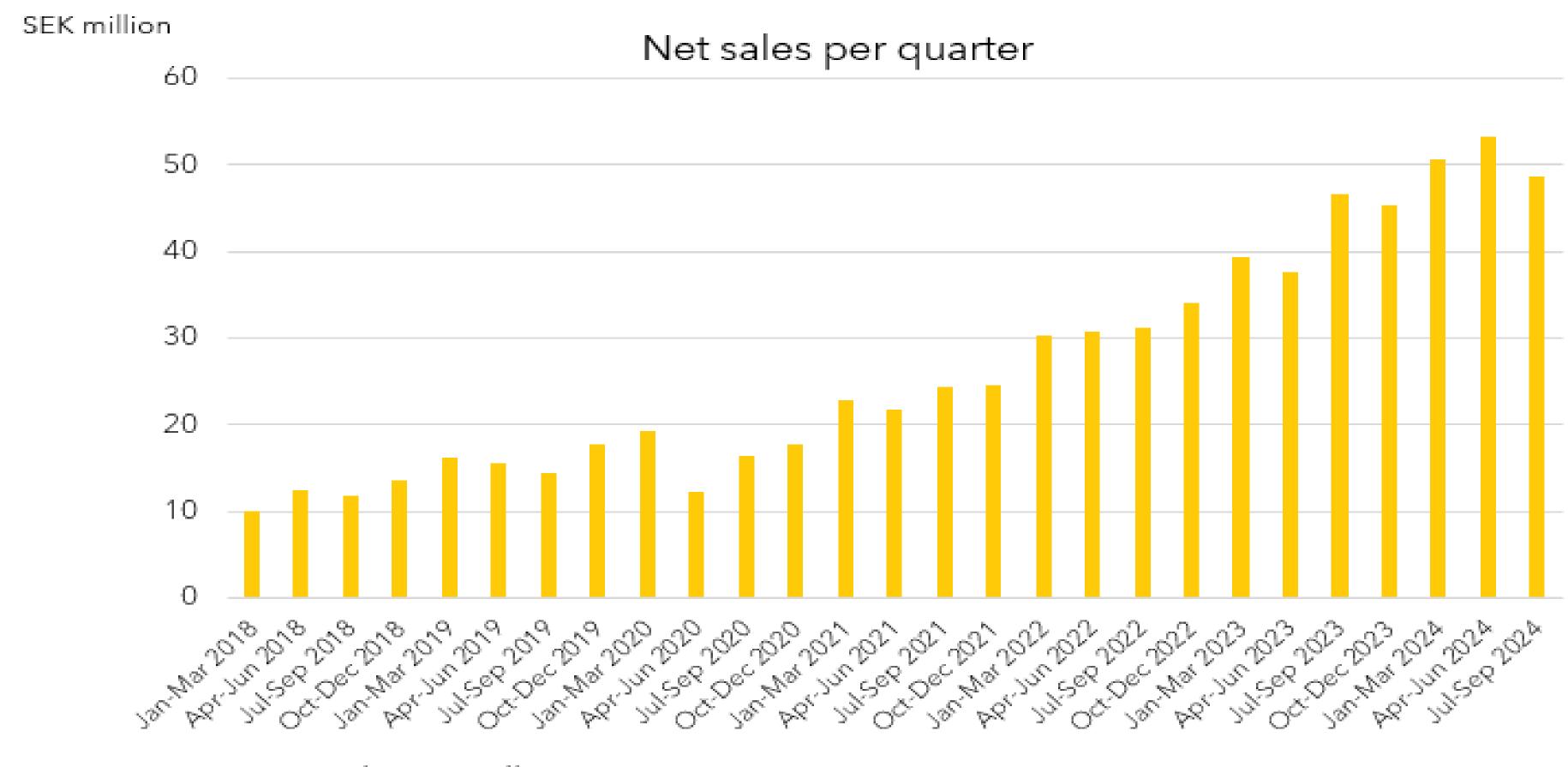
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DVSIR

- Focus on investments for future growth
 - Sales 48.7 MSEK (46.7 MSEK)
 - Growth + 4% (all organic)
 - Growth local currency + 7 %



Quarterly net sales





Sales by region

	Q 3	Q 3		Q1-Q3	Q1-Q3		Q4 2023-	Q4 2022-	
MSEK	2024	2023	Dev %	2024	2023	Dev %	Q3 2024	Q3 2023	Dev %
EMEA	45.2	38.9	16.3	138.0	108.8	26.8	178.6	141.1	26.6
Asia Pacific North & South	2.1	1.6	32.9	5.5	4.3	28.9	7.3	5.1	43.4
America	1.5	6.2	-76.7	9.2	10.8	-14.8	12.2	11.9	2.4
Total	48.7	46.7	4.4	152.7	123.9	23.3	198.1	158.1	25.3



Sales by sales channel

MSEK	Q3 2024	Q3 2023	Dev %		Q1-Q3 2023	Dev %	Q4 2023- Q3 2024	Q4 2022- Q3 2023	
Direct sales	36.7	34.8	5.3	114.6	96.9	18.2	148.5	125.4	18.4
Distributor sales	12.1	11.8	2.0	38.1	27.0	41.3	49.6	32.7	51.9
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 - -23.7 MSEK (-7.8)



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 - -23.7 MSEK (-7.8)
- Strong financial position
 - 171 MSEK in liquidity
 - No debts



DVSIR

Highlights Q3 and Commercial outlook

Update on Thermo Fisher Scientific

- In just a couple of years Devyser has become known and recognised across the transplant community
- Significant research work presented at ASHI
- First data on upcoming 2025 product launches presented at ASHI

"In conclusion, direct comparison of two NGS-based dd-cfDNA quantification methods demonstrated that the new Devyser kit has a slightly lower LOD and overall is a robust assay en par with the established CareDx assay for monitoring dd-cfDNA as an adjunct biomarker in organ transplanted patients"



Seeking FDA clearance with Thermo Fisher Scientific

- Today, patient samples are sent to external testing service providers which means that turnaround times are longer than they need to be
- This also means that those external service providers are collecting the revenues from the significant current reimbursement i.e. 2800USD per test
- Enabling hospitals to take back testing into their laboratory will improve turnaround times and not least enable hospitals to benefit from this significant revenue that todays is sent to a few large companies in the industry





Dvysi

North America update

- Devyser Genomic Laboratories in Atlanta is on track with:
 - Getting our first tests reimbursed (Mol Dx registered, Z-codes assigned, working towards establishing coverage)
 - Our engagement with Cyted follows plan
- **Cystic Fibrosis** is a key focus of ours in the the US market. With the IVD agreement announced with Illumina we now have the ability to seek FDA approval also for this product.
- RHD testing looks highly promising and could be a gamechanger for Devyser in the US. ~15% of americans are rh negative / 3.7m births annually means 550k patients to be tested yearly and we have made significant in roads to this market

China

- On October 27th we received the news of our Devyser Compact product being regulatory approved by the China NMDA
- Since the strengthening of regulatory restrictions a few years back related to market approval our business had been quite slow - but we're now expecting this to resume
- China is a very large market with a lot of potential for several of Devysers products and this approval will accelerate the Devyser expansion into the region of Asia Pacific

Rest of world update

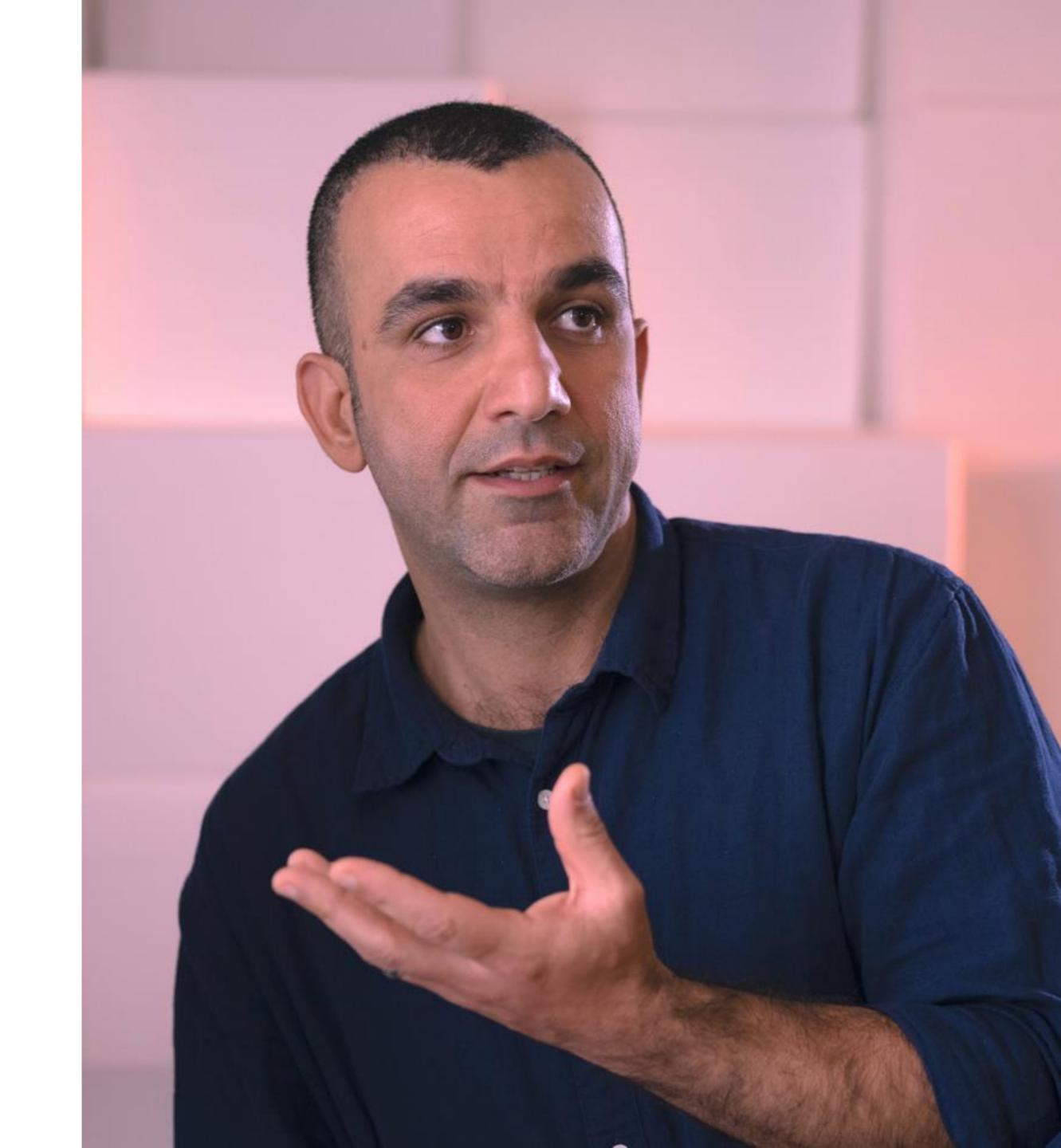
- Both our direct and indirect business is following plan in both Q3 and YTD.
- We continue to expand market share across segments and markets
- Even in Italy where we for several of our products already are the clear market leader we are delivering double digit growth YTD and won additional tenders during Q3
- Our funnel is very healthy and we're confident around the growth outlook

Going forward

- Focus on North America and Europe
- Optimistic about sales development
- Continue our way to profitability

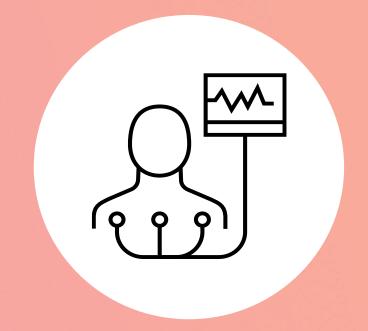


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Our diagnostic areas

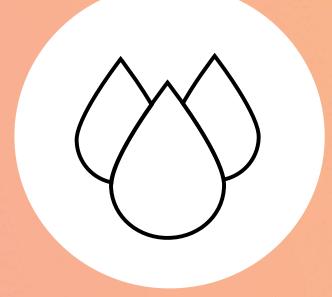


Posttransplantation



Oncology

HEREDITARY DISEASES



Hematology



Reproductive Health



Cystic Fibrosis



Cardiovascular Diseases

Product portfolio by disease area

