

DISCLAIMER

In General. This disclaimer applies to this presentation and any oral comments of any person presenting it. This document, taken together with any such oral comments, is referred to herein as the "Presentation". This Presentation has been prepared by DiaSorin S.p.A. ("DiaSorin" or the "Company" and, together with its subsidiary the "Group"). The Presentation is being furnished to you for information purposes only and for use in presentations of the industrial plan of the Group.

No distribution of this Presentation. This Presentation is being furnished to you solely for your information and may not be reproduced, in whole or in part, or redistributed to any other individual or legal entity.

Verbal explanation. This Presentation has to be accompanied by a verbal explanation. A simple reading of this Presentation without the appropriate verbal explanation could give rise to a partial or incorrect understanding.

No offer to purchase or sell securities. The information, statements and opinions contained in this Presentation are for information purposes only and do not constitute a public offer under any applicable legislation or an offer to sell or solicitation of an offer to purchase or subscribe for securities or financial instruments or any advice or recommendation with respect to such securities or other financial instruments.

Rounding. Due to rounding, numbers presented throughout this Presentation may not add up precisely to the totals provided and percentages may not precisely reflect the absolute figures.

Miscellanea. This Presentation has been prepared on a voluntary basis. DiaSorin is therefore not bound to prepare similar presentations in the future, unless where provided by law. Neither the Company nor any member of the Group nor any of its or their respective representatives, directors, employees or agents accept any liability whatsoever in connection with this Presentation or any of its contents or in relation to any loss arising from its use or from any reliance placed upon it.

Piergiorgio Pedron, the manager responsible for the preparation of the company accounting documents for DiaSorin S.p.A., declares that, pursuant to Article 154-bis, paragraph 2, of the Legislative Decree February 24, 1998, no. 58, to the best of his knowledge, the accounting information included in this Presentation correspond to document results, books and accounting records.

FORWARD-LOOKING STATEMENTS

This document contains forward-looking statements that are based on current expectations, estimates, forecasts and projections about the industries in which DiaSorin operates and the beliefs and assumptions of the management of DiaSorin. In addition, the management of DiaSorin may make forward-looking statements orally to analysts, investors, representatives of the media and others. In particular, among other statements, certain statements regarding future financial performance, the achievement of certain targeted metrics at any future date or for any future period, trends in results of operations, margins, costs, return on capital, risk management and competition are forward-looking in nature. These statements may include terms such as "may", "will", "expect", "could", "should", "intend", "estimate", "anticipate", "believe", "remain", "on track", "design", "target", "objective", "goal", "forecast", "projection", "outlook", "prospects", "plan", or similar terms. Forward-looking statements are not guarantees of future performance and are, by their nature, subject to inherent risks, uncertainties and assumptions that are difficult to predict because they relate to events and depend on circumstances that may or may not occur or exist in the future and, as such, undue reliance should not be placed on them.

Forward-looking statements do not take into account any additional effects that may arise from impacts on the global market in which DiaSorin operates and, more generally, on the macroeconomic scenario, also following any eventual governmental measures related to the spread of COVID-19 and any potential delay in the vaccination campaign.

Actual results may differ materially from those expressed in forward-looking statements as a result of a variety of factors, including: the impact of the COVID-19 pandemic, the ability of the Group to create and launch new products successfully; changes in the global financial markets, general economic environment and changes in demand for diagnostic/ healthcare/life sciences products, which is subject to cyclicality; changes in local economic and political conditions, changes in trade policy and the imposition of global and regional tariffs or tariffs targeted to the diagnostic/healthcare/life sciences industry, the enactment of tax reforms or other changes in tax laws and regulations; the Group's ability to offer innovative, attractive products; various types of claims, lawsuits, governmental investigations and other contingencies, including product liability and warranty claims, investigations and lawsuits; material operating expenditures in relation to compliance with health and safety regulations; the including product liability and warranty claims, investigations and lawsuits; material operating expenditures in relation to compliance with health and safety regulations; the including product liability and warranty claims, investigations and lawsuits; material operating expenditures in relations to compliance with health and safety regulations; the including product liability and warranty claims, investigations and lawsuits; material operation in the diagnostic/healthcare/life sciences industry, which may increase due to consolidation; the Group's ability to fund its defined benefit pension plans; the ability to access funding to execute the its business plans and improve its own businesses, financial condition and results of operations; the Group's ability to realize anticipated benefits from joint venture arrangements; disruptions arising from political, social and economic instability; commercial risk due the fact that the Group operates in a market characterized by the presence of large competitors; ris

FORWARD-LOOKING STATEMENTS

Any forward-looking statements contained in this document speak only as of the date of this document and DiaSorin disclaim any obligation to update or revise publicly forward-looking statements. Further information concerning the Group and its business, including factors that could materially affect the Group's financial results, are included in DiaSorin's reports and filings with CONSOB and Borsa Italiana.

No update. The information and opinions in this document is provided to you as of the dates indicated and DiaSorin does not undertake to update the information contained in this document and/or any opinions expressed relating thereto after its presentation, even in the event that the information becomes materially inaccurate, except as otherwise required by applicable laws.

Non-IFRS and Other Performance Measures. This document contains certain items as part of the financial disclosure, which are not defined under IFRS. Accordingly, these items do not have standardized meanings and may not be directly comparable to similarly-titled items adopted by other entities. DiaSorin management has identified a number of "Alternative Performance Indicators" ("APIs"). These APIs (i) are derived from historical results of DiaSorin and are not intended to be indicative of future performance, (ii) are non-IFRS financial measures and, although derived from the financial statements, are unaudited and (iii) are not an alternative to financial measures prepared in accordance with IFRS. The APIs presented herein include EBIT¹, EBITDA², adjusted EBITDA³, Net Financial Position⁴ and Free Cash Flow⁵. These measures are not indicative of historical operating results, nor are they meant to be predictive of future results. These measures are used by the management to monitor the underlying performance of the business and operations. Similarly entitled non-IFRS financial measures reported by other companies may not be calculated in an identical manner, consequently the measures reported in this document may not be consistent with similar measures used by other companies. Therefore, investors should not place undue reliance on this data.

¹ EBIT is defined as the "Operating Result" net of interests and taxes – ² EBITDA is defined as the "Operating Result", gross of amortization and depreciation of intangible and tangible assets. EBITDA is a measure used by the Company to monitor and evaluate the Group's operating performance and is not defined as an accounting measure in IFRS and therefore shall not be considered an alternative measure for assessing the Group's operating result performance. -³ Adjusted EBITDA, excluding extraordinary costs and expenses incurred in the Luminex transaction announced on April 11, 2021 - ⁴ The Net Financial Position is defined as the algebraic sum (positive balance sheet labilities) of cash and cash equivalents and other current financial liabilities and non-current financial liabilities. -⁵ Free Cash Flow is defined as the set of means available to the Company and is equal to cash flows deriving from operating activities net of interest received or paid, and net of investments of fixed assets.



AGENINA



WHO WE ARE

DIASORIN IS A GROUP COMMITTED TO DEVELOPING PRODUCTS

THAT REQUIRE COMPLEX TECHNOLOGIES AND HIGH-RISK RESEARCH.

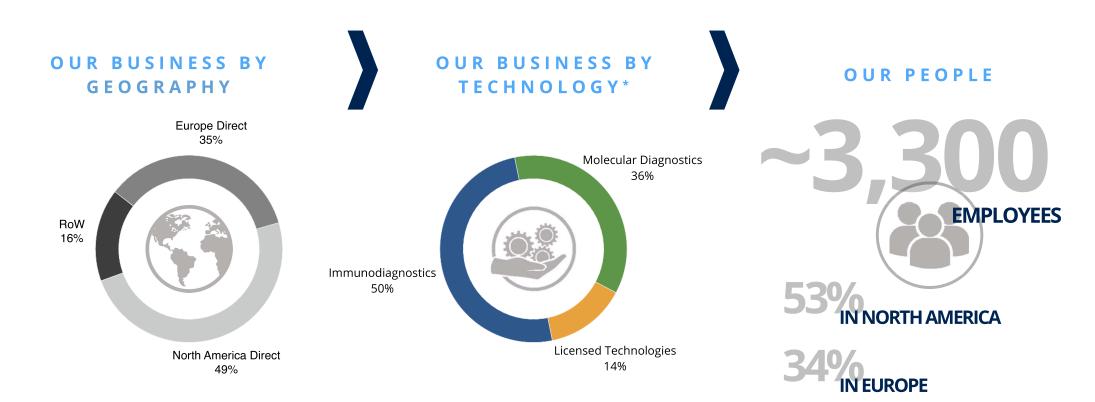
WITH THE ACQUISITION OF LUMINEX, WE ARE NOW A LEADER IN A WIDER

SPECTRUM OF INNOVATIVE TECHNOLOGIES:

IN OTHER WORDS, WE NOW ARE

SPECIALIST³

THE NEW DIASORIN AFTER LUMINEX ACQUISITION



BROADER PRESENCE IN NORTH AMERICA, WITH A STRONGER MOLECULAR DIAGNOSTIC BUSINESS

GROUP OVERVIEW WE WALK THE TALK SETTING

MAIN BUSINESS ACHIEVEMENTS (1/2)

TARGETS

PRODUCTS LAUNCHED

Value Based Care (VBC) initiatives

Launch and increase in the adoption of VBC tests









Immunodiagnostic Tests



New Immunodiagnostic & **Molecular Diagnostic** tests





Molecular Diagnostic Tests



WE WALK THE TALK



MAIN BUSINESS ACHIEVEMENTS (2/2)

TARGETS

PROGRAMS LAUNCHED / UNDER DEVELOPMENT

Decentralization Trend

LIAISON® XS **POC PLATFORMS**



Full-steam launch following approval of key tests









China Manufacturing

OPENING OF NEW PLANT IN CHINA BY 2023

On track to strengthen DiaSorin positioning as a local manufacturer





M&A and Partnerships



ACQUISITION OF KEY TARGET COMPANIES































DECENTRALIZATION VS. CONSOLIDATION TRENDS: A QUICK REMINDER



Need for better performance of routine laboratory services



OPPOSITE FORCES ARE (STILL)
RESHAPING THE LABORATORY
SPACE AND THE IVD INDUSTRY





Need for higher efficiency in patient management



GROUP OVERVIEW

WE WALK THE TALK

THE DECENTRALIZATION TREND ACCELERATED



Roadblocks to performing diagnostic tests for COVID-19 outside of healthcare facilities have been removed.

TESTS CAN BE PERFORMED EVERYWHERE, EVEN IN THE STREET, OR AT HOME.

The importance of Healthcare decentralization leads to additional funding for decentralized hospital locations.

2 examples:

"RECOVERY AND RESILIENCE FACILITY FOR ITALY"

- Funding proximity healthcare services (Mission 6)

€18.5 BN 2021-2026

"U.S. RURAL HOSPITAL FINANCING THROUGH THE AMERICAN RESCUE PLAN ACT"

- Reimbursement for rural healthcare providers

\$8.5 BN 2021

GROUP OVERVIEW WE WALK THE TALK

WHAT DID THE PANDEMIC MEAN FOR IVD COMPANIES: KEY TRENDS



INCREASED PUBLIC AWARENESS

• **Central role of diagnostic testing** had an impact on patients' and governments' perceptions



DECENTRALIZATION OF SPECIALTIES

- Need for COVID diagnostic tools has brought specialty testing to smaller-size labs
- Higher polarization:
 - Importance of high-throughput platforms to manage high volumes of tests (Hub & Spoke model)
 - **PoC technologies** that made *near patient testing* practical and affordable



TECHNOLOGIES/ PLATFORMS

- Widespread use of PCR technology, including in smaller hospital labs
- Lateral flow technology showed its limitation

GROUP OVERVIEW

WE WALK THE TALK

BUSINESS OPPORTUNITY

IMPACT OF THE PANDEMIC IN OUR KEY REGIONS





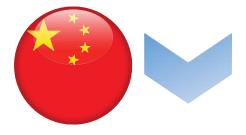
- 1. to increase testing capacity in order to meet demand
- 2. to develop new technologies (e.g. RADx® initiative)
- FDA empowered to bring new assays to the market more efficiently (e.g. MeMed)







 Acknowledgement of the importance of using Multiplexing technology for differential diagnosis (e.g. Respiratory infections)



Accelerated path to independence also in the Healthcare Sector

Strengthening of local players
 (importance of being "China-for-China"
 players) and increased price pressure
 (e.g. Reimbursement Cuts and
 Centralized Procurement)

 No COVID-19 diagnostic test developed by non-Chinese companies has been approved by CFDA

WE WALK THE TA

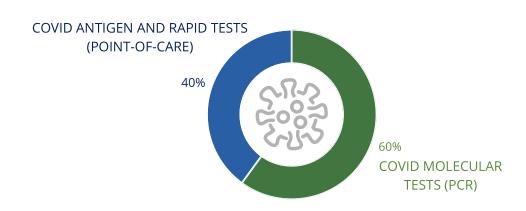
IMPACT ON THE HEALTHCARE SYSTEMS: THE COST OF COVID TESTING

2021E IVD MARKET ~100 \$/BN

+40% VS. PRE-PANDEMIC



2 MAJOR REVENUE CONTRIBUTORS



IN THE LAST 12 MONTHS COVID-RELATED IVD PRODUCTS HAVE GENERATED ~35 \$/BN OF THE OVERALL IVD MARKET TURNOVER



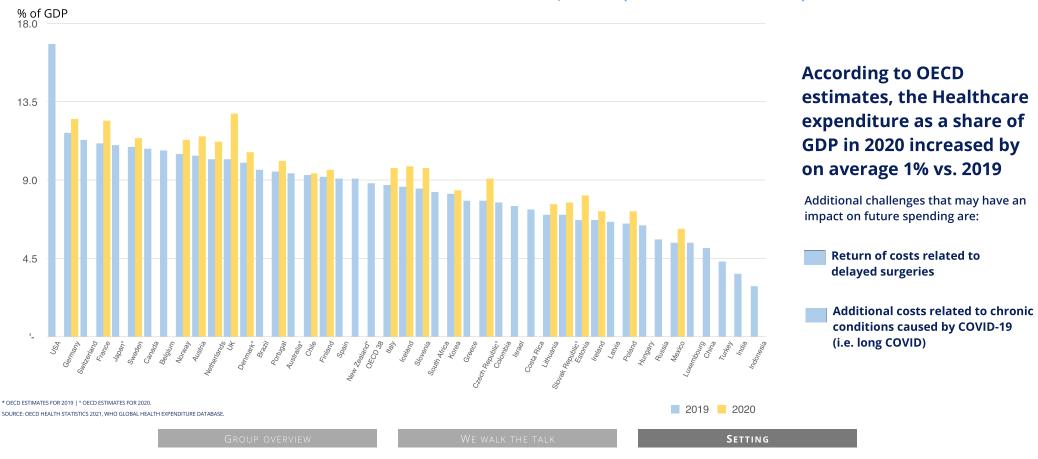
> 150 \$/BN ANNUAL COST FOR HEALTHCARE SYSTEM TO PERFORM COVID TESTS

Group overview

Ne walk the talk

THE COST OF COVID-19 TREATMENT INCLUDING TESTING

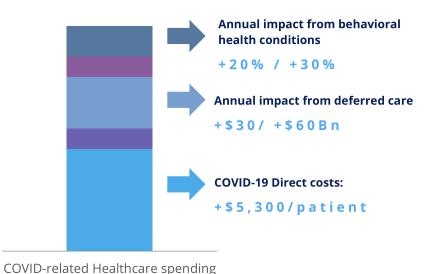
HEALTHCARE EXPENDITURE AS A SHARE OF GDP, 2019 (OR NEAREST YEAR) AND 2020



COVID-19 COSTS IN THE U.S. HEALTHCARE SYSTEM AND IMPACT ON PRIVATE HOSPITALS

ESTIMATED IMPACT ON U.S. HEALTHCARE SYSTEM¹





ESTIMATED IMPACT ON U.S. PRIVATE HOSPITALS²



8.1%

March 2020 probability of default for U.S. Hospitals

\$100 BN

Coronavirus Aid, Relief and Economic Security Act funding for Health Care (HC) providers

\$8.5 BN

Reimbursement for rural HC providers for lost revenue and additional expenses due to COVID-19 as part of 2021 \$1.9 trillion American Rescue Plan Act

WHAT'S NEXT?

GROUP OVERVIEW

WE WALK THE TALK

HOW SHOULD DIAGNOSTIC COMPANIES ADDRESS THESE NEW CHALLENGES

DIAGNOSTIC PLAYERS WILL BE REQUIRED TO SHIFT AND ADAPT THEIR APPROACH IN ORDER
TO ADDRESS THE CHALLENGES AND CHANGES THAT ARE EMERGING WITHIN THE HEALTHCARE SPACE



Platforms with higher throughput/sq.ft. and flexibility to adapt to different testing volume needs









Innovative and reliable platforms that offer near patient solutions with high sensitivity/specificity tests

GROUP OVERVIEW

WE WALK THE TALK



KEY TECHNOLOGIES OVERVIEW

SPECIALIST3

DiaSorin



Luminex



IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LICENSED TECHNOLOGIES

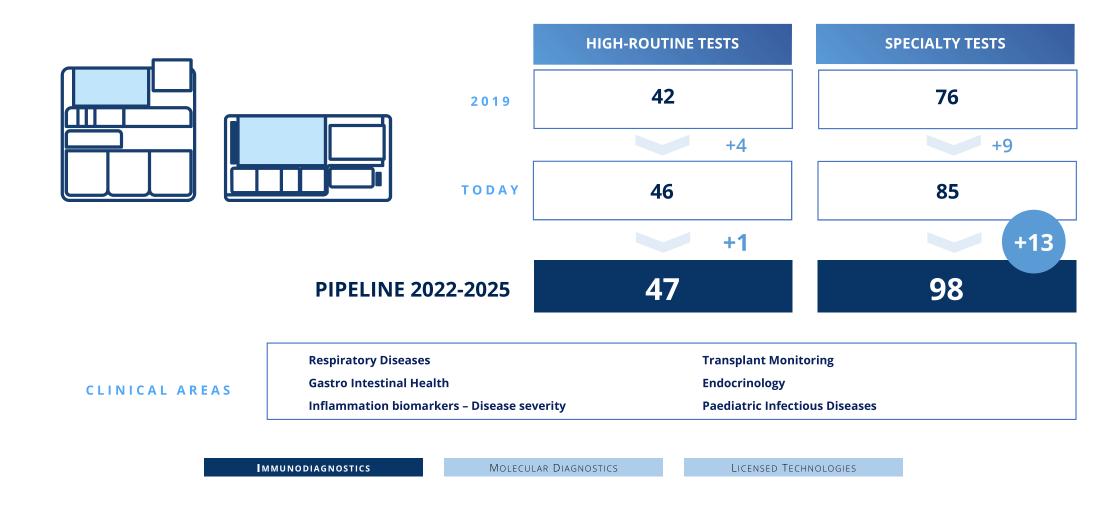


IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LICENSED TECHNOLOGIES

KEEP FOCUSING ON SPECIALTIES







LIAISON® MeMed BV

IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LICENSED TECHNOLOGIES







Headquarters:

Haifa, Israel | Boston, USA

Funding & grants >\$200M

From leading investors, insurers, US DoD and EU Commission

Landmark FDA clearance

The leader in the emerging field of Advanced Host-Response Technologies

The body's immune system is built to communicate what's going on.

Our mission is to listen.

...which leads to some of the biggest healthcare challenges of our time





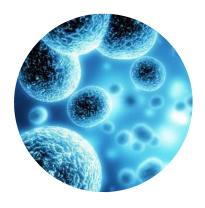
Conventional tests are important but insufficient to effectively aid patient management



Prolonged Time To Results



Inaccessible infection sites



Often, no pathogens are detected



Undetected bacterial co-infections

Confidential

The vision



A few minutes



Small blood volume



Decoding fever and more...

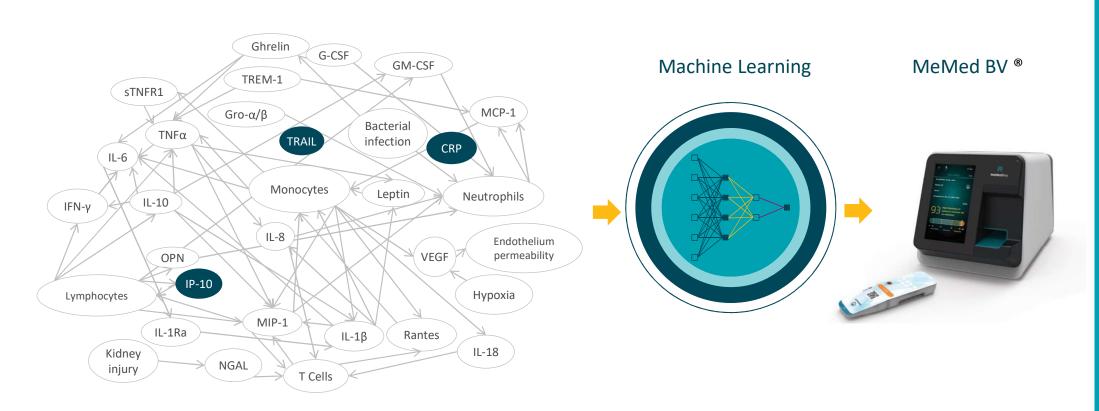


feMed. Proprietary and Confidential Information. This document and its content may not be reproduced, disclosed or passed on to third parties except with the explicit prior consent of

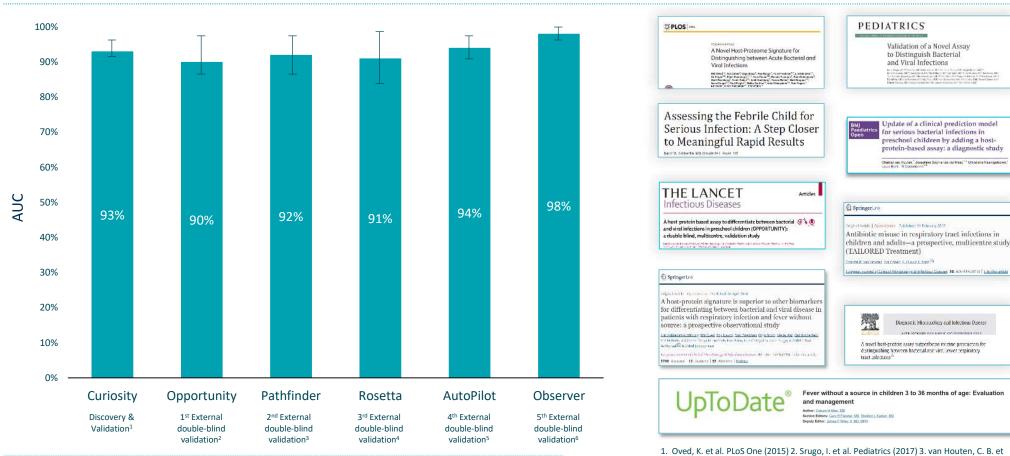
The paradigm: decoding the body's immune response



MeMed BV® uses machine learning to computationally integrate three host proteins (TRAIL, IP-10, CRP) in minutes



BV® high performance independently confirmed in unprecedented blinded validation and real-world evidence of >20,000 patients



MeMed, Proprietary and Confidential Information. This document and its content may not be reproduced, disclosed or passed on to third parties except with the explicit prior consent of MeMed

 Oved, K. et al. PLoS One (2015) 2. Srugo, I. et al. Pediatrics (2017) 3. van Houten, C. B. e al. Opportunity: Lancet Infect. Dis. (2016). 4. The Rosetta study (in preparation) Data on file 5. The AutoPilot study. Data on file; 6. The Observer study. Data on file Landmark FDA Clearance of 1st Technology to Aid in Distinguishing between Bacterial and Viral Infections Using the Body's Immune Response





Indications

Differential diagnosis Bacterial VS Viral infection



Children (>3 months) & adults



Suspected acute infections
(LRTI, FWS, URTI, systemic infections etc.)



end samples collected at hospital admission



Can complement direct viral detection tests by identifying bacterial-viral co-infection

Limitations include immunocompromised patients

MeMed BV[®] is measured on MeMed Key[®] - enabling central lab precision at the point of need.



Diasorin and MeMed partnership - unique complementary capabilities







MeMed BV® on Liaison® receives CE-Mark (Nov 2021)

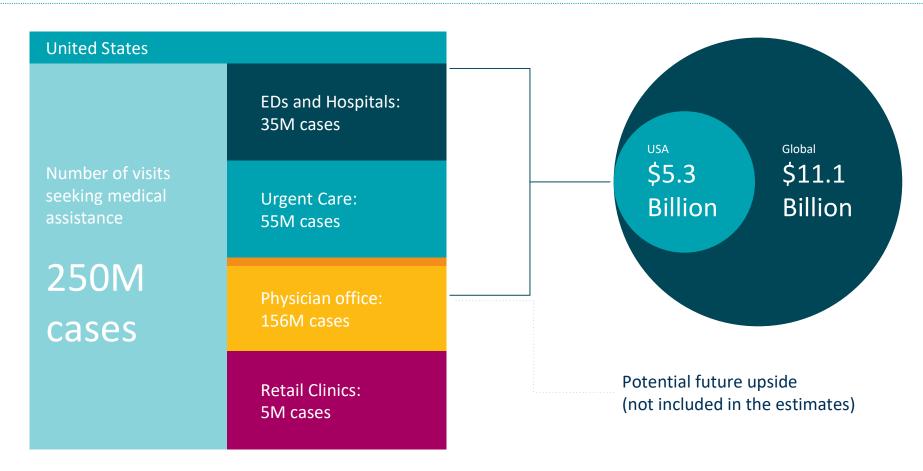


PRESS RELEASE

DIASORIN LAUNCHES LIAISON® MEMED BV®, THE FIRST HIGH THROUGHPUT BLOOD TEST TO DIFFERENTIATE BETWEEN BACTERIAL AND VIRAL INFECTIONS, IN COUNTRIES ACCEPTING THE CE MARK



Targeting >\$11B market opportunity



Cherry et al. 2008; Fagnan 1998; Schappert and Rechtsteiner 2011; Fortune Business Insights; COLA; BCBS; Market Data Forecast; Becker's Hospital Review; UCA 2018 Benchmark Survey; internal analysis

Impacting patient outcome and health economics



1 Source: Schneider et al. Cost Impact Analysis of Novel Host-Response Diagnostic for Patients with Community-Acquired Pneumonia in the Emergency Department, 2021. JME



The immune system is built to communicate what's going on.

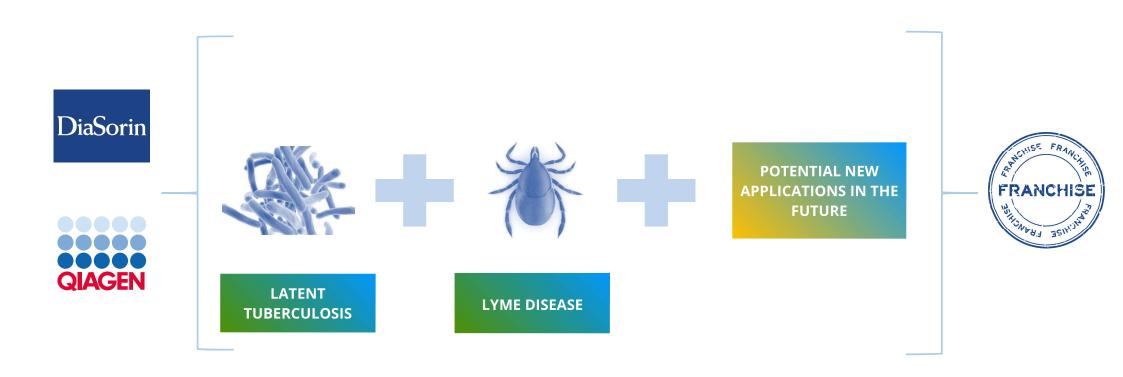








LIAISON® QUANTIFERON® FRANCHISE



IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LATENT TUBERCULOSIS UPDATE



SUCCESS OF THE PROGRAM

PROGRAM ALMOST COMPLETED



> 480 existing customers



PROGRAM IN PROGRESS



> 150 existing customers over an estimated Total Addressable Market of 500+ accounts

4

Hospitals strategy through LIAISON® XS

IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LIAISON® LYMEDETECT®





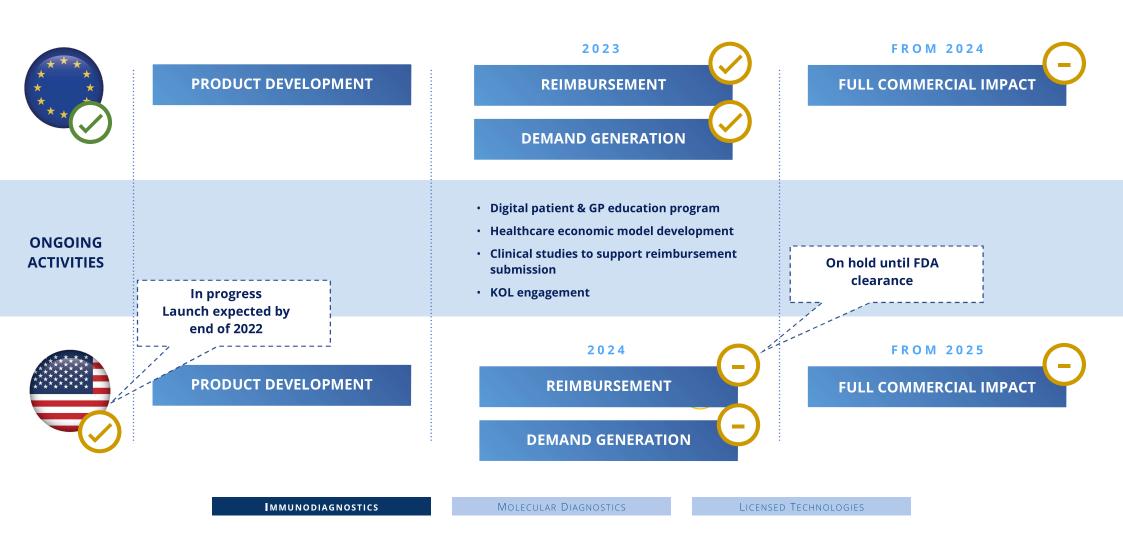
QIAGEN QUANTIFERON TECHNOLOGY AND DIASORIN SEROLOGY TESTING COMBINED TO PROVIDE INCREASED SENSITIVITY IN ACUTE INFECTION DRIVING ANTIBIOTIC THERAPY DECISION



IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LIAISON® LYMEDETECT® PROGRAM STATUS



LIAISON® IQ



Immunodiagnostics Molecular Diagnostics

PHARMACY MARKETS IN USA AND ITALY ARE SIGNIFICANTLY DIFFERENT....







Pharmacy chains dominate the market (5 biggest chains cover one third of the total stores¹)

Mostly independent pharmacies (pharmacy chains are developing)



Regulatory framework is established

• CLIA-waived pharmacies can run IVD (both Molecular and Immuno)

Regulatory framework in evolution. COVID pandemic accelerated the process

- Capillary blood tests (fingerprick) are allowed starting from 2021²
- Swab tests (beyond COVID-19) not allowed yet



Reimbursement for tests allowed in pharmacies operating under a valid CPA³

No reimbursement (out of pocket)



Under a CPA, pharmacists can deliver care functions such as initiating, modifying, or discontinuing drug therapy and ordering and interpreting laboratory tests

Pharmacists can't interpret laboratory tests and can't dispense therapies to patients

IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

... SO IS THE ROLE OF PHARMACISTS IN DIAGNOSTICS DELIVERY

MARKET

~20,000 pharmacies

1/3 in rural setting, where pharmacists have a key role in the community

Growing number of pharmacies (contrary to the rest of Europe)



OPPORTUNITY

Pharmacists can leverage diagnostics to:

- Effectively prescribe OTC supplements to support the monitoring of some chronic conditions.
- Provide first screening on potential diseases to be further investigated by a Medical Doctor



Support OTC supplements / monitoring of chronic conditionsFerritin, Vitamin D, Folate, Vitamin b12, C-reactive protein, ...

Provide first screening layer to address patient doubts on potential pathologies to be further investigated by the MD

Celiac disease, Allergy, D-dimer, ...



IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LIAISON® XS PROGRAM RELOADED



IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LIAISON® XS PROGRAM RELOADED: HOSPITALS STRATEGY IN THE U.S.











TOTAL ADDRESSABLE MARKET ~1,200 HOSPITALS

POTENTIAL PRODUCTS' PIPELINE

★ QFT

Latent TB, Lyme Total, G & M

ANAEMIA

B12, Ferritin, Folate

SEPSIS PCT

★ MEMED BV

★ VITAMIN D

HYPERTENSION

250H + 1,25

Renin, Aldosterone

★ GI PANEL Calpro, Elastase, H.pylori

COVID TrimericS GROWTH нGH, IGF-1

> INFECTIOUS DISEASES EBVs, Toxo, CMV, MMRV, HSV

FERTILITY

Estradiol, Progesterone, Prolactin, Testosterone, LH, HCG, FSH

HEPATITIS & RETROVIRUSES

IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LIAISON® XXL



IMMUNODIAGNOSTICS MOLECULAR DIAGNOSTICS LICENSED TECHNOLOGIES

LIAISON® XXL

CURRENT SITUATION WITH LIAISON® XL

POTENTIAL FUTURE WITH LIAISON® XXL

~70% of existing

DiaSorin installed base is placed stand-alone (single instrument)

LIAISON® XL







LIAISON® XXL (Single - module)









LIAISON® XXL (Double – module)

IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

APPROACH TO CONSOLIDATION AND DECENTRALIZATION TREND





















POINT OF CARE

Expansion in Point-Of-Care

LOW-MEDIUM THROUGHPUT

Migration from LIAISON® & Hospital Strategy

MEDIUM-HIGH THROUGHPUT

Strengthening of existing installed base + acquisition of new customers

VERY HIGH THROUGHPUT

Large Labs requiring more efficient footprint and higher throughput platforms



SMALL-TO-MID VOLUMES

MID-TO-LARGE VOLUMES

MEGA VOLUMES



IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS



TESTING NEEDS SERVED BY DIFFERENT MOLECULAR DIAGNOSTIC TECHNOLOGIES

NEED	SOLUTION
Identify pathogen confirming existing clinical suspicion	Single target determination through single/low plex technology

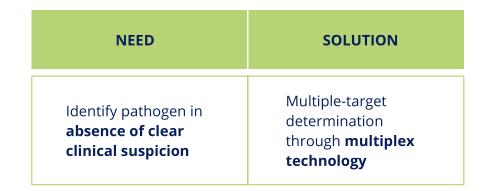
Example: one run to differentiate between COVID-19 and Flu A & B











Example: one run to screen for respiratory pathogens



Up to 4 different pathogens

Up to 40 different pathogens

MARKET TRENDS

SINGLE/LOW PLEX

Mature market (low-single digits growth ex-COVID) with a wide spectrum of platforms (from single to high-throughput)

Clear distinction between high-throughput application (e.g. HIV, HCV, etc.) and specialty testing

Pandemic pushing decentralized testing needs

Pandemic increasing adoption among smaller hospitals, mostly in Europe

MULTIPLEX

Market growing consistently double-digits in the last 5 years

Market driven by reimbursement:

large adoption



resistance

Limited number of panels (# 5)

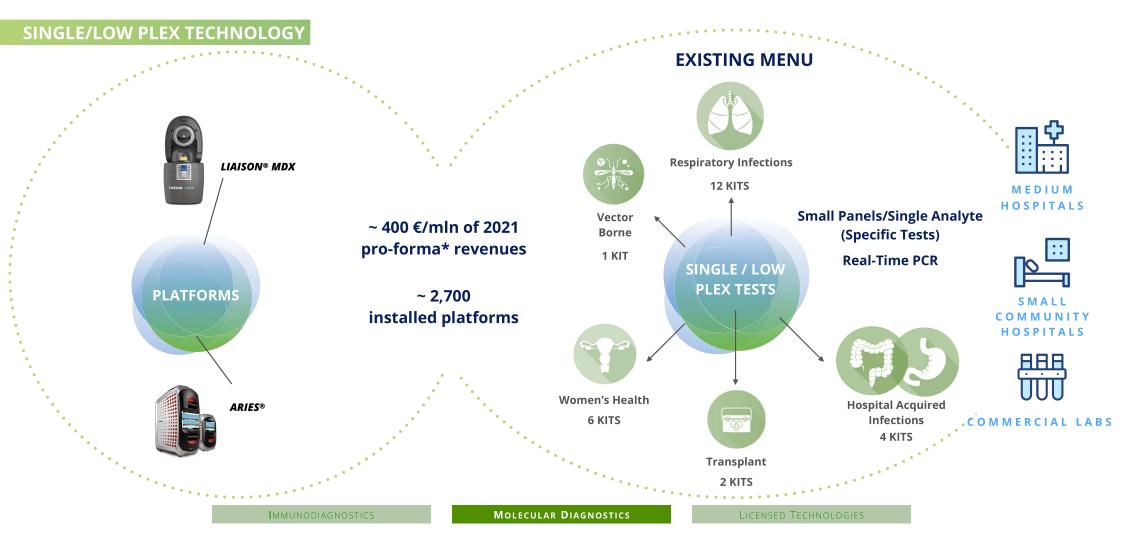
Price and reimbursement pressure

Pandemic pushing adoption among smaller hospitals

IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

MOLECULAR DIAGNOSTICS OFFER



^{* 2021} pro-forma revenues include 12 months of Luminex business contribution

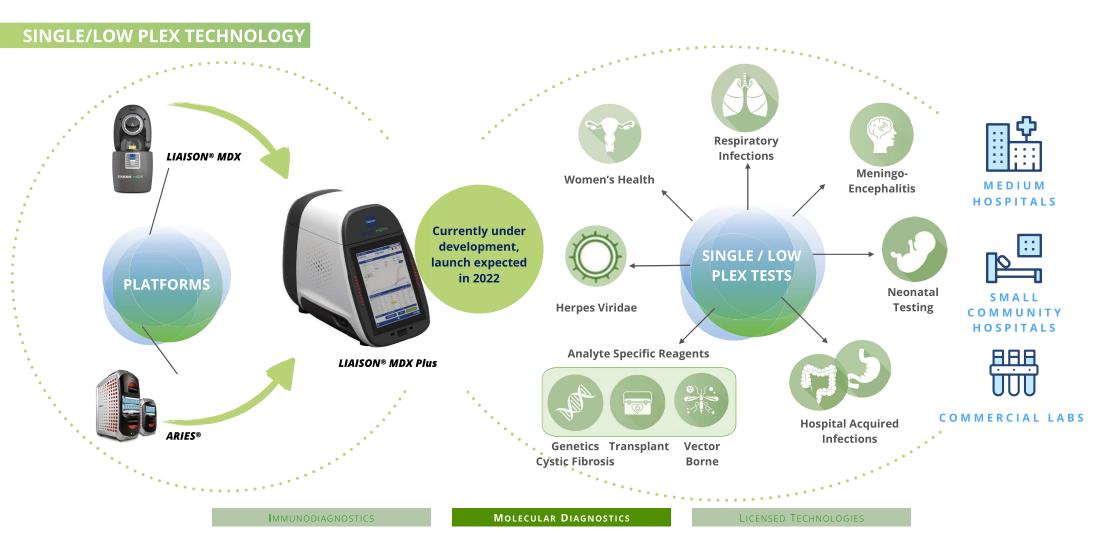
LIAISON® MDX PLUS



| MMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LAUNCH OF LIAISON® MDX PLUS





LIAISON® NES



IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LIAISON® NES

WIDE MENU AVAILABLE ON A RELIABLE,
EASY-TO-USE AND PROFESSIONAL
POINT-OF-CARE PLATFORM, FITTING WITH
THE DECENTRALIZED SETTING'S NEEDS









Technology on par with the laboratory offerings

Immediate result reporting near the patient

Designed for decentralized settings, focus on ease of use

Main Target Clients/Channel







IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

POINT OF CARE BENEFITS











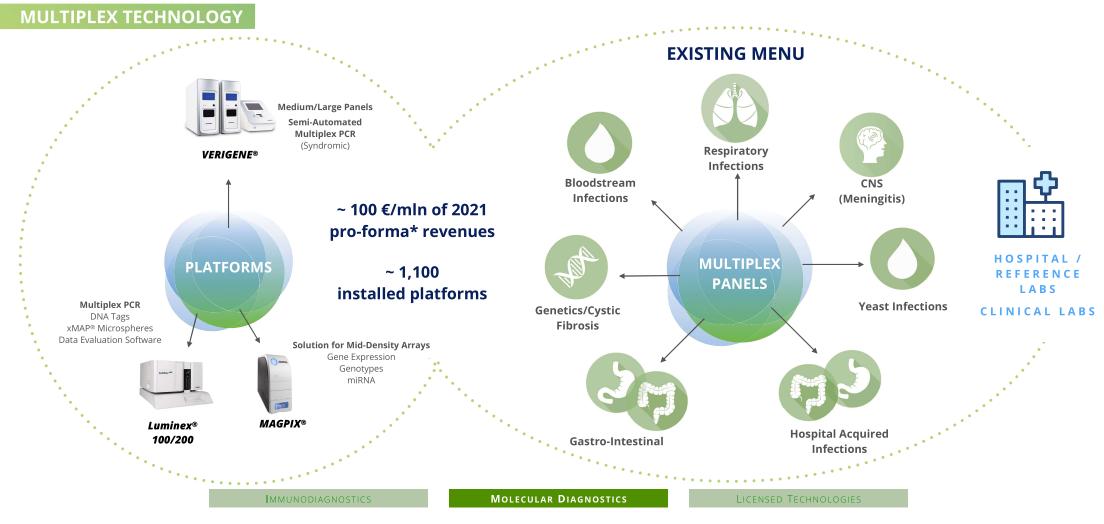
IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS



IMMUNODIAGNOSTICS MOLECULAR DIAGNOSTICS

MOLECULAR DIAGNOSTICS OFFER



^{* 2021} pro-forma revenues include 12 months of Luminex business contribution

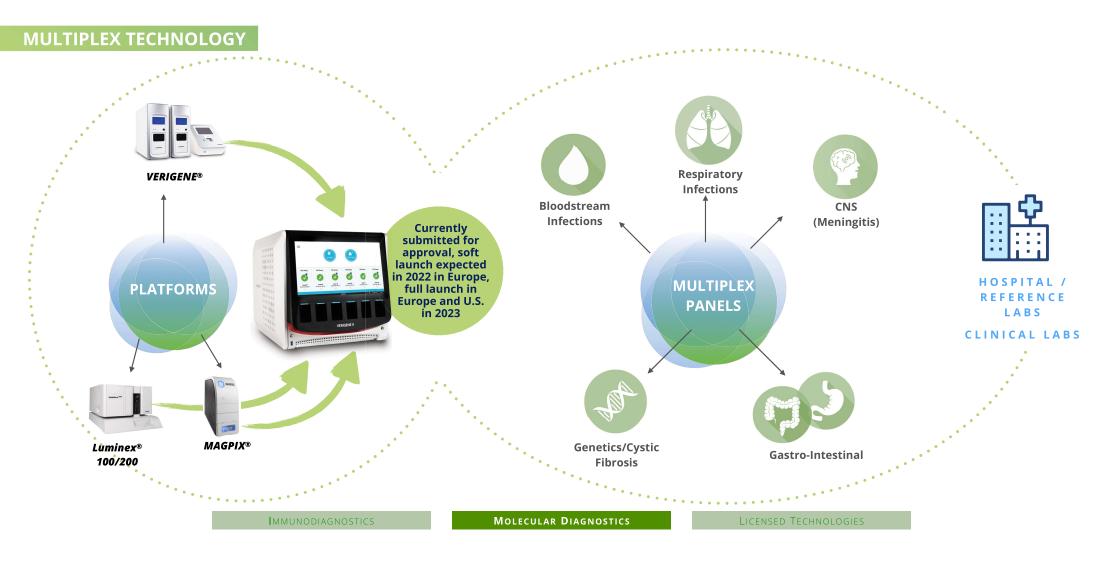
LIAISON® PLEX



IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LAUNCH OF LIAISON® PLEX





FLEX THE PLEX

Key differentiating features

Users choose Full Panel or Flex Testing, unmasking additional results with Flex Credits

Flex customers view fewer targets at a reduced cost, but can unmask additional results as needed using Flex Credits

Customers pay only for tests selected



Flexibility & Scalability

System can scale up for Low, Mid or High-volume settings across Hospitals, IDN Core Labs and Regional Reference Labs



Economic value: Flex testing

Flex enables cost control by:

- fitting into multiple Respiratory Testing algorithms
- empowering user to set up testing panels that meet patient's needs and work within reimbursement landscape



APPROACH TO CONSOLIDATION AND DECENTRALIZATION TREND

LIAISON® NES

LIAJSON° MDX+

LIAJSON° Plex









POINT-OF-CARE

Expansion in Point-of-Care

SINGLE-LOW PLEX

Menu expansion & instrument enhancement

MULTIPLEX

Expansion in Syndromic

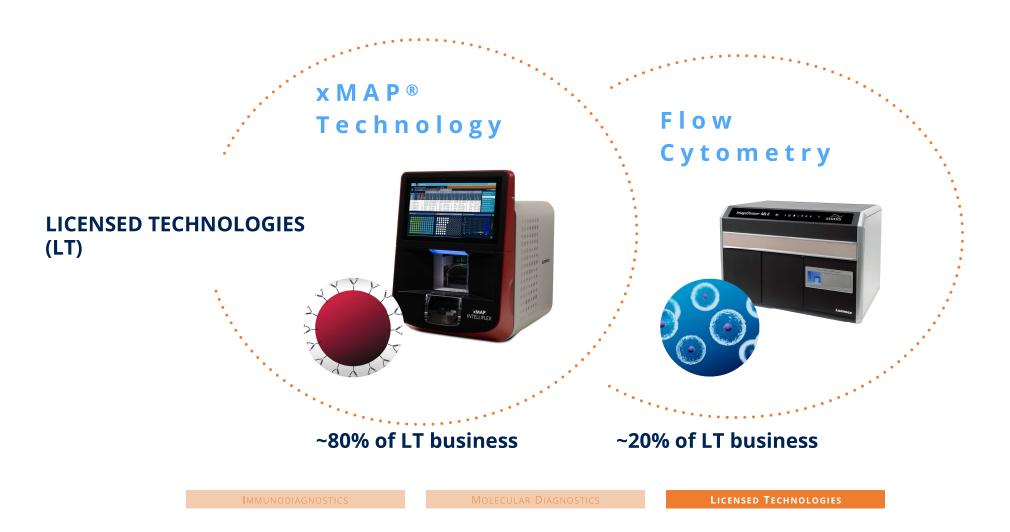


LOW-MID VOLUME



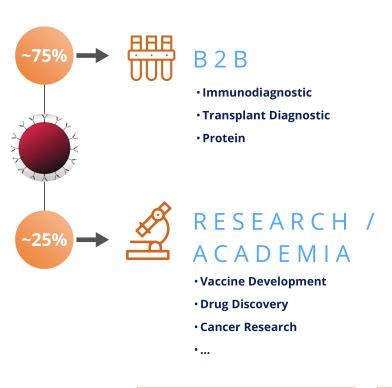


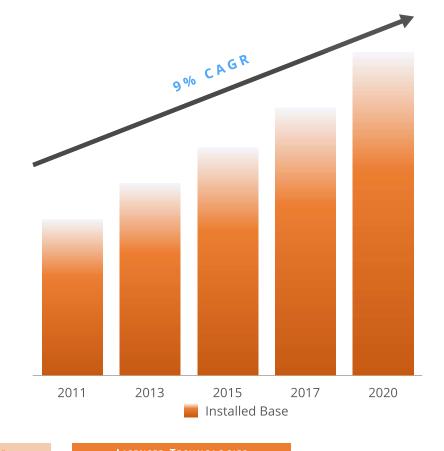
LICENSED TECHNOLOGIES OVERVIEW



LICENSED TECHNOLOGIES: DEEP DIVE ON xMAP® TECHNOLOGY MARKET

25+ YEARS OF EXPERIENCE IN MULTIPLEXING WITH THE LUMINEX XMAP® VERSATILE MICROSPHERES





IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

XMAP® TECHNOLOGY: PARTNERSHIP OPPORTUNITIES



EXISTING LONG-TERM PARTNERSHIPS...

- \cdot 80+ Partners investing in the technology
- Long-term contractual partnerships
- Opportunities in large markets

...AND ADDITIONAL OPPORTUNITIES

New Business Development

XMAP® TECHNOLOGY: INNOVATION AT THE EDGE FOR PARTNERS THROUGH THOUSANDS OF KITS AND CUSTOM ASSAYS

IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS

LICENSED TECHNOLOGIES: THE XMAP INTELLIFLEX® PLATFORM



RELIABLE, SENSITIVE RESULTS

CUSTOMIZATION AND FLEXIBILITY

MAXIMUM DATA/ANSWERS
500 ANALYTES PER WELL; UP TO 2 PARAMETERS PER ANALYTE

NO OTHER MULTIPLEX PLATFORM COMBINES LOW- AND HIGH-PLEX CAPABILITIES, QUICK TIME TO RELIABLE RESULTS, AND THE ABILITY TO ACQUIRE DATA FOR TWO PARAMETERS PER ANALYTE SIMULTANEOUSLY

IMMUNODIAGNOSTICS

MOLECULAR DIAGNOSTICS



MMUNODIAGNOSTICS MOLECULAR DIAGNOSTICS LICENSED TECHNOLOGIES

LICENSED TECHNOLOGIES: DEEP DIVE ON FLOW CYTOMETRY

>10,000 INSTRUMENT PLACEMENTS. THOUSANDS OF PUBLICATIONS ON THE TECHNOLOGY



IMAGE-BASED FLOW CYTOMETRY



BROADEST SPECTRUM OF FLOW CYTOMETRY SOLUTIONS

- **√** Intuitive
- √ Innovative
- **√** Flexible



ImageStreamx® Mk II



Amnis FlowSight®



Amnis® CellStream®

MICROCAPILLARY FLOW CYTOMETRY







Guava® easyCyteTM/ easyCyteTM HT

KEY NEXT OPPORTUNITIES

IMAGING FLOW CYTOMETRY

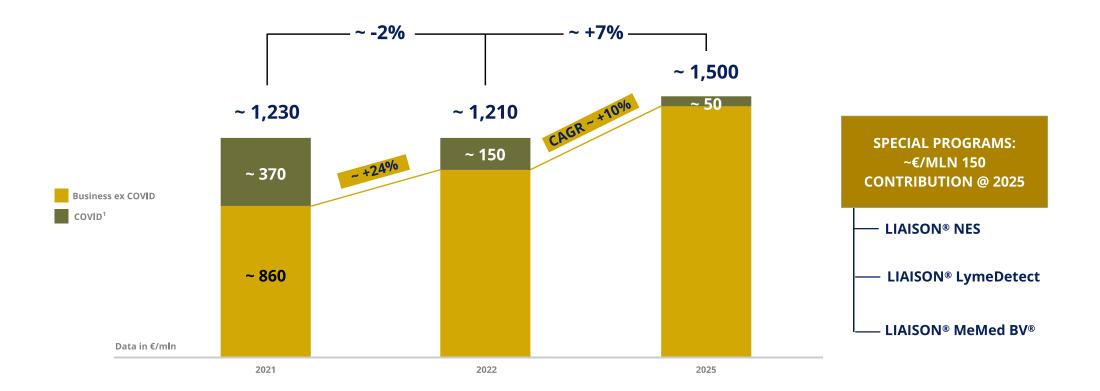
- Launch of the new "ImageStream Next" platform
- "Truth sets" of normal cell images to facilitate detection of abnormalities via existing AI & **Machine Learning technology**
- · New acquisition or partnership opportunities to foster recurring revenue stream
- **Licensing** of technology or instrumentation to third parties outside current fields and customer base

MICROCAPILLARY FLOW CYTOMETRY

- Licensing or Partnership with third parties outside current fields and customer base
- Launch of bacterial cell count & viability **products** to open new markets and foster instrument placement opportunities



2022 OUTLOOK & 2025 GUIDANCE @CER*: TOTAL SALES

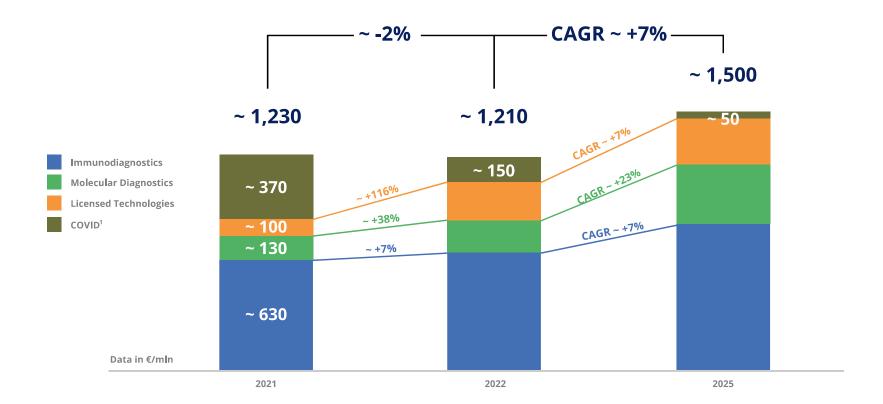


FINANCIAL OUTLOOK

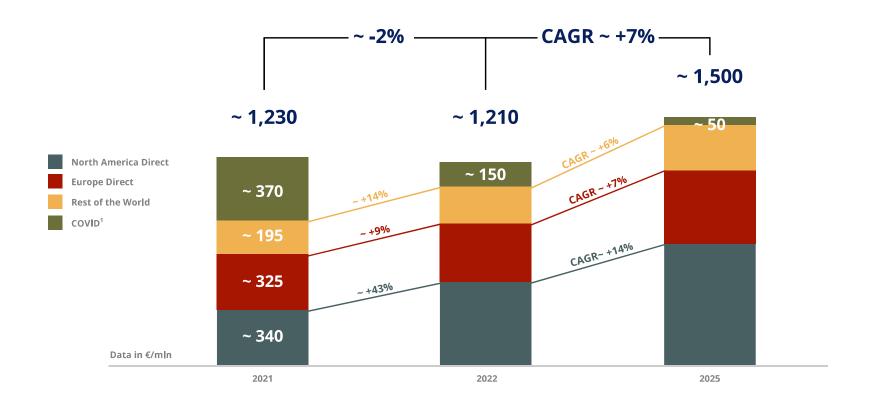
*IN ORDER TO ALLOW COMPARABILITY ACROSS YEARS, ALL FINANCIAL DATA HAS BEEN RESTATED AT CONSTANT EXCHANGE RATE (WITH REGARDS TO THE US DOLLAR 1.16 USD PER EUR)

1. COVID BUSINESS DOES NOT INCLUDE REVENUES FROM MULTIPLEXING RESPIRATORY PANELS THAT ALSO DETECT SARS-COV-2.

2022 OUTLOOK & 2025 GUIDANCE @CER*: SALES BY TECHNOLOGY



2022 OUTLOOK & 2025 GUIDANCE @CER*: SALES BY GEOGRAPHY



REVENUES COMPARISON: 2019 ACTUAL - 2025 GUIDANCE @CER*



GROUP SYNERGIES AFTER LUMINEX ACQUISITION @CER*

MAIN DRIVERS OF COST SYNERGIES:

Platform Consolidation

Geographic Footprint Rationalization

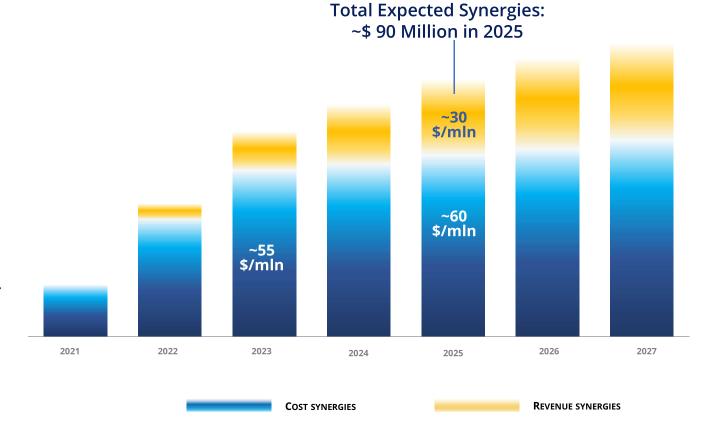
Operations and Supply Chain Optimization

Organization Integration and Right-Sizing

MAIN DRIVERS OF REVENUE SYNERGIES:

U.S. Hospital Cross-Selling

Leveraging DiaSorin Commercial Footprint Outside the U.S.



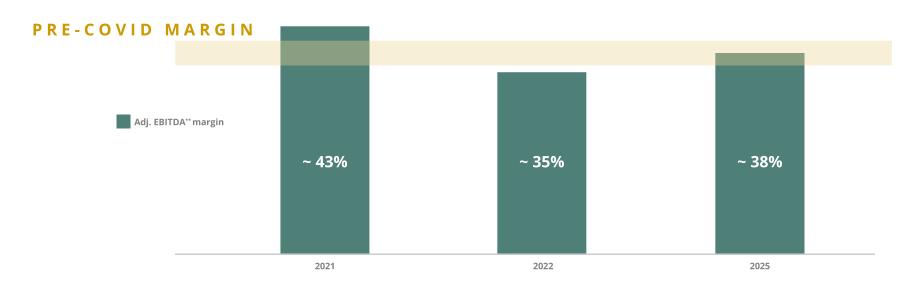
FINANCIAL OUTLOOK

Indicative graphic only, not to scale

2022 OUTLOOK & 2025 GUIDANCE @CER* ADJUSTED EBITDA** MARGIN

- High operating leverage driven by COVID volumes
- · Gross Margin dilution due to product mix
- Reduction of COVID Volumes lower operating leverage
- First full year inclusive of Luminex results
- · Partial synergies' achievement

- Gross Margin improvement partially offset by royalties on partnership products and product mix
- · Full synergies' realization
- Negligible COVID revenues



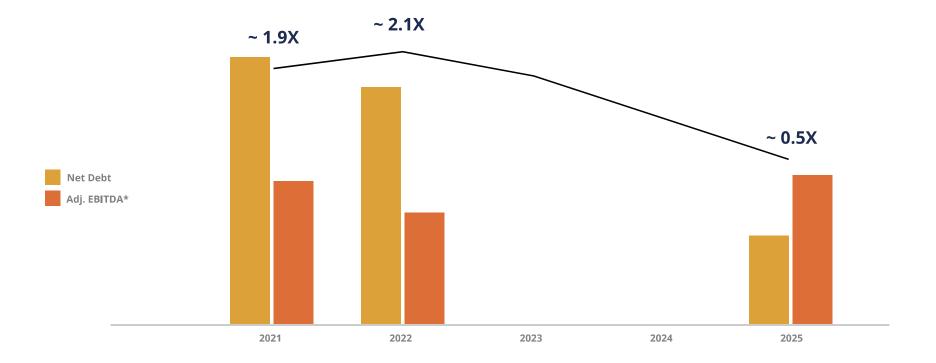
FINANCIAL OUTLOOK

*in order to allow comparability across years, all financial data has been restated at constant exchange rate (with regards to the US Dollar 1.16 USD per EUR)

** Without non-recurring Luminex acquisition and integration costs

Indicative graphic only, not to scale

NET DEBT/ADJUSTED EBITDA* @CER**



FINANCIAL OUTLOOK

**In order to allow comparability across years, all financial data has been restated at constant exchange rate (with regards to the US Dollar 1.16 USD per EUR)

2022 OUTLOOK & 2025 GUIDANCE @CER*

	2021e	2022e	2025 e	Notes
Revenues	~€ 1,230 mln	~€ 1,210 mln	~€ 1,500 mln	22-25 CAGR ~ +7% with Covid ~ +10% ex Covid
Adjusted EBITDA** margin	~43%	~35%	~38%	22-25 Adj. EBITDA** CAGR ~ +10%
Free Cash Flow	~€ 300 mln	Cumulative 22-25: ~€ 1,100 mln		Cumulative 22-25 Capex: ~€ 450 mln
Net Debt / Adjusted EBITDA**	~1.9x	~2.1x	~0.5x	

^{*}In order to allow comparability across years, all financial data has been restated at constant exchange rate (with regards to the US Dollar 1.16 USD per EUR)

^{**} Without non-recurring Luminex acquisition and integration costs