

Disclaimer

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This document contains forward-looking information and statements about GRIFOLS based on current assumptions and forecast made by GRIFOLS management, including pro forma figures, estimates and their underlying assumptions, statements regarding plans, objectives and expectations with respect to capital expenditures, synergies, products and services, and statements regarding future performance. Forward-looking statements are statements that are not historical facts and are generally identified by the words "expected", "potential", "estimates" and similar expressions.

Although GRIFOLS believes that the expectations reflected in such forward-looking statements are reasonable, various known and unknown risks, uncertainties and other factors could lead to material differences between the actual future results, financial situation, development or performance of the Company and the estimates given here. These factors include those discussed in our public reports filed with the Comisión Nacional del Mercado de Valores and the Securities and Exchange Commission, which are accessible to the public. The Company assumes no liability whatsoever to update these forward-looking statements or conform them to future events or developments. Forward-looking statements are not guarantees of future performance. They have not been reviewed by the auditors of GRIFOLS.

Agenda

Time (all in CET)	Topic		
10:30-11:00 am	Breakfast		
11:00-11:05 am	Welcome	Nuria Pascual	
11:05-11:15 am	Introductory Remarks	Raimon Grifols	
11:15-11:50 am	Plasma	Peter Allen	
11:50-12:00 am	Egypt and Canada	Daniel Fleta	
12:00-12:45 am	Biopharma	William Zabel, Joana Sàbat	
12:45 am-1:00 pm	Diagnostic	Antonio Martínez	
1:00-2:15 pm	Lunch		
2:15-2:45 pm	Innovation	Albert Grifols Coma-Cros, César Cerezo, Carter Keller	
2:45-3:30 pm	Biotest	Joana Sàbat, Jörg Schüttrumpf	
3:30-4:00 pm	Financials	Alfredo Arroyo	
4:00-4:15 pm	Final Remarks	Víctor Grifols Deu	
4:15-5:00 pm	Q&A		



GRIFOLS 2022 Investor & Analyst Day

INTRODUCTORY REMARKS

Strengthening Our Mission to Improve Patients' Life Sustainably

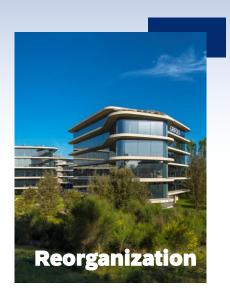
Raimon Grifols, co-CEO

Grifols has supported patients amid unprecedented times through the generosity of its donors and commitment of its 25,000 employees

Reinforcing our mission through a sustainable approach, strengthening...











Expanding and Diversifying Our Plasma Network, Capacities and Capabilities







Expanding and Diversifying Our Plasma Network, Capacities and Capabilities

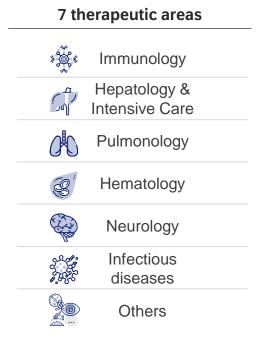
	Today	Progress
Broadening our global footprint of 400+ centers	312 89	+90 centers in 2 years
Strengthening our fractionation capacity	22 ML/year	+7 mL in 2 years

Innovation Ecosystem to Drive Growth and Profitability



Bolstered With the Biotest Acquisition

Innovation ecosystem Internal research New Strategic alliances platforms **GRIFOLS** Investigator Investee -sponsored companies research External research collaborations





3 key projects to drive revenue growth and margin expansion

Fibrinogen

Trimodulin

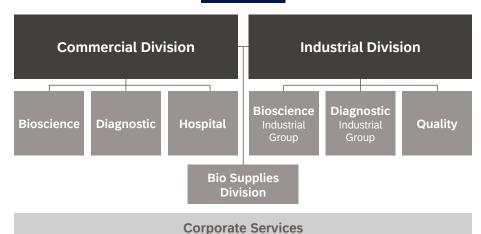
Cytotect®

Streamlining Our Organizational Model...



Previous structure





... To Strengthen the Core

The "New Grifols"

New structure

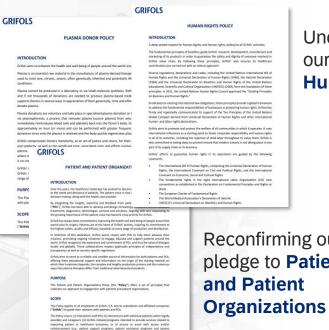


- **Creation of full-fledged Business Units**
- Enhanced effectiveness and operational efficiencies
- Stronger governance model
- **Accountability** over execution
- Less operational complexity
- Greater speed and agility through organizationwide services
- Reduce time-to-market reaction

Operating With a Sustainable Approach

The First in the Industry to Have a Set of 3 Policies Reflecting Our Commitments

Pioneers in developing a Plasma **Donors Policy**



Underscoring our respect for **Human Rights**

Reconfirming our pledge to Patients and Patient

30 Goals for 2030

across the organization

- √ 50% of women in Senior Management positions
- ✓ Achieve 100 training hours/employee per year
- ✓ Increase number of social outreach initiatives and social initiatives investment by 50%
- Donate 240 million international units of clotting factor medicines to support hemophilia patients in developing countries
- ✓ Increase by 10% financial contributions to the Jose Antonio Grifols Lucas Foundation
- ✓ Achieve >60-80% of total spending on suppliers assessed by ESG criteria

Donors and Patients: At the Heart of Our Operations







Social Value

For PATIENTS

€28,600M+

positive impact of 4 main plasma proteins (IG, Albumin, Alpha-1, pd Factor VIII)

For DONORS

€1,800M+

Physical and psychological wellbeing

Financial stability

Educational expenses

Better health

€20M+

dedicated to patient advocacy groups and organizations

For LOCAL COMMUNITIES

€700M+

Note: The Social Return on Investment (SROI) method aims to gain a deeper understanding of an organization's social, environmental and economic impact. The SROI method offers Grifols a valuable cost-benefit analysis. The SROI uses individual assessments to measure the change in stakeholders' lives because of Grifols' activities. The SROI framework is not a financial framework and derives from the social accountancy. Therefore, the framework attempts to put a number on qualitative issues, and it will always show approximate values.



Enhancing Our Operations' Multiplier Effect

Impact on Society 💢 👣 💆 🎏





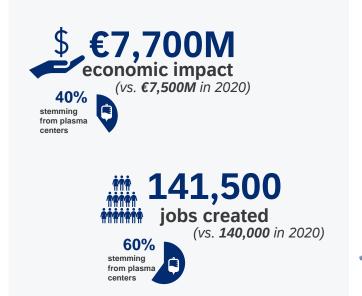




Our Foundations



Socio-economic Impact





Note: Grifols determined the socio-economic impact of its operations in terms of wealth generation and job creation in the United States, Spain, Germany and Ireland. The input-output framework is an accounting instrument that represents all production and distribution operations of an economy in a given timeframe. This model enables observing the different flows of intersectoral transactions in a specific economy in a reference year. In addition, it allows us to observe a series of effects about the production of the system, which appear broken down between the direct or initial, indirect and total effects, which suppose the sum of the previous ones.



Our Team Is in Our Top Priorities

Grifols Talent













Gender equality



60% women

65% promotions are women

41%

senior managers are women

Adjusted Pay Gap

Spain

3.2%1

U.S.

2.1%²

Germany

0.5%

Ireland

0.1%

Talent development



Diversity and inclusion

772 employment of people

+29%

vs. 2020

52%

with disabilities

+11%

aged 30-50 years old

nationalities vs. 2020



¹ Average gender gap for similar jobs in Spain of 40.2%

² Average gender gap for similar jobs in the U.S. of 33.2%

Building On Our Ambitious 2030 Environmental Goals

Environmental Responsibility 👼 🖏





-55% (from -40%) GHG emissions per unit of production



Mobility decarbonization in business travel and commutes



energy efficiency per unit of production



Circular economy continue implementing new measures



100% (from 70%) electricity consumption from renewable energies



Biodiversity protect it in our properties

Recognized Among the World's Most Sustainable Companies















The new Sant Cugat (Barcelona) office building and auditorium awarded **LEED Gold certification** (Leadership in Energy & Environmental Design)

GRIFOLS 2022 Investor & Analyst Day

PLASMA

Leveraging a Unique Combination of Plasma Capacity & Capabilities to Accelerate Performance

Peter Allen, EVP Plasma U.S.

A Unique Combination of Plasma Capacity and Capabilities

Ensuring Sustained Growth and Profitability Through Three Pillars



Expanding global plasma collectionswith the highest
standards of quality
and safety



Leveraging
integrations through
 technology,
standardization and
 efficiencies



Donor & Employee experience excellence



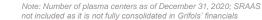
Successfully Executing Our Strategic Plan...

2020



Contract Manufacturing Agreement: Canada

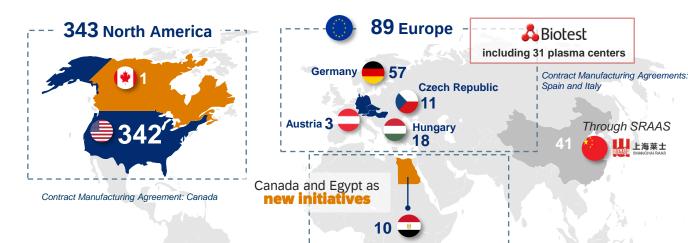




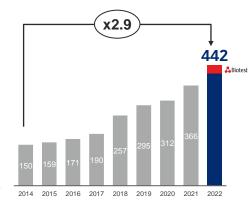


... to Consolidate a Broad Global Footprint of 400+ Centers

By the end of 2022



Expansion of our plasma-center network



Note: Number of plasma centers expected by the end of 2022; SRAAS not included in the 400+ figure as it is not fully consolidated in Grifols' financials

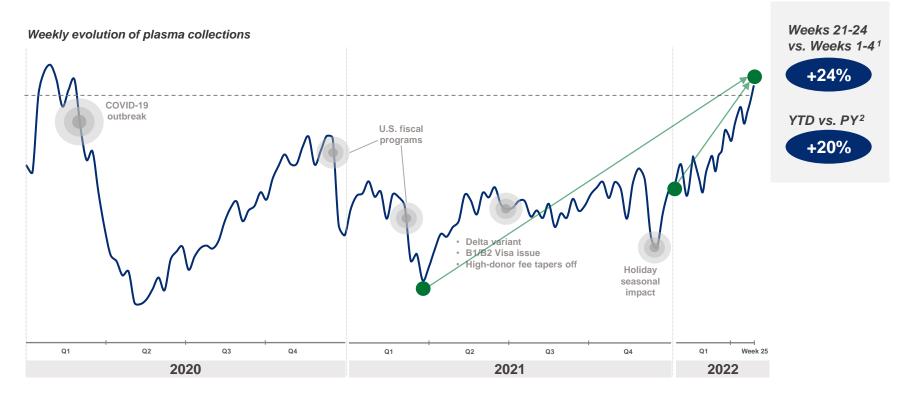
Through SRAAS

Middle East and Africa



Accelerating Plasma Collection Throughout 2022

On Track to Satisfying Growing Plasma Demand



¹ Comparing first 4 weeks of 2022 with last 4 weeks of 2022 (as of week 25)



² Comparing first 25 weeks of 2022 with first 25 weeks of 2021



Accelerating Plasma Collection Throughout 2022

On Track to Satisfying Growing Plasma Demand

• Tailwinds

- Current macro-economic context drives momentum
- Contribution from new and recently acquired centers
- Regular centers return to pre-pandemic weekly levels
- New donor compensation scheme
- Higher talent retention
- New plasmapheresis devices roll-out process to end by year-end – increasing donation yields and efficiencies
- · Digital marketing enhancement
- Both COVID-19 headwinds and U.S. government stimulus impacts are behind us

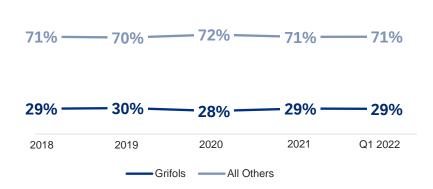
Headwinds

- B1/B2 visa restrictions on U.S southern border continue to impact donation levels.
 Plasma collections upside if these constraints are ultimately lifted
- Donor compensation still high
- Persistence of inflationary and labor pressures



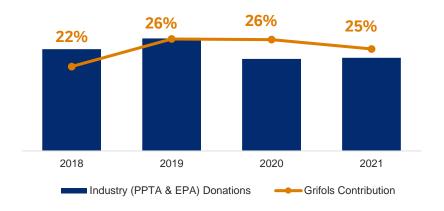
Global Plasma Market Position Supports Leadership in Plasma-Proteins Market Share

Open plasma centers market share evolution (% of open plasma centers)



Note: both graphs including U.S. and EU data

Plasma volumes market share evolution (Industry donations)





Rapid Inorganic Expansion Over the Past 4 Years

A Strong Integration Response Led to Greater Efficiency Gains











35

Q1 2018

Haema. KEDRION Q2 2019

- Operations
- Quality and safety
- Shared services systems
- Donor management applications and communications
- Technologies
- Human resources
- Culture

- Common SOPs1
- Similar technologies
- Standardized policies
- Uniformed practices

- Best practices adoptions
- Common training development and execution
- Leverage on the center of excellence



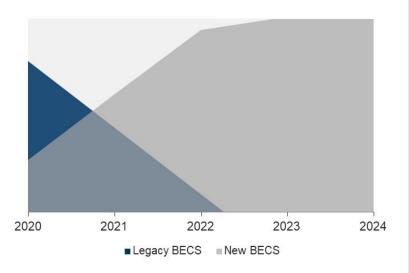


Leveraging Technology to Optimize Ongoing Efficiencies

Increasing Business Process Optimization

Digital Transformation

BECS migration



Benefits

- Improved documentation and workflow
- Automated paperless manufacturing records using handheld computing devices
- Streamlined process flows and reduction of manual verification steps thanks to software automation
- Labor optimization via paperless and process automation
- Donor screening devices that reduce cycle time and allow for automated data entry

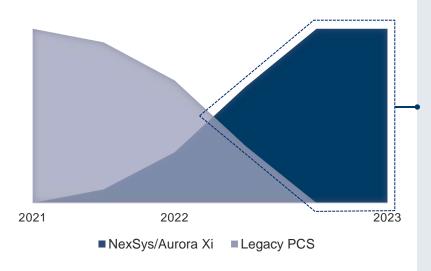


Leveraging Technology to Optimize Ongoing Efficiencies

New Devices Enhance Donor and Employee Experience

Digital Transformation

Device migration



✓ Benefits

- NexSys designed with data management, safety features and usability enhancements to improve yield and efficiency:
 - Integrated touch screen
 - Integrated scanner
 - Integrated to BECS
- Aurora Xi with greater storage capacity; updated algorithms provide faster separation process to decrease donation cycle time and improve yield:
 - More precise controls and settings of device operating capabilities optimize collection volume while efficiently managing donation supplies



Leveraging Our Capabilities

Process Enhancement From Different Approaches

Process efficiencies

- Reduce complexities in the donor center towards standardization of our systems of record and infrastructure
- Better access to donors and potential donors through online appointment and communication platforms
- Rapid donor care management



Donor-center network management

- Diversify the donor demographics to expand the unique donor count
- Focus on donor retention and donation frequency by high service delivery



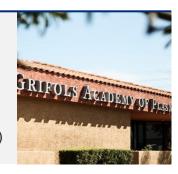
One global plasma organization

- Share expertise and best practices across Grifols plasma organizations
- Implement standardization across plasma organizations preserving local environment
- Orientation to global change and innovation



Talent development

- Academy continues to expand virtual training programs and partnerships
- 350+ graduates from Academy's leadership development program
- 50k+ training hours (all modalities)

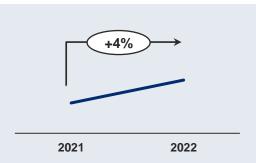




Leveraging Technology to Optimize Efficiencies

Delivering on Improvements Derived From Optimizing Diversification

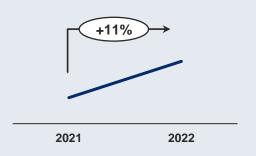
Donor Retention Rate



Technology delivering better targeting for donor recruitment, while enhancing donor experience and loyalty:

- · Legacy BECS transitioned out
- Electronic System **Appointment**
- Donor visibility for upcoming donations

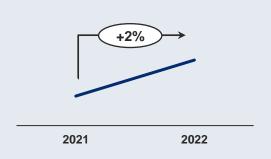
Donations per Employee Hour



Talent development, effectiveness and efficiency initiatives have led to a **higher donations processed per employee** due to:

- Process standardization
- Efficiency gain in donor registration process
- Paperless process

Average Yield



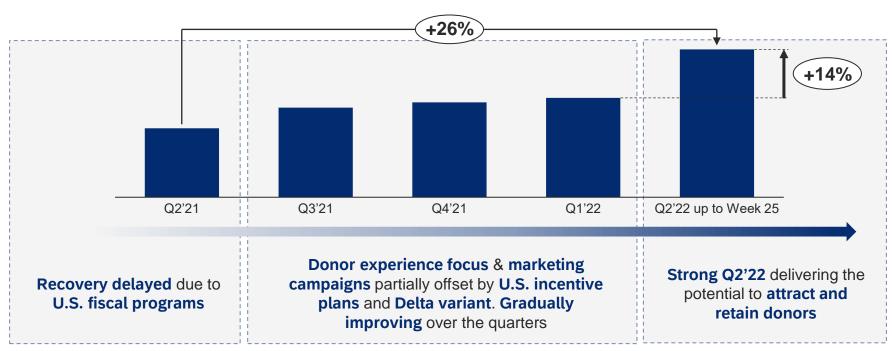
Average liters of plasma per unit are increasing (**higher yield**) as a result of **technology**



Average Liters per Center Demonstrate Positive Momentum

Communication Programs Delivering the Potential

Average Liters per Center*



^{*} Regular centers excluding U.S.-Mexican Border





Employee and Donor Experience Drive Our Mission

Mastering Both Experiences Enables Achieving the Average Liters per Center Target

Employee experience

Substantially improve employee engagement through technology, tools and training

Donor experience

Delight donors at every touchpoint resulting in continual and reliable collections



Higher Donor and Employee Engagement and Retention

Delivering on the employee and donor experience makes **the difference**, what ensures Grifols can withstand competition and market fluctuations

Employee Experience Directly Impacts the Donor Experience

4

Grifols' Employees Proudly Help Patients Around the World

PRE-EMPLOYMENT and ONBOARDING

IN-CENTER EXPERIENCE

EMPLOYMENT TENURE









Reputation for industry recruiting, onboarding, wages and benefits

Strong training programs and a focus on team building, ownership, and work-life balance

Opportunities for advancement, engagement, progressive training and skill enhancement

How does employee experience impact the donor experience?

1

When employees **feel part of a team** and that **they matter** they treat donors in the same way

Well-trained and well-rested employees are more confident and positive and will delivering a more consistent and exceptional donation experience

Long-term employees establish meaningful connections with donors, that stay loyal because they want to continue to see their favorite employees



Success Depends Upon the Donor Experience

Donors Feel Good Knowing Their Plasma Helps Patients Around the World



Multi-channel communications with strong digital presence targeting donor segments

Memorable experiences

Donation centers become a destination

Ongoing dialogue and predictive digital communications inspire donor promotion

Powered by Grifols technology and digital infrastructure

Donors' online activity, behavior and in-center experience are captured and analyzed, providing a 360-degree real-life view about the donor



The Development of Donor Experience

Technology and Data Infrastructure Support Donor Experience

Collect data at every touch

Transactional / Operations data

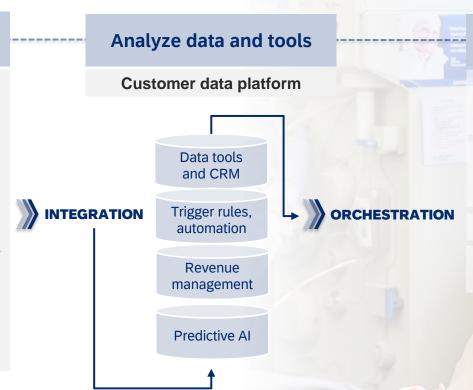
Market, demographic, consumer data

Online & mobile activity data (UX)

Marketing campaign data

Donor research/ segmentation

Customer service & feedback data



Informs and elevates donor experience

Donor direct content (email, SMS, apps, servicing tools, chatbots)

Future donor content (website, social media, search listings, advertising)

In-center experience

Culture of service and employee experience





Building Momentum Through Our U.S. Recruitment Campaigns

Successfully Encouraging People to Donate Plasma

In **2020**, we launched the **industry's first national campaign**, doubling new donor numbers at the height of COVID. Since then, U.S. coverage continues to grow

- ✓ Leveraged donor segmentation data to target attractive future donor audiences
- ✓ Expanded across broadscale channels: digital, video ads, streaming TV and radio, local media and sponsorships
- ✓ Launched lead generation and lead nurturing emails, capturing nearly 250k prospects, with higher conversions, engagement and frequency









Give Your Light

Answer The Call

Show Your Good Side

2020

2021

2022

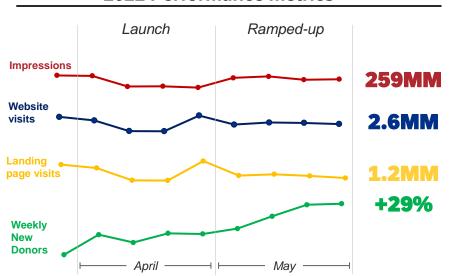
Moving forward...

- Continue to build awareness and engagement with future donors, using uninterrupted recruitment campaigns
- Gained momentum, building on continued learnings to expand and refine recruitment messaging and tactics
- Expand **digital presence** across channels to secure market leadership, reaching and converting future donors to new donors

Building Momentum Through Our U.S. Recruitment Campaigns

Successfully Encouraging People to Donate Plasma

2022 Performance metrics





Omni-Channel Marketing Plan Social Media Programmatic Audio & Video Meredith Partnership (online publications) & NFL Digital NBCU Partnership

✓ Benefits

- Raised broad awareness on the benefits and need of plasma donations
- Recruited new donors and recapture donors who lapsed in plasma donations
- Reinforced messaging of life saving medicines and donor compensation benefits
- Delivered recruitment diversity within targeted demographics

GRIFOLS



In **2021**, we launched the **first German national campaign** to highlight benefits for donors, increasing new donor numbers during the pandemic. The number of new donors has since increased significantly

- ✓ New approach: modern, young and eye-catching
- ✓ Key visuals: show pleasing images or benefits related to donating plasma
- ✓ Target group: young people and individuals facing economic difficulties in pandemic times
- ✓ Multichannel: online, digital, social ads, streaming TV and radio, out-of-home ads, public transport
- √ Key element: landing page with appointment system

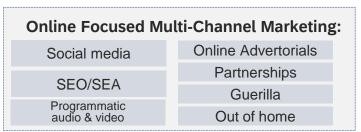


Moving forward...

- Expand digital tools and insights of donors needs and motivation for data-driven marketing campaigns
- Delimitation from competitors and leadership by using donor-experience technologies and services
- Greater awareness of plasma donation is indispensable to produce life-saving medicines



- ✓ Extension of the targeted demographics from young to older
- Expansion of leadership position in social media & online among competitors
- ✓ Development of data-driven marketing along the customer journey
- ✓ Implement a wide range of new technology/systems and upgrade donor app



2021 Performance "Deserve More Than Respect"



Monthly New Donor Performance



2022 Investor & Analyst Day

Leveraging a Unique Combination of Plasma Capacity & Capabilities to Accelerate Performance

Double-digit plasma growth addresses expected plasma needs for 2022 to meet strong underlying demand

Current macro-economic context drives momentum on plasma collections

Successful execution of expansion plan through strong integration response

The reorganization will enable to fully capture benefits of **scale** and **integration**

Reinforcing **Donor and Employee Experience** to gain further efficiencies while ensuring sustainable growth

Continue to build plasma awareness and donor marketing campaigns to support further plasma donations



CANADA AND EGYPT

Responding to the Global Need for a Sustainable Plasma Model

Daniel Fleta, Chief Industrial Officer

Responding to the Global Need for a Sustainable Plasma Model

Game-Changing Initiatives to Become a Blueprint for Other Countries

PLASMA AS A **STRATEGIC ASSET**



"The World Health Organization urges state members to take all the necessary steps to establish, implement and support nationally-coordinated, efficiently-managed and sustainable blood and plasma programs according to the availability of resources, with the aim of achieving self-sufficiency, unless special circumstances preclude it"



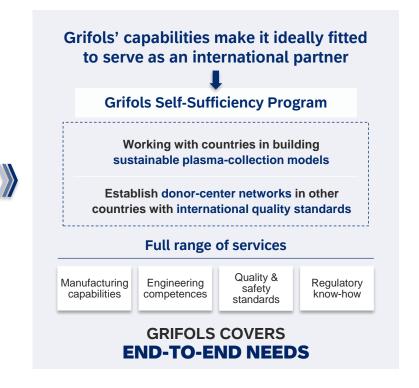
COVID-19 pandemic

Increased emphasis on the importance of plasma supply and its limitations



Growing demand for plasma-derived therapies

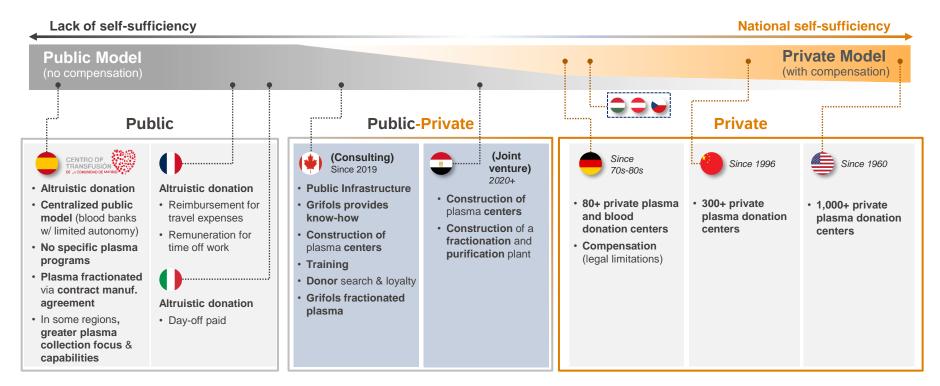
New indications and higherprevalence diseases Diagnosis and prescriptions are increasing



¹ World Health Organization (WHO) (https://www.who.int/publications/i/item/WHA63.12)

Responding to the Global Need for a Sustainable Plasma Model

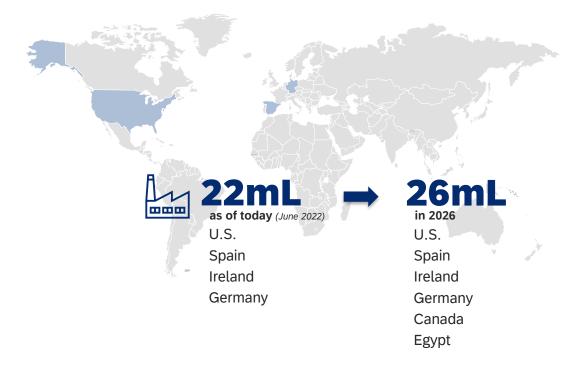
Self-Sufficiency Management Models





Ensuring Sufficient Capacity

Fractionation Capacity Global Footprint





Canada: Leveraging Our Experience and Capabilities...



Well-Positioned to Satisfy the Country's Needs

Canadian plasma-derived therapies market



Grifols is the best positioned company in Canada

2021 IgG Kg / inhabitants



Mature and stable market with high consumption of IgG levels

... but Canadian government seeks to be more self-sufficient

Deep experience in the Canadian market, with more than 30 years as the primary fractionator for Canada's blood operators



Only large-scale domestic manufacturing facility and uniquely positioned for continued growth across full platform of service offerings



Grifols has a **local fractionation**, purification and fill & finish plant

Source: Global Market Database, 2021 preclosing



Grifols Canada Therapeutics (GCT)

Strategic Location





Construction of a New State-Of-The-Art Plant



Canada Site Facts

✓ Acquired by Grifols October 1, 2020

Located in **Technoparc** *Montreal*, Quebec

Plot: **63,134 m**² Built: **19,865 m**²

3-story building

Manufacturing, warehouse & QC lab and admin building



Canada **Current Facility**





Canada **Expansion Project Construction**



+6,600 m²

Phase I August 22

Albumin Purification

Phase II Sept '23

Fractionation Expansion

Phase III Nov '24

Warehouse Expansion

Gamunex Expansion



Albumin Conversion – Phase I, August 2022













Egypt: Delivering On Our Promises Through a Pioneering Model

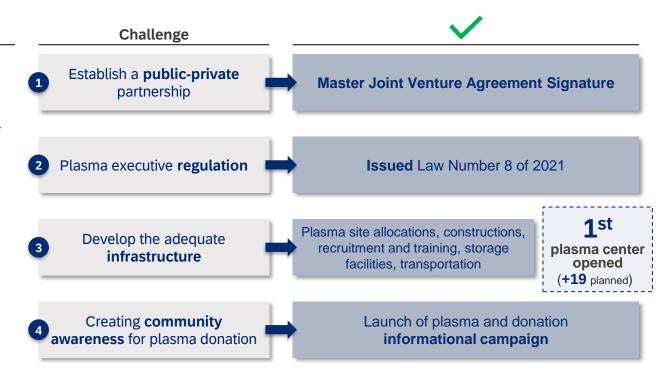
Overcoming Challenges to Strengthen Egypt's Healthcare System

Egyptian plasma-derived therapies market

National mandate to develop the healthcare system

Plasma-derived medicines needed for its **growing population**

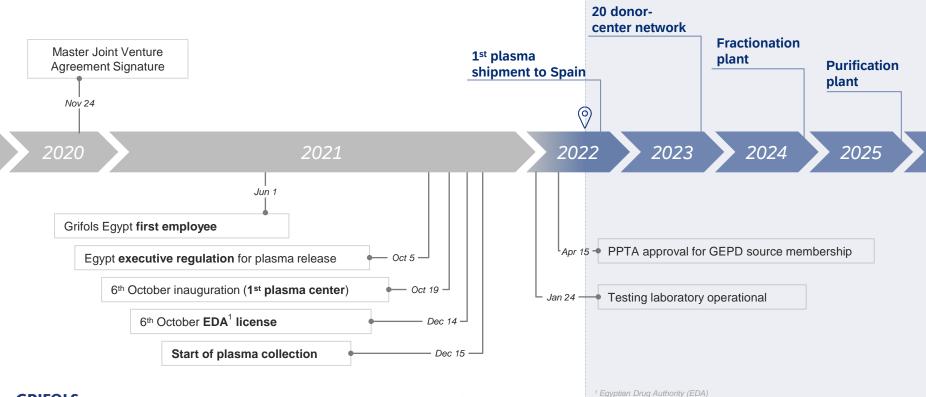
Government committed to the country's **self-sufficiency**



Achievements Clear the Path for Future Milestones



Year 1: Grifols Successfully Laid the Foundations of the Egypt Project



First EDA¹ Licensed Donor Center in Egypt



Paving the Way for the First Integrated Plasma-Supply Platform in Africa





Donor center, plasma testing labs, plasma freezers warehouse, Grifols Academy for Plasmapheresis and plasma operations offices





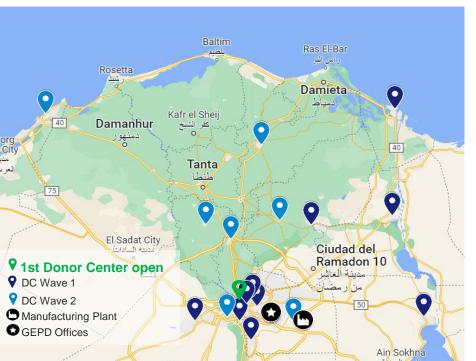
¹ Egyptian Drug Authority (EDA)

Canada Egypt

Plans on Track for the Construction of 20 Donor Centers

10 Plasma Collection Centers Projected in Wave 1 (10+ in Wave 2)







Plans on Track for the **Manufacturing Site** Construction



Responding to the Global Need for a Sustainable Plasma Model

Key Takeaways

Self-sufficiency is crucial to guarantee the sustainability of plasma-derived medicines supply

Grifols positions to serve as a best-in-class partner through private or public/private initiatives to support a sustainable plasma model

Grifols' unique **know-how** and leadership in **engineering** and **operations** represents a **competitive advantage** to execute these projects

Milestones in **Egypt** and **Canada** reflect how Grifols **delivers on its promises**

The implementation of international quality standards strengthens the safety & efficacy of our medicines

Best positioned to capture new opportunities...

Becoming a **blueprint** for **other countries** to increase plasma self-sufficiency levels

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BIOPHARMA

Poised for Performance in an Essential and Growing Market

Bill Zabel, EVP Commercial Operations

Joana Sàbat, SVP Global Marketing & Market Access



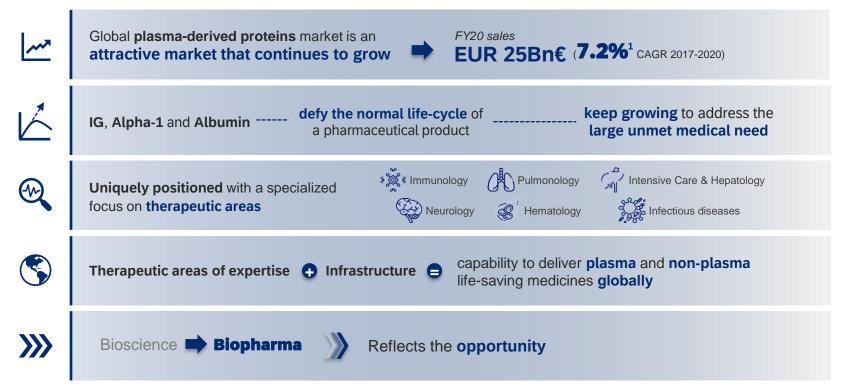
Poised for performance...





Grifols Is Well Positioned to Get Back to Growth

Driven by Biopharma Strengths



¹ The Worldwide Plasma Proteins Market, MRB, February 2022







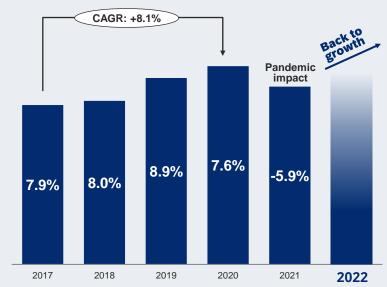
Underlying Demand Supports Further Growth

Core growth products

show positive trend in Q1'22 vs. Q1'21

IgG	Back to growth, driving Biopharma sales recovery
Albumin	Strong start of the year, driven by sales in China and U.S.
Alpha-1	Increasing market share and sustained growth during 2021 and 2022
HyperRAB®	Consolidated market leader, growing year over year

Bioscience Revenues (EUR in millions, growths at cc)







Main geographies

show significant recovery in Q1'22 vs. Q1'21



Recovering high-single-digit growth



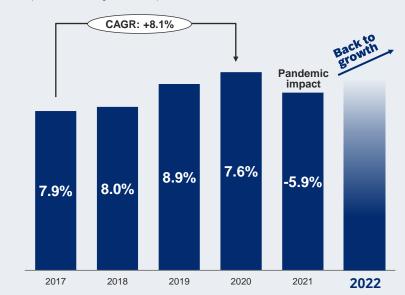
Strong albumin growth driving **double-digit**-growth



Sustained growth

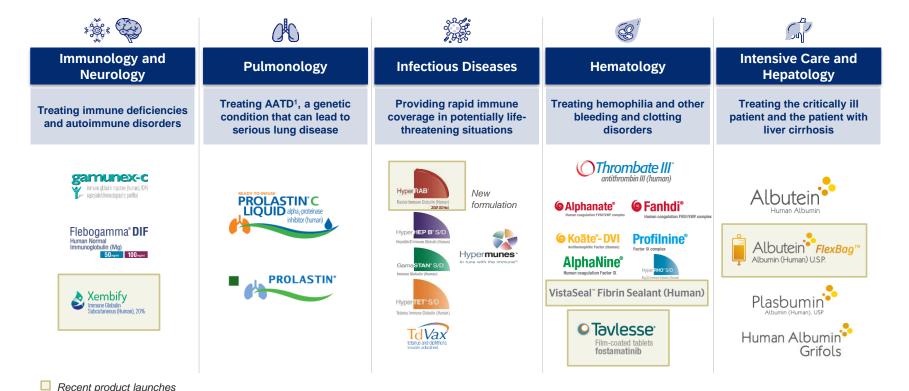
Bioscience Revenues

(EUR in millions, growths at cc)



Building on Biopharma Portfolio With Recent Product Launches

Deep Knowledge and Capabilities Coupled With a Focus on Therapeutic Areas



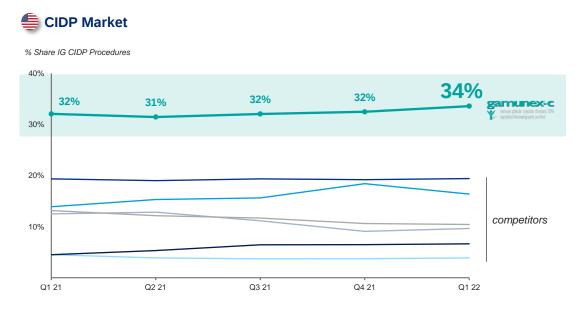






Leading an Increasingly Competitive U.S. CIDP¹ Market

Gamunex®-C Recognized as the #1 Prescribed IG for CIDP





Source: LexisNexis Trending Data, Medical Claims only; and Gamunex®-C and GammaKed share a J-code ¹ Chronic Inflammatory Demyelinating Polyneuropathy (CIDP)





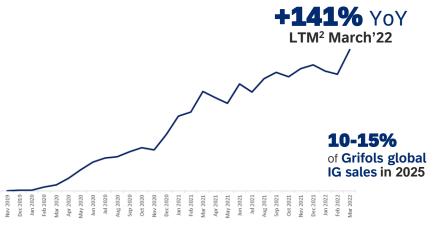


Xembify® Continues to Gain U.S. Market Share



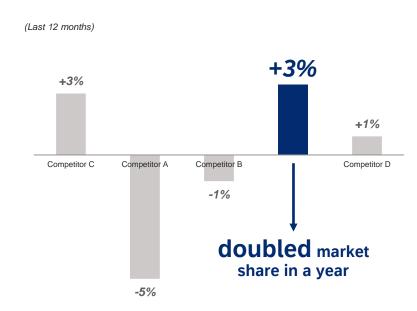
Supporting Patients by Providing a New Solution That HCPs1 Rely On





Over **1,000 U.S. HCPs** have prescribed Xembify[®] since its launch and **87% continue to prescribe** it

... and market share evolution in PI³



Source: Grifols Internal data

³ Primary Immunodeficiency (PI)



¹ Health Care Practitioners (HCPs)

² Last Twelve Months (LTM)

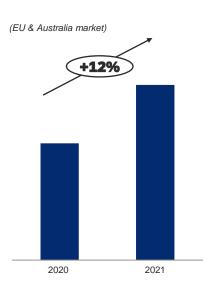




... While Expanding to New Geographies Outside the U.S.

SCIG Represents 23% of Volume in EU/Australia

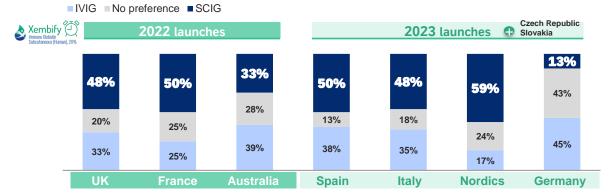
SCIG continues to grow



Sources: (i) Grifols Database provisional data 2021 (accessed May'22); (ii) Grifols ATU Study 2022. Base: All HCPs. Germany (n=40), France (n=40), Italy (n=40), Spain (n=40) UK (n=40) Nordics (n=29) and Australia (n=36)

Launching Xembify® in several European countries throughout 2022 and 2023 leveraging on increasing growth of SCIG preference over IVIG

Preferred route of administration for IG therapy by market (% HCPs)





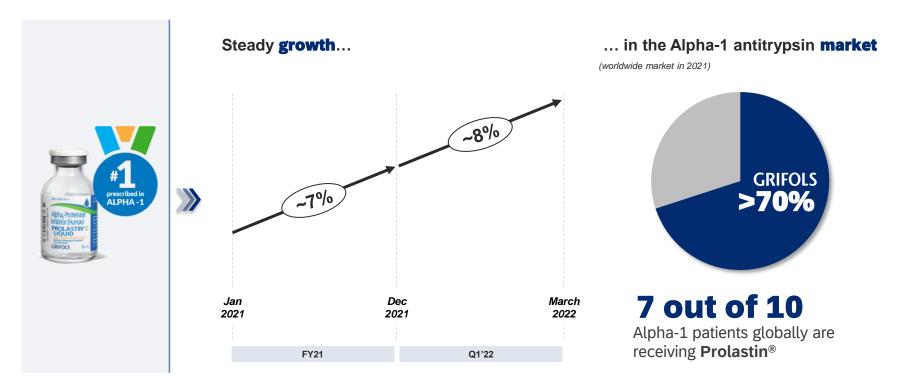
helping IG franchise to grow high-single-digit YoY





Leading the Alpha-1 Market for 35 Years

Prolastin^{®1} Shows Steady Growth in 2021 and Q1'22



Source: Grifols Internal data

¹ Prolastin[®] includes all Grifols Alpha-1 brands







Leading the Alpha-1 Market for 35 Years

Prolastin^{®1} Shows Steady Growth in 2021 and Q1'22



Sustained and reliable product supply...

Helping ensure HCPs and patients have access to Prolastin® when and where it is needed



Context

Competitor alpha-1 product shortage in Q4'20-Q1'21 challenged treatment continuity in EU



Grifols **was ready** to immediately **put in place all efforts**, guaranteeing **Prolastin**[®] to patients

... backed by Patient Support Programs

Patient health management programs provide personalized care and home infusion

- ✓ Personalized support and benefits beyond therapy
- ✓ Significant improvements in compliance with therapy and health outcomes after intervention year²

² Sandhaus RA, Turino G, Brantly ML, et al. The diagnosis and management of alpha1-antitrypsin deficiency in the adult. J COPD Foundation. 2016;3(3):668-682



¹ Prolastin[®] includes all Grifols Alpha-1 brands

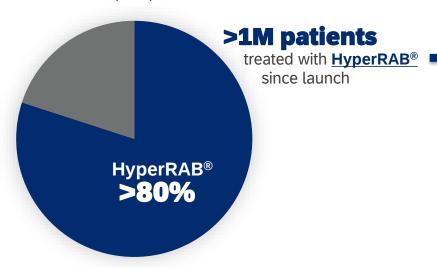




HyperRAB® Driving Hyperimmunes Portfolio

Record Sales in 2021

Rabies IG Market (2021)



Comprehensive HRIG¹ Solution

First and only high-potency HRIG that enables the delivery of more of the total dose per mL at the wound site, regardless of the wound size or patient weight ^{2,3,4}



Graph source: Grifols Database provisional data 2021 (accessed Mav'22)

⁴ Manning SE, Rupprecht CE, Fishbein D, et al; Advisory Committee on Immunization Practices Centers for Disease Control and Prevention (CDC). Human rabies prevention—United States, 2008: recommendations of the Advisory Committee on Immunization Practices. MMWR Recomm Rep. 2008;57(RR-3):1-28



¹ Human rabies immune globulin (HRIG)

² HyperRAB (rabies immune globulin [human]) Prescribing Information. Grifols.

³ Siegel J. Rabies Immune Globulin: Ensuring Administration Safety. Pharmacy Practice News Special Edition. 2019: 47-52.



Albutein FlexBag™ Successful Launch

Designed, Engineered and Manufactured by Grifols



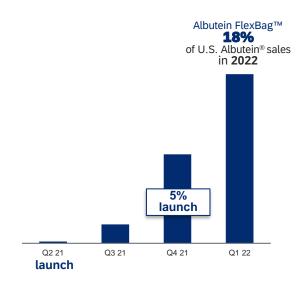
High degree of satisfaction among customers...

- ✓ First and only 5% 500mL offers convenience
- ✓ Availability of both 5% and 25% strengths



- ✓ Storage and portability vs. vials
- ✓ Ease of use/administration vs. vials
- ✓ Environmentally-friendly design

... reflected in **revenues** backed by fast adoption in the U.S.



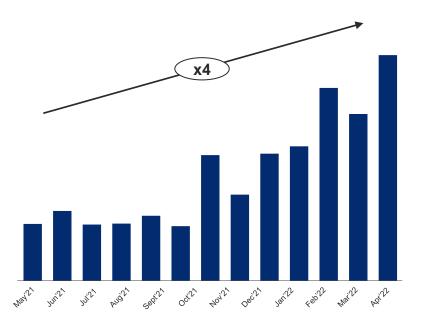
¹ Albutein® includes all Grifols albumin brands



Taylesse® Continues to Gain Traction

Successfully Launched in Multiple European Countries, Including EU41

○ Taylesse Sales volume (last 12 months)



First launching country **Germany**

On track throughout 2021

Launches proceeded during the pandemic

Exceeding volume targets in Spain and Italy

progress for

12 more countries
in 2022-2023

Sources: Internal data; (i) Price and Reimbursement; (ii) Health Technology Assessment ¹ Includes France, Germany, Italy and Spain





Vistaseal® Market-Leading Fibrin Sealant in the U.S.

Recovery of Elective Surgical Procedures Following Pandemic-Related Interruptions





Poised for performance...



... in an essential and growing market





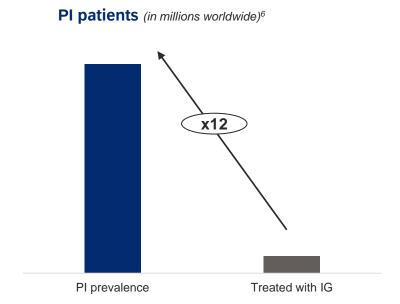


Further Growth in IG Market Driven by Progress in Pl¹ Diagnosis

Operating in an Essential and Growing Market

PI underdiagnosis facts

- Average diagnosis 12.4 years²
- Undiagnosed nh → 70-90%³
- **500,000** undiagnosed 🛍 🗐 4 (NIH estimates)
- >400 related-genetic defects⁵



⁶ Grifols Internal Source



¹ Primary Immunodeficiency (PI)

² Routes J, et al. J Clin Immunol. 2016;36:450-461

³ Primary Immunodeficiencies (PID) – Driving Diagnosis for Optimal Care in Europe, European Reference Paper

⁴ Modell V. Immunol Res. 2007;38(1-3):43-47

⁵ French MA, Tangye SG. J Infect Dis. 2019;jiz230. doi:10.1093/infdis/jiz230





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- Average diagnosis 12.4 years²
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 To-90%³
- 500,000 undiagnosed 🏥 = 4 (NIH estimates)
- >400 related-genetic defects⁵

Grifols Strategic Imperative Truncating time to diagnosis and treatment

Uniquely positioned to leverage our expertise in IG replacement therapy and diagnostics to improve the rate and time to an appropriate PI diagnosis



Grifols is actively partnering with insurers and patient groups to apply artificial intelligence programs to identify patients who may benefit from laboratory or genetic testing to detect the possibility of an underlying immune deficiency exists

¹ Primary Immunodeficiency (PI)

² Routes J. et al. J Clin Immunol, 2016:36:450-461

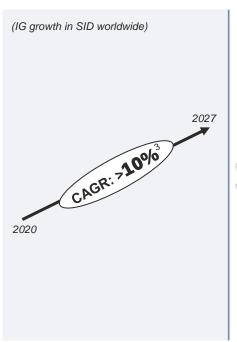
³ Primary Immunodeficiencies (PID) – Driving Diagnosis for Optimal Care in Europe, European Reference Paper

⁴ Modell V. Immunol Res. 2007;38(1-3):43-47

⁵ French MA, Tangye SG. J Infect Dis. 2019;jiz230. doi:10.1093/infdis/jiz230

SID Market Continues to Expand in Hematological Malignancies

Plans on Track for Xembify® to Treat New Key Indications e.g., CLL¹ and HGG²



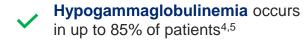


IG growth in SID

Increased use of SID-inducing medication in oncological and autoimmune indications



Extended survival rates of cancer patients



Infections are major cause of death in patients with CLL

Grifols has a Clinical

Development Plan for

HGG and CLL with

Xembify®

Continued **commitment** to SIDs patients $\mathring{\parallel}\mathring{\mathring{\parallel}}$



⁵ Seppänen M. Clin Exp Immunol. 2014;178(Suppl 1):10-13.



¹ Chronic Lymphocytic Leukemia (CLL)

² Hypogammaglobulinemia (HGG)

US PPTA Distribution Data

⁴ Friman V et al. Hematol Oncol. 2016;1-12: doi 10.1002/hon.2323. [Epub ahead of print]





Accelerating AATD¹ Diagnosis as Our Primary Goal

Further Expanding Our Testing Program Worldwide

AATD underdiagnosis facts

Avg. delay between the onset of symptoms and diagnosis



2.7² avg. by **p**

avg. number of **physicians seen** by **patients before diagnosis**



Patients present with common respiratory symptoms^{3,4}



Diagnosis does not depend on clinical presentation alone Laboratory testing is the only way to make the diagnosis^{2,5,6,7}

Grifols Strategic Imperative Truncating time to diagnosis and treatment Grifols' worldwide testing program (since 2006) >205,000 >16.000 >950,000 tests deficient severe patients performed patients found diagnosed (vs. >850,000) (vs. >190.000) (vs. >15.000) in Jun'2021 in Jun'2021 in Jun'2021

⁷ Global Initiative for Chronic Obstructive Lung Disease. Global Strategy for the Diagnosis, Management, and Prevention of Chronic Obstructive Pulmonary Disease. 2021 Report



¹ Alpha-1 Antitrypsin Deficiency (AATD)

² Campos MA, Wanner A, Zhang G, et al. Trends in the diagnosis of symptomatic patients with alpha1-antitrypsin deficiency between 1968 and 2003. Chest. 2005;128(3):1179-86.

³ World Health Organization. Alpha 1-antitrypsin deficiency: memorandum from a WHO meeting. Bull World Health Organ. 1997;75(5):397-415.

⁴ McElvanev NG, Stoller JK, Buist AS, et al. Baseline characteristics of enrollees in the National Heart, Lung and Blood Institute registry of α1-antitrypsin deficiency, Chest, 1997;111(2):394-403.

⁵ ATS, ERS. American Thoracic Society/European Respiratory Society statement: standards for the diagnosis and management of individuals with alpha-1 antitrypsin deficiency. Am J Respir Crit Care Med. 2003;168(7):818-900.

⁶ Sandhaus RA, Turino G, Brantly ML, et al. The diagnosis and management of alpha1-antitrypsin deficiency in the adult. J COPD Foundation. 2016;3(3):668-682.





Accelerating AATD¹ Diagnosis as Our Primary Goal

Dedicated Efforts to Increase the Diagnosis Ratio

AATD underdiagnosis facts

 Avg. delay between the onset of symptoms and diagnosis



• 2.7² b

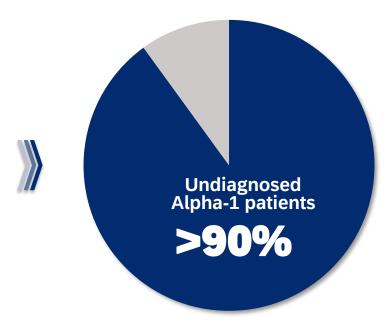
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Potentially Increasing Albumin Opportunity in Critical Care

Supporting Regular Albutein®1 Use as a Disease Modifying Treatment

Hepatology facts

of global population with **NAFLD**², the most common cause of chronic liver disease³

>14 million

cirrhosis cases

2nd leading cause of years lost of working life⁴

Liver transplant is the only procedure available today to avoid further complications and death

global market growth potential stemming from regular albumin use (>€400M)

Grifols Hepatology Program

Albumin medicine role in hepatology is explained by albumin blood levels (albuminemia), supporting its regular use as a disease modifying treatment



critical care area



48.9M cases of sepsis leading to 20% of all global deaths⁵

No conclusive guidance on the choice of fluid and resuscitation protocols exists as of today6

Albumin 1-time use to recover hypovolemia evolving to albuminemia-base treatment (Albios Balance/Ariss)

HES7 threat as a result of EMA's recent decision to suspend marketing authorization in EU



¹ Albutein[®] includes all Grifols albumin brands.

² Non-alcoholic fatty liver disease

⁴2021 EASL-Lancet Liver Commission on Liver Disease

⁵ WHO Sepsis Fact Sheets

^{6 2021} Surviving sepsis campaign

⁷ Hvdroxvethvl-starch (HES)

Poised for Performance in an Essential and Growing Market

Key Takeaways

Global plasma-derived proteins market is an **attractive market** that continues to **grow**

IG, Alpha-1 and albumin defy the normal life-cycle of a pharmaceutical product and keep growing

Strong key brands: Gamunex®, Xembify®, Prolastin®, Albutein® and HyperRAB®

Building on Biopharma portfolio by adding new proteins with a specialized focus on therapeutic areas

Expansion of **PI** and **SID** markets expected to **outpace potential erosion from new technologies**

Grifols' Diagnostic offers an unparalleled opportunity to continue increasing diagnosis rates and support Biopharma growth

GRIFOLS 2022 Investor & Analyst Day

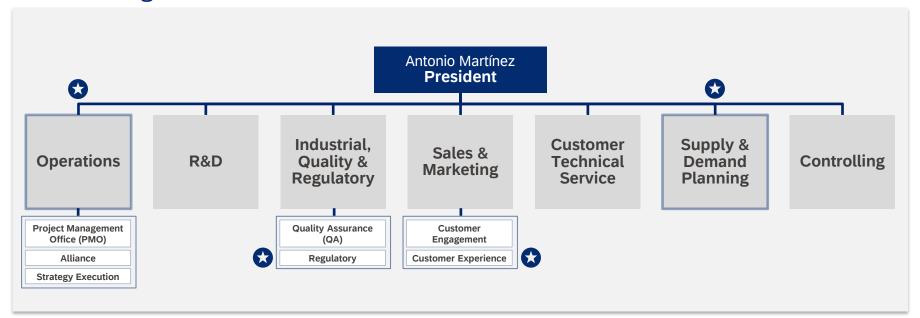
DIAGNOSTIC

Evolving Business Model to Strengthen Competitiveness

Antonio Martínez, President Diagnostic

A New Leadership to Accelerate Diagnostic

Streamlining the Business Unit to Increase Effectiveness and Drive Efficiencies



Focus and deep knowledge within the Diagnostic Business Unit

Changes implemented in the Business Unit



Diagnostic Mission

Improving care for donors and patients by providing laboratorians with innovative diagnostics tools

Growing the Business Based On Three Strategic Pillars



Consolidate and strengthen current business



Support Biopharma



Regulatory innovation & digital transformation







Combining Capabilities to Accelerate Growth

Broad Portfolio of Platforms, Know-How and Global Presence

Platforms

CLIA¹ Labs in U.S. & Europe **R&D** capabilities & manufacturing facilities of In Vitro Diagnostic (IVD) devices



CDMO² capabilities for therapeutic proteins



Regulatory Knowledge

Extensive regulatory knowledge in IVD and Pharma fields



Over The Counter (OTC) experience



Global Sales & Customer Technical Support

Sales & technical service in key countries



- ² Contract development and manufacturing company (CDMO) 3 Transcription Mediated Amplification (TMA)
- ⁵ Next Generation Sequencing (NGS)
- ⁶ Polymerase Chain Reaction (PCR)

Proprietary/Partnership Technology













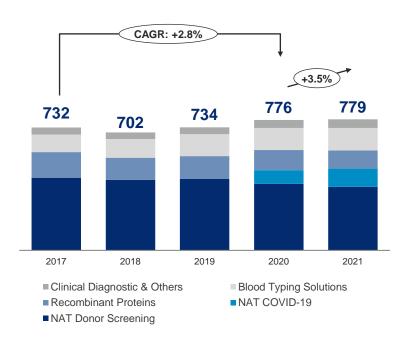




Diagnostic Division Reported Solid Growth in 2021

Driven by COVID-19 Innovation and Strong Underlying Business

Revenues (EUR in millions)



Core business of **NAT** achieved **sustained growth** and continued to **consolidate** in the main markets (U.S. and China)

NAT testing addressed the emergency of new pathogens (Zika, COVID-19) and represents an **opportunity** for **new segments**

Constant growth in Blood Typing Solutions (BTS)

Note: growths at constant currency (cc), which excludes exchange rate fluctuations vs. PY









Increasing Presence in Key Markets

Strategic Partnerships and Alliances to Drive Further Growth



Strategic long-term partner for Diagnostic:

- Consolidation of current business
- Innovation in partnership with a leading testing player
- Growth: developing new assays for global expansion













Immunohematology Market Delivering Strong Growth

Expanding Grifols' Blood Collection Capabilities



New red blood cells manufacturing facility in San Diego

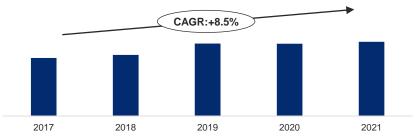




BTS constant growth led by DG Gel® System sales and fueled by Erytra Eflexis®

Worldwide market penetration

>7,000 instruments **50M** gel cards achieved in 2021













Supporting Biopharma Through High-ROI¹ Programs

Successful Alpha-1 Testing Strategy



Grifols' testing program continues to accelerate diagnostic globally





🖟 AlphalD

Validated, Convenient and Efficient Diagnostic Solution

- Cost-effective
- FDA approved
- Scientific Societies Support







>16,000 severe patients diagnosed



(figures since 2006)



Life-saving therapy for those patients who need it

² Chronic Obstructive Pulmonary Disease (COPD)



¹ Return On Investment (ROI)

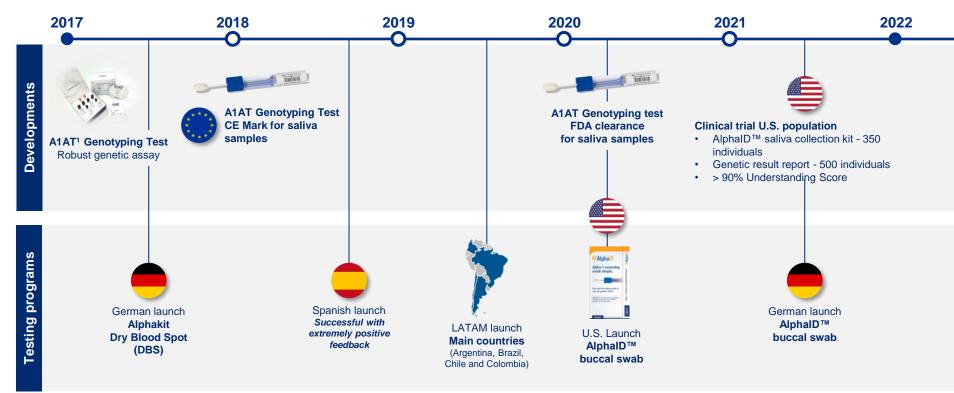






Our Journey to an Innovative Program...







¹ Alpha-1-Antitrypsin (A1AT)







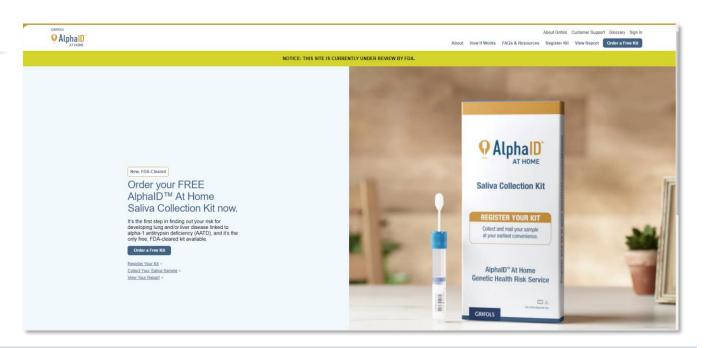
... To Achieve a New Milestone

Over The Counter 510(k) Filing Submitted to FDA: AlphaID™ at Home

May 2022

FDA SUBMISSION

The program is ready to launch once cleared



Over The Counter innovative solution to expand the U.S. screening program







Expanding the Model to Support Biopharma Growth

While Embracing Future Initiatives Including Biotest Projects

Basis for innovative testing strategies



Alpha-1 patients







Beyond









³ Inherited AT deficiency (hATd)

⁴ Antithrombin III (ATIII)







Grifols, Already Compliance With New In Vitro Diagnostics **Medical Devices European Regulation (IVDR)**

First Certifications for the New IVDR Received in December 2021



Grifols on track with the initial IVDR timing

Additional transition period (until May 2024) not needed



Promonitor Quick IFX

one of the **first near-patient testing** devices approved under the IVDR

Grifols will ensure on-time delivery of IVDR certification for all relevant products





Digital Transformation Promotes Higher Performance

Advanced Software Solutions and a New Customer Portal



Key strategic solutions to streamline customer operations

Blcodstream

SOFTWARE

New Bloodstream software as a single point management system for the NAT testing laboratory

Blood Typing Manager

New middleware solution connecting blood typing data across laboratory network system to improve patient care, enhance operational efficiency and minimize risks



Digital platform for technical documentation of our products:

Going Green



- All information, one site
- · Quick and easy access
- · Granted up-to-date documentation
- · Digital safety environment



Evolving Business Model to Strengthen Competitiveness

Key Takeaways

Strengthened leadership to increase effectiveness and drive efficiencies

Broad portfolio of platforms, assets and global presence to outperform competition and grow value

Secured U.S. key large customer while poised to increase penetration to fast-growing Chinese market through strategic alliance with SRAAS

Unique diagnostic capabilities to expand Grifols' Biopharma core markets and beyond

Grifols Alpha-1 **OTC breakthrough testing** to achieve FDA regulatory **remarkable milestone**

Enhanced field **digitalization** to support **further growth**



GRIFOLS 2022 Investor & Analyst Day

INNOVATION

Accelerating Innovation: Gaining Focus and Speed in Our Pipeline

Albert Grifols Coma-Cros, Chief Innovation Officer César Cerezo, SVP Drug Development Carter Keller, SVP GigaGen

Accelerating Innovation Through 3 Key Objectives

Further evolving our innovation organization

Gaining focus and speed in our pipeline (from past to present)

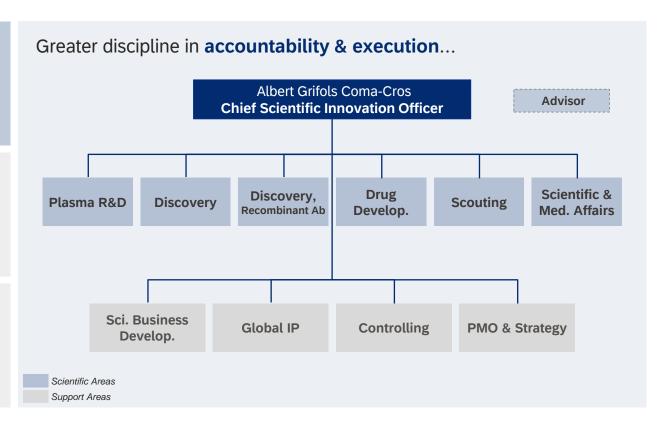
Building new innovation models (paving the future)

Focus on Scientific Areas While Elevating Controlling and PMO¹

Further evolving our innovation organization

Gaining focus and speed in our pipeline (from past to present)

Building new innovation models (paving the future)



¹ Project Management Office

Rationalizing Portfolio & Establishing Result-Oriented Processes

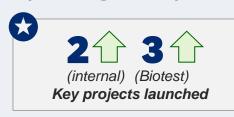
Further evolving our innovation organization

Gaining focus and speed in our pipeline (from past to present)

Building new innovation models

(paving the future)

Optimizing and re-prioritizing our R&D portfolio...





Ð

... while adopting a result-oriented governance



Clear direction with a 2030 Ambition target



Stronger business-weight (i.e., commercial, financial return) into project approval and prioritization processes



Increased weight of milestones achievement into bonus scheme

New Approaches: Developing a 360-Degree Innovation Ecosystem

Further evolving our innovation organization

Gaining focus and speed in our pipeline (from past to present)

Building new innovation models (paving the future)

Driving **growth**...

Digital & Data

OALKAHEST°

Bioinformatics platform

Digitalization of R&D organization

China

上海菜士 SRAASH SHANGHAI RAAS

R&D collaboration agreement

Strategic Alliances & Initiatives



Precision medicine (ATIII in sepsis collaborations)

ETHICON

Plasma innovation outlicensed (new projects)



Scouting for synergistic collaborations (increased efforts)

Balanced Risk-Value Pipeline

	Discovery	Pre-Clinical	Phase 1	Pha	se 2	Phase 3	Phase 4 / Regulatory	LCM	
.		recIG Spike in PdIG with enriched recombinant libraries (PI)				IVIG-PEG	Xembify® Europe Xembify® Bi-weekly dose	Xembify® Prefilled syringes	
lmmunology ﴾ پُرُپُرُدِ ا	y ++					IVIgG Next Gen PI			
		,				Xembify® in CLL			
Hepatology	/ ++					PRECIOSA D.Cirrhosis (Alb.20%)		FlexBag [®]	
Intensive Ca	are					APACHE ACLF (Alb 5%)		US, EUR	
Pulmonolog	gy	Alpha-1 AT Non-cystic fibrosis bronchiectasis		Alpha-1 AT 15% (SC) AAT deficiency			SPARTA - Prolastin-C®	Prolastin® EU 4-5gr vials	
(g) Hematology	+			ATIII in Sepsis***		Fibrinogen Cong. Deficiency & severe hypofibrinogen Fibrinogen Acquired Deficiency IVIGG Next Gen - ITP	Fostamatinib** ITP – Refractory patients		
Others	**	GIGA 564 GIGA 2328 Anti-CTLA-4 Anti-CTLA-4 mAb Oncology mAb Oncology		AKST4290 nAMD & DR			Fibrin Sealant Biosurgery Pediatric Use		
		GIGA 2339 HBV Recombinant hyperimmune Ig				Trimodulin sCAP			
្ត្រូវិទ្ធិ Infectious វិទ្ធិទី Diseases	+++					Cytotec® Pregnancy (CMV infection)			
Neurology	+++	AKST 1220 CADASIL		GRF6019 AD GRF6021 PD with Dementia	ABvac40 AD AKST4290 PD	AMBAR-Next			
Plasma projects Non-plasma projects Biotest projects Detailed nex									

^{*} Project of AlbaJuna (Grifols' invested company); ** Licensed rights from Rigel Pharmaceuticals in EU and other countries; *** Partnership with Endpoint Health



Notable Progresses in 4 Core Projects





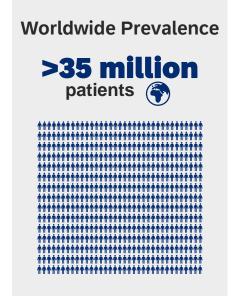


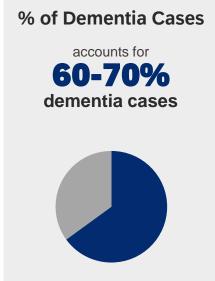


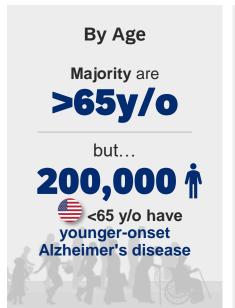
Alzheimer's Disease Therapy: An Urgent Unmet Need

5th Leading Cause of Death Worldwide

Progressive disease in which dementia symptoms gradually worsen over several years
No cure to date but research continues, and treatments are available to alleviate symptoms









Sources: Alzheimer's Disease International, Worlds Alzheimer's Report 2015, https://www.alz.org/Alzheimers-dementia/what-is-dementia



R&D Efforts Encompass the Entire Development Path

Innovation Strategy in Neurogenerative Diseases (ND)

Discovery

Plasma Proteome Chronokines

Understanding of plasma at the molecular level in healthy and disease states

Discovery of **chronokines** proteins with biological impact that changes with age

Pipeline expansion of therapeutic plasma fractions, small molecules, antibodies

Explore **therapeutic benefit** of plasma proteins in a wide range of CNS¹ disorders

Clinical Development Programs

Plasma Fractions & Small Molecules

GRF-6019

GRF-6021

Proprietary plasma fraction inducing neurogenesis

2 clinical programs

in AD² and PD³ patients with dementia in phase II

Active Immunotherapy

Aβvac40

vaccine against Aβ40

Placebo-controlled AB1601 trial in 120 very early AD and MCI⁴ patients in phase II



Two-Pronged Strategy

Plasma exchange

Albumin replacement

Remove toxic moleties from systemic and brain compartments

Immunomodulatory effects on AD pathological mediators

Restore pharmacological properties of Albumin

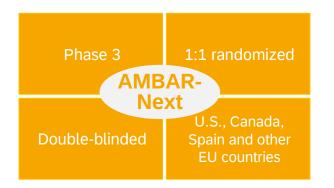


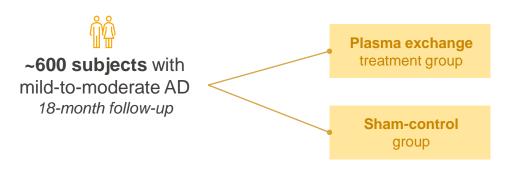
¹ Central Nervous System (CNS); ² Alzheimer's Disease (AD); ¹ Parkinson Disease (PD); ⁴ Mild Cognitive Impairment (MCI)



AMBAR-Next: Slowing Clinical Decline of Mild-to-Moderate AD

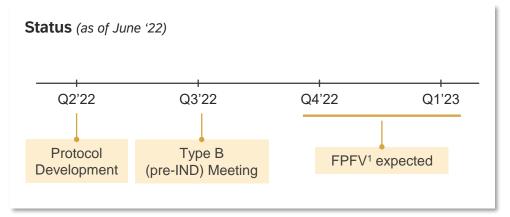






Centers and countries





¹ First Patient First Visit (FPFV)



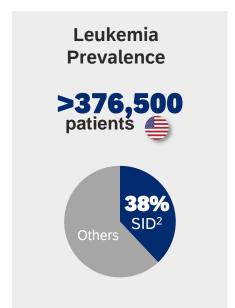


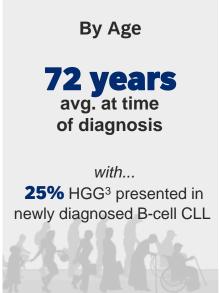


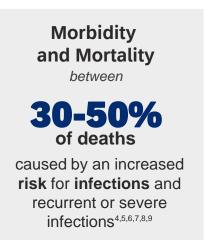
Targeting the Fastest-Growing Patient Segment for IG Therapy

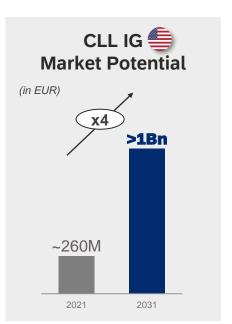
Chronic Lymphocytic Leukemia (CLL)

1 Accumulation of monoclonal, mature CD5+B cells in the peripheral blood, bone marrow, and secondary organs¹ No definitive cause has been established; targeted therapy regimens proven as efficacious treatments









¹ Burger JA. ² Secondary Immunodeficiencies (SID); ³ Hypogammaglobulinemia (HGG); ⁴ Wintrobe MM, Hasenbush LL.; ⁵ Hansen MM.; ⁶ Morra E, Nosari A. Montillo M.: 7 Hamblin AD, Hamblin TJ, 8 Morrison VA.: 9 Dhalla F, Lucas M, Schuh A, Bhole M, Jain R, Patel SY, et al



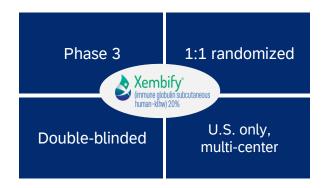




GigaGe & HBV

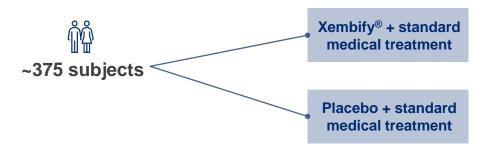
Xembify® to Prevent Infections in CLL Patients

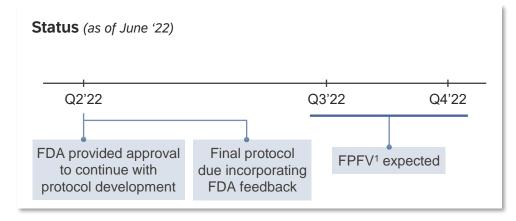
Treating Patients with Concomitant HGG and Recurrent or Severe Infections



Centers and countries







¹ First Patient First Visit (FPFV)

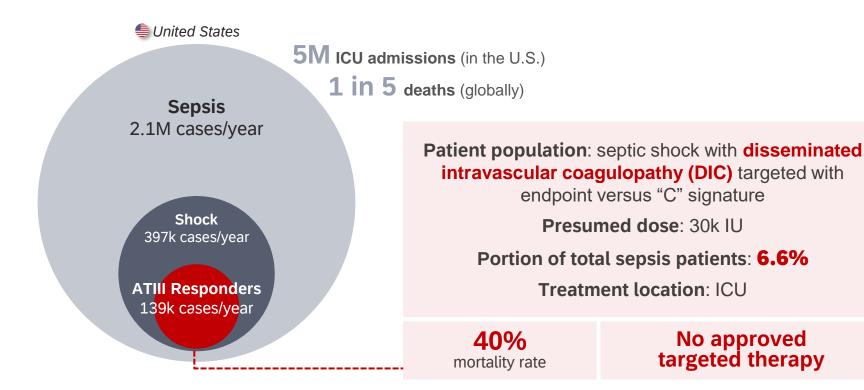




Giga(

ATIII in Sepsis: Very High Burden of Healthcare Costs

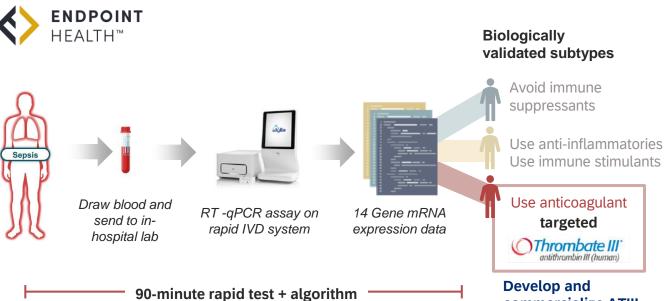
Outlook for U.S. Potential if Treatment Proven





Disruptive Diagnostic to Identify Thrombate III Responders

Global Collaboration Agreement and Solid Partnership With Endpoint Health



- Developed by Endpoint Health
- Distributed/sold by Endpoint Health + IVD partner
- Novel assay, system and algorithms / IP protected
- FDA authorized system and sample collection device

commercialize ATIII for severe sepsis

Gene expression array (Biocartis device)

- IVD test using small blood sample
- 15 marker array developed and validated in over 900 sepsis patients
- Gene expression array evaluating innate and adaptive immune response, complement, adaptive suppression and coagulation









Building the Future to Treat Severe Sepsis

Next Steps



Solid partnership with Endpoint Health (EPH)



- Supplying ATIII to conduct clinical trials
- Manufacturing capacity
- Advice on clinical development and regulatory activities (led by EPH)

Currently

EPH is waiting results of an observational study, which results will...

- Guide the pre-IND package submission to the FDA
- Help inform the design of the upcoming Phase II study

Moving forward Evaluate the feasibility of the Phase III study design with regards to endpoints and mortality event rates

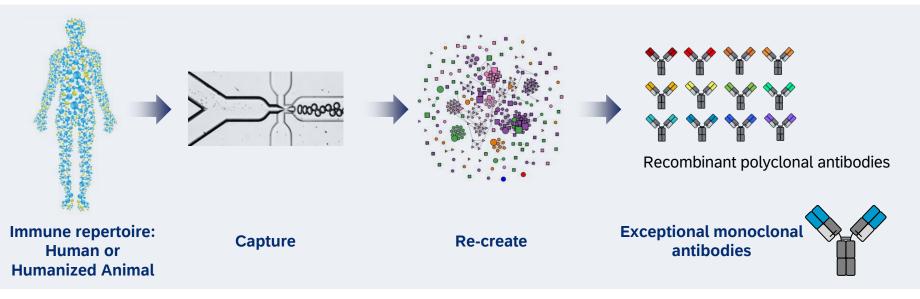


GigaGen Captures & Recreates Complex Antibody Repertories



World-Leading Immune Genomics Technology for Drug Discovery



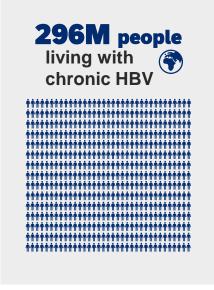


GigaGen & HBV

Chronic HBV¹: Widespread Disease With Significant Morbidity

Despite Currently Available Therapies and Vaccines

USD 3.8Bn² worldwide market

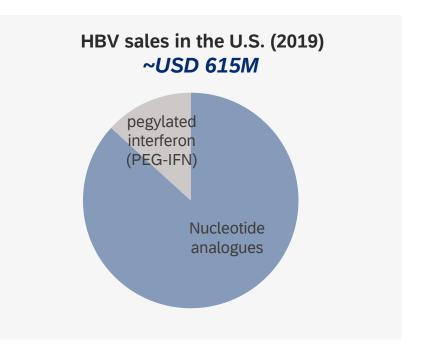


1.5M

newly infected patients per year³

20% of patients go on to develop cirrhosis & hepatocellular carcinoma

~820,000deaths per year⁴



⁴ Hepatitis B Foundation (hepb.org)



¹ Hepatitis B Virus (HBV)

² Global Data report: Hepatitis B Virus Infection – Global Drug Forecast and Market Analysis to 2029

³ WHO (https://www.who.int/news-room/fact-sheets/detail/hepatitis-b)

Anti-HBV Recombinant Polyclonal



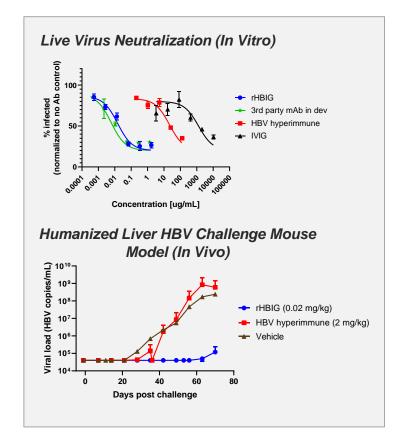


rHBIG

- Recombinant polyclonal antibody therapy (rHBIG) targeting HBV surface antigens
- Composed of >1,000 antibodies
- Derived from antibody sequences from human donors following HBV vaccine booster
- Polyclonal therapy ensures retained targeting of the many genotypes present and the large number of escape mutant variants that arise

Potency

Neutralizing potency of > 1,000x that of plasma-derived HBV hyperimmune therapy In vitro and in vivo models demonstrate efficacy potential GMP manufacturing initiation planned for Q4 2022







GigaGen's Antibody Platform Potential

rHBIG¹: the First Chapter



Recombinant Polyclonal Antibody Therapies

- Completely **new class of therapeutics** uniquely enabled by the GigaGen platform
- Capability that Grifols alone possesses
- Additional recombinant polyclonal therapeutics for infectious disease, immunology, transplant and other areas will be enabled with rHBIG's success

Monoclonal Antibodies

- Some indications are best addressed with monoclonal antibody treatments
- GigaGen's platform rapidly identifies rare, highly potent antibodies with unique activities from natural immune systems







Accelerating Innovation: Gaining Focus & Speed in Our Pipeline

Key Takeaways

Gaining focus and speed in our pipeline by rationalizing our portfolio and adopting a more result-oriented governance

Building new innovation models in digital and data, China and precision medicine

Seeking **U.S. indication for Xembify**® **in CLL**, the fastest growing patient segment for IG therapy

Leading the mild-to-moderate
Alzheimer's space with AMBARNext, a confirmatory trial for our
AMBAR program

Partnering with Endpoint Health to treat severe sepsis with ATIII, an innovative precision medicine approach

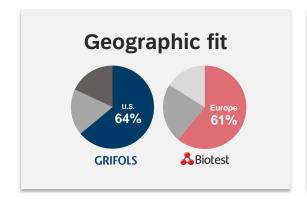
GigaGen's novel technology to create recombinant polyclonal antibody drugs for a wide range of infectious diseases

GRIFOLS 2022 Investor & Analyst Day



Maximizing Opportunities for Both Companies...

Grifols and Biotest: A Transformational Transaction







Broader commercial reach





Plasma collection and manufacturing capacity



Pipeline acceleration of key novel products



... Accelerating Grifols' Innovation Pipeline

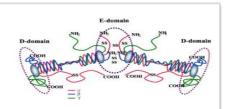
Bringing in **Precision Therapies**, Fibrinogen and Trimodulin, and Cytotect®

		Discovery	Pre-Clinical	Phase 1	Pha	se 2	Phase 3	Phase 4 / Regulatory	LCM
***			recIG		 		IVIG-PEG	Xembify ® Europe	Xembify [®]
չ∰્ર≰ Immun	nunology	++	Spike in PdIG with enriched recombinant libraries (PI)				IVIgG Next Gen PI		Prefilled syringes
			,				Xembify® in CLL	Bi-weekly dose	
	Hepatology/	++				PRECIOSA D.Cirrhosis (Alb.20%)		FlexBag®	
Intensive Care		***			/ 		APACHE ACLF (Alb 5%)		US, EUR
Puli	lmonology		Alpha-1 AT Non-cystic fibrosis bronchiectasis		15%	a-1 AT (SC) eficiency		SPARTA - Prolastin-C®	Prolastin® EU 4-5gr vials
(g) Her	matology	+			ATIII in \$	Sepsis***	Fibrinogen Cong. Deficiency & severe hypofibrinogen Fibrinogen Acquired Deficiency IVIGG Next Gen - ITP	Fostamatinib** ITP – Refractory patients	
Oth	hers	++	GIGA 564 GIGA 2328 Anti-CTLA-4 mAb Oncology mAb Oncology		AKST4290 nAMD & DR	AKST1210 ESRD-CI		Fibrin Sealant Biosurgery Pediatric Use	
0.4- 1-6-	lafa atia ca		GIGA 2339		1		Trimodulin sCAP		1
9 74	ectious eases +++		HBV Recombinant hyperimmune Ig		C		Cytotect® Pregnancy (CMV infection)		
Neu	urology	+++	AKST 1220 CADASIL		GRF6019 AD GRF6021 PD with Dementia	ABvac40 AD AKST4290 PD	AMBAR-Next		
* Project of AlbaJuna (Grifols' invested company); ** Licensed rights from Rigel Pharmaceuticals in EU and other countries; *** Partnership with Endpoint Health Plasma projects					Non-plasma projects	Biotest projects			

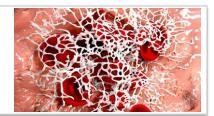
GRIFOLS

Fibrinogen: An Unparalleled Opportunity to Boost Revenue per Liter and Expand Margins

Fibrinogen is an **abundant**, **large**, and **complex** protein, which makes it ideally suited to be isolated from human plasma. It **accounts** for **95% of all coagulation factors** in the blood



Fibrinogen **promotes platelet aggregation**. It is **converted to fibrin** which forms the connecting glue in **blood clots**



Fibrinogen concentrate is a highly pure preparation of human fibrinogen that can be used to safely replace the absence or deficit of fibrinogen safer, much faster and with greater efficiency and precision than the current options of fresh frozen plasma or cryoprecipitates



Treating Acquired and Congenital Fibrinogen Deficiency

Playing an Essential Role in Blood Clotting and Hemostasis

Congenital deficiency

- Very rare, inherited bleeding disorder in which the body's ability to form blood clots is impaired
- Fibrinogen is used for treatment and prophylaxis of bleeding episodes in these patients

Deficiency **0-1 g/L**

Acquired deficiency

- Body's own fibrinogen lost due to major bleeding during elective surgical procedures or unexpected trauma events
- Fibrinogen is the first coagulation factor missing in major blood loss
- Replacement of lost fibrinogen is critical to restore effective hemostasis

Normal levels 2 -4 g/L

Deficiency

Massive bleeding

- · Major surgeries
- Postpartum hemorrhage
- Trauma

Source: IMS Data 2-2017, Internal Critical bleeding market research, LEK Critical Bleeding market assessment 2017, and Biotest market research







Clotting Assessment: Critical to Monitor Deficiency

Monitoring Patient Coagulation Status Allows Administering Precise Therapies

Point-of-Care (PoC) diagnostic...



... bringing in positive market implications

Prediction of massive transfusion requirement

Creation of **goal-directed** and **individualized** coagulation algorithms that may **improve patient outcome**

Distinguish the most important coagulation deficiencies (including fibrinogen)







ROTEM®

- Goal-oriented treatment protocols proposed for bleeding management in surgery and trauma patients
- · Rapidly developing field of acute medicine



Grifols is uniquely positioned to leverage its deep Diagnostic expertise

Most widely-used devices...

purce: Benes, Jan et al. "Viscoelastic Methods of Blood Clotting Assessment - A Multidisciplinary Review." Frontiers in medicine vol. 2 62. 14 Sep. 2015, doi:10.3389/fmed.2015.00062



Promising Results in Both Clinical Trials

Congenital FD¹

Phase I/III study: Largest clinical trial in congenital fibrinogen deficiency worldwide

Treatment of adults and children

Results confirm high expectations regarding efficacy and safety...

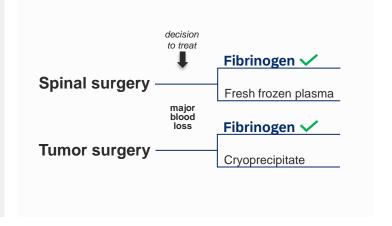
- Expected pharmaco-kinetics and -dynamics (Phase I), excellent efficacy and safety (Phase III)
- 175 bleeding events (BEs) treated in 36 patients of all age groups
- Overall hemostatic response assessments of 175 BEs demonstrated a treatment success in nearly all cases
- Study completed

Acquired FD¹



Phase III study in severe spinal surgery and pseudomyxoma peritonei (tumor) surgery

- Non-inferiority study compared to standard of care (fresh frozen plasma or cryoprecipitate)
- Interim analysis with 120 subjects (June '22) confirms planned patient number
- Recruitment ongoing 150 of 200 patients recruited
- Other interim analysis to define final sample size expected in December '22

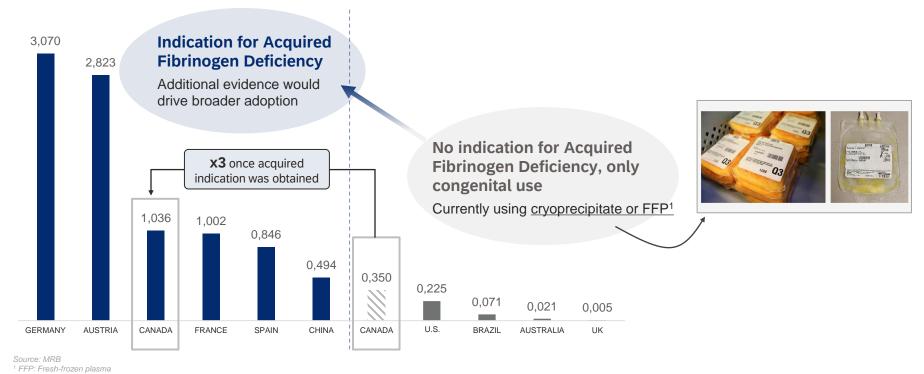


¹ FD: Fibrinogen deficiency



Acquired Fibrinogen Deficiency Indication: Broader Adoption and Opportunity

Fibrinogen consumption per capita (g/1000 inhab.)







Seizing Fibrinogen's Large Market Opportunity

Current Trends Underscore Its Potential While Further Benefiting Patients



Improved Point-of-Care diagnostics

 Increasing knowledge, availability and reimbursement worldwide

(NEW!) Precision Therapy



Increased awareness

- European and Canadian guidelines
- Diagnostic companies and plasmaderived medicines companies increasing education on Precision Therapy



Change of market dynamics

 Shortage of cryoprecipitate during pandemic driving fibrinogen concentrate usage with improved customer experience in the U.S. Expected CAGRs 19-27 (volume)





If Acquired Deficiency indication is obtained (based on Canada example)





Source: MRB, November 2021



Trimodulin: A Potential Blockbuster

Unique Polyvalent IG Composition



Important functions

- Recognizes pathogens, also part of innate immune system
- IgM and IgA can be secreted and are present directly on the pulmonary surface
- Strong anti-inflammatory effects by acting through cellular receptors
- Scavenge virulence factors, such as lipopolysaccharides which lead to inflammation
- Binds and modulates activated coagulation factors, complement factors, and cytokines

	IVIG	Trimodulin	
IgM	-	~23%	
IgA	-	~21%	
IgG	≥95%	~56%	
Patient target	e.g., patients with immunodeficiency	Clinical development in severe COVID-19 and sCAP ¹	

¹ Severe community acquired pneumonia: Severe CAP (sCAP) is usually defined as CAP that requires admission to the intensive care unit (ICU)

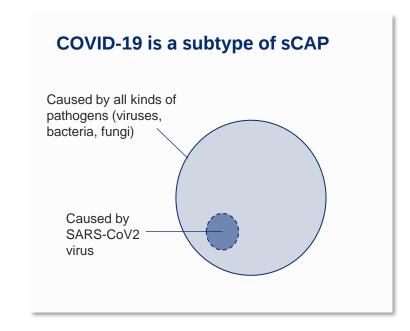
sCAP1: A Leading Cause of Illness and Death Worldwide2

A High Unmet Medical Need

Pneumonia requiring supportive therapy within a critical care environment

23-58%^{4,5} mortality range

no significant changes over the past decades despite the availability of improved broad-spectrum antibiotics



¹ Severe community acquired pneumonia: Severe CAP (sCAP) is usually defined as CAP that requires admission to the intensive care unit (ICU)

⁵ Sirvent et al. 2013, Med. Intensiva 37:308e 15.



² Wunderink 2014, N Engl J Med 370;6.

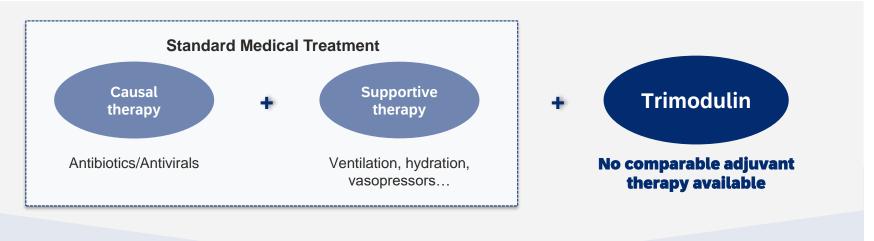
³ Chest Journal (CHEST) (chestnet.org)

⁴ Woodhead, 2006, Critical Care 10:S1, p3.



Changing the sCAP¹ Treatment Paradigm

Trimodulin Improves Patient Response With Further Decrease in Mortality



- ✓ ▼ overshooting immune-mediated tissue damage
- ✓ ▼ risk for lung inflammation, severe sepsis, septic shock, respiratory and multi-organ failure
- ✓ Prevention of secondary infections



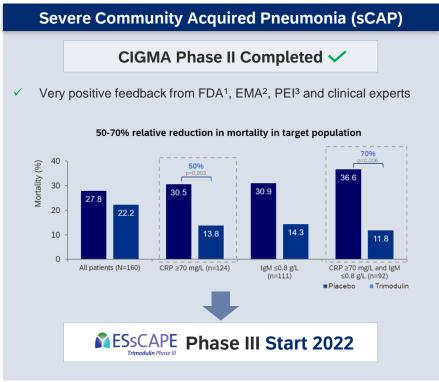
¹ Severe community acquired pneumonia: Severe CAP (sCAP) is usually defined as CAP that requires admission to the intensive care unit (ICU)

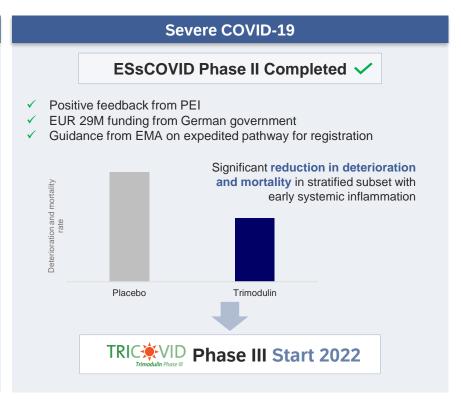


Cytotect® Pregnancy

Extensive Development Program in Severe Pneumonia

Patient Groups and Disease Stages for Optimal Therapy Identified in Phase II Studies





¹ U.S: Food Drug and Administration; ² European Medicines Agency; ³ Paul-Ehrlich-Institute;



Cytot Pregn

Leveraging on Phase II Results to Start Phase III Trials in 2022



(Start 2022)



(Start 2022)

Randomized, placebo-controlled, double-blind, multi-center, phase III trials investigating the efficacy and safety of Trimodulin in adult hospitalized subjects

SCAP patients



480-780 subjects

- Subjects on invasive mechanical ventilation (within <12h)
- Subjects with inflammation (CRP >70 mg/L)
- SARS-CoV-2 negative



COVID-19 patients



334 subjects

- Subjects on low-flow oxygen, high-flow oxygen, NIV
- · Subjects with early systemic inflammation





Grifols' Capacities Help Address the Opportunity

Potential Addressable Markets

Potential treatment

OPPORTUNITY

sCAP

Adjunctive treatment of patients with severe Community Acquired Pneumonia (sCAP) who require invasive mechanical ventilation and have CRP¹ >70 mg/L

SCAP market size350,000patients

initial target population
>80,000 patients/year

Significant **upside** due to **higher price** depends on clinical trials' data

TRIC-

N ESSCAPE

COVID-19

Add-on therapy to standard of care in adult patients with severe COVID-19 and CRP ≥50 mg/L

- Early approval will be beneficial for faster sCAP approval
- Development risk covered by public funding
- Even with vaccination, >20,000 patients/year in EU are expected over the next three years

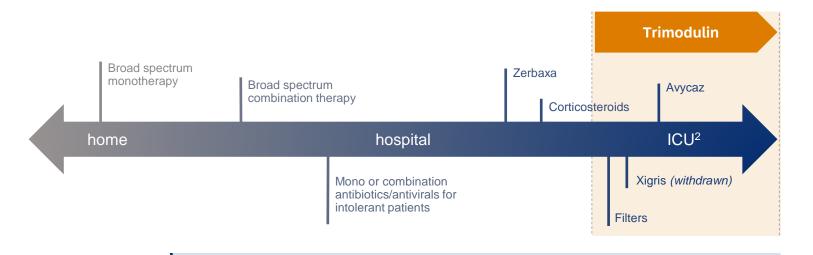
Source: Biotest market research

1 C-reactive protein (CRP)



No Direct Competitors for Trimodulin in sCAP¹

Harnessing the Opportunity to Address a Critical Medical Need





No direct competitors: no IgM enriched immunoglobulins on the market or in clinical trials

High medical need: high mortality rate despite antibiotics or antivirals

Commercial advantageous pricing opportunity

¹ Severe community acquired pneumonia: Severe CAP (sCAP) is usually defined as CAP that requires admission to the intensive care unit (ICU)

² Intensive Care Unit (ICU)



CMV¹ Infection: A Disease With Large Unmet Medical Need

No Approved Treatment for This Disease

The most common congenital infection in developed countries

0.3-2.4% prevalence in newborns

Infected women may transmit the virus to the fetus



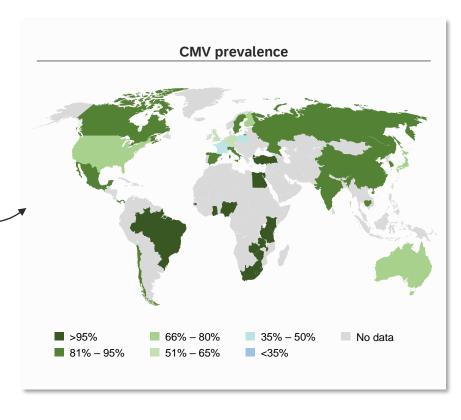


Lower rates





- Hearing loss
- Intellectual disabilities
- Premature birth
- Development delays
- Death



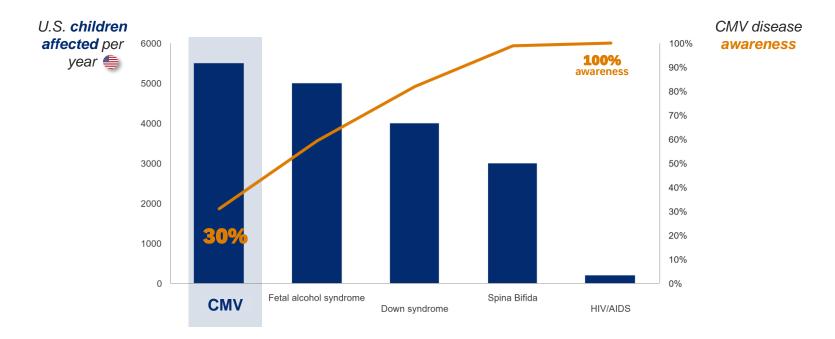
¹ Cytomegalovirus (CMV)





Very Low Awareness of CMV Infection in Pregnancy

Potential Effective Treatment to Increase Awareness and Diagnosis









Cytotect®: Tackling CMV¹ With a High-Potential Specialty Protein

First Treatment to Prevent Transmission from the Mother to the Unborn Child

It **binds to CMV** and **avoids infections** of host cells and presents CMV particles for phagocytosis



Modulates and interacts with immune cells (dendritic cells, monocytes, B- and T-cells), exerting a positive immunological balance

Anti-CMV antibodies in Cytotect® are actively shuttled through the placenta

These CMV-specific antibodies **block** the **infection from all CMV genotypes** and from **virus variants** that are resistant to virostatics



Clinical Experience Suggests Good Efficacy of Cytotect®

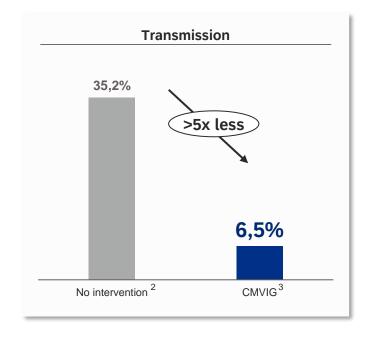
Significant Transmission Reduction When CMVIG1 Is Administered

Previous prospective, observational study has described success of **study protocol**



Adequate criteria for Phase III leveraging on previous experience:

- Inclusion and diagnosis criteria revised
- Accelerating treatment start
- Adjusting the dose
- Establishing right intervals



³ Treatment group (n=153), Maternal-fetal transmission at gw20: 6,5% (10/153). Kagan et al. Ultrasound Obstet Gynecol 2021; 57: 560-567



¹ CMVIG: Cvtomegalovirus Immunoglobulin

² No intervention group (n=108), Maternal-fetal transmission at gw20: 35,2% (38/108). Kagan et al. Ultrasound Obstet Gynecol 2019; 53(3): 383-390







PreCyssion Trial: Prevention of Maternal-Fetal CMV Transmission

Following Primary Maternal Infection With Gestational Age ≤ 14 Weeks



Objective:

Demonstrate efficacy and safety of Cytotect® in preventing maternal-fetal transmission of CMV



Study Design

- Pivotal, clinical Phase III
- Open-label
- Single-arm
- Prospective
- Multicenter
- · With historical control group
- 80 subjects 13 of 80 patients recruited (as of June 15, 2022).

 Recruitment dependent on the course of the pandemic (hygiene measures reduce CMV transmissions).





Experts Unveil the Uniqueness of Anti-CMV

Success of Cytotect® Represents an Important Opportunity

Number of births per year





Neonates with congenital CMV infection (0.5% - 1.0% of all newborns)



Pediatric infectious disease **experts acknowledge Hyper anti-CMV** and antivirals are the **only therapeutic alternatives** to treat these cases

Important opportunity behind phase III success

Diagnostic tools could be developed, leveraging on Grifols' expertise, to accelerate treatment adoption



Grifols' Scale and Commercial Strength: Key to Unfold the Full Value of a New Generation of Plasma Medicines

	Fibrinogen	Trimodulin	Cytotect® Pregnancy
Expected market approval	(combined approval: acquired & congenital deficiency)	\$\rightarrow\$ 2024-2025	2024-2025
Key considerations	 Results of AdFIrst study for acquired FD¹ (interim results June '22) and completed phase I/III trial in congenital FD will serve as the basis for the combined approval in Europe FDA meeting in H2'22 to provide guidance on acceptance of clinical data for BLA 	 Study duration dependent on interim analysis results Approval timelines assuming earlier TRICOVID trial success 	 As for now, only European market approval is considered Intention in the mid-term to seek U.S. market opportunity
¹ FD: Fibrinogen deficiency			

Grifols' Scale and Commercial Strength: Key to Unfold the Full Value of a New Generation of Plasma Medicines

Significantly improving revenue per liter and margins					
USD 30-50M	USD ~300M	USD ~30M			
 Greater commercial reach Complementary presence in important markets (US) Ability to develop new markets 	 Scalability of production volumes Full market potential of the main indications becomes addressable 	 Commercial reach and promotion Diagnostic capabilities 			
USD 400-800M	USD 1-2 Bn	TBD			
Leveraging on Diagnostic capabilities Expanding into other geographies (e.g., China)	Peritonitis/lymphopenic sepsis: USD >1Bn each Transplant: USD >30M Pediatric sepsis: USD 80M Multi-drug-resistance: USD 0.2Bn	Potential upside from U.S. Substitution USD > 200M			
	 USD 30-50M Greater commercial reach Complementary presence in important markets (US) Ability to develop new markets USD 400-800M Leveraging on Diagnostic capabilities Expanding into other geographies	 USD 30-50M Greater commercial reach Complementary presence in important markets (US) Ability to develop new markets Leveraging on Diagnostic capabilities Expanding into other geographies (e.g., China) Scalability of production volumes Full market potential of the main indications becomes addressable USD 1-2 Bn Peritonitis/lymphopenic sepsis: USD >1Bn each Transplant: USD >30M Pediatric sepsis: USD 80M 			

GRIFOLS

Grifols and Biotest: A Transformational Transaction to Maximize Both Opportunities

Acquisition **enhancing** Grifols and Biotest **opportunities** while **accelerating pipeline**

Expanding and diversifying plasma sourcing by adding 31 European centers and increasing revenue footprint in EMEA

Shared **values** and **culture** based on strong **family heritage**

Two precision therapies
(fibrinogen and trimodulin) and a
new indication for Cytotect®
with an expected market approval
by 2024-2025

Unique opportunity to **boost** revenue per liter and expand margins

Contributing to increase fibrinogen opportunity up to **USD 400-800M** and trimodulin to **USD 1-2Bn**

GRIFOLS 2022 Investor & Analyst Day

FINANCIALS

Investments in Plasma and Innovation Pipeline to Accelerate Growth and Profitability

Alfredo Arroyo, CFO

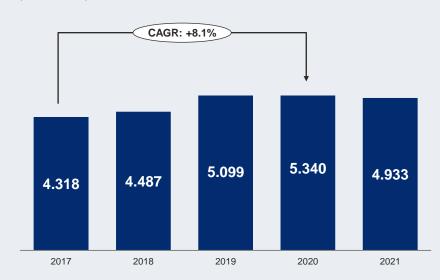
Strong Underlying Fundamentals

Committed to Long-Term Growth

- Global player in the attractive plasma market
 (7.2%¹ CAGR₂₀₁₇₋₂₀₂₀ for the industry) with strong position in
 the U.S.
- Leaders in diagnostic solutions to screen whole-blood and plasma
- Strengthening global presence in China through SRAAS strategic alliance, Europe, Middle East and Africa, and Canada (operations in 100+ countries and subsidiaries in 30+)
- Largest plasma-center network (400+) and industrial capacity (22mL+) to support underlying demand
- Reinforced pipeline to deliver a balanced risk-value portfolio

Grifols revenues

(EUR in millions)



Note: CAFR at constant currency (cc), which excludes exchange rate fluctuations vs. PY

¹ The Worldwide Plasma Proteins Market, MRB, February 2022

Doubling-Down Efforts to Increase Performance in the Past 2 Years

Reinforced **plasma capabilities** and **capacities**, supporting acceleration of **plasma collections**

Biotest investment bolstering innovation, adding two unused plasma proteins

Global footprint while strengthening presence in Europe, China, Canada and Egypt

CAPEX and **R&D** investments to support future growth

Commitment to achieve deleverage

Divested non-strategic assets while executing on structural cost savings

Full Year Performance Marked by COVID-19

2021 Key Financials

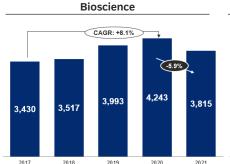
(EUR in millions)	H1'21	H2'21	2021
Revenue	2,537	2,397	4,933
% Growth	+2.3%	(9.8%)	(3.7%)
Gross Margin	1,114	849	1,963
% Margin	43.9%	35.4%	39.8%
R&D	159	196	355
SG&A	507	555	1,062
EBITDA	635	327	962
% Margin	25.0%	13.6%	19.5%
EBITDA Adjusted	637	377	1,014
% Margin	25.1%	15.8%	20.6%
Group Profit	267	(78)	189

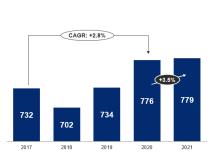
The decline in plasma volumes caused by COVID-19 was the primary driver of extraordinary financial impacts

- Lower collections and higher cost per liter impacted revenues and margins particularly in H2 2021
- Strong underlying demand backed by price increases
- Higher R&D and SG&A due to the integration of new companies, transaction and restructuring costs, and inflationary pressures

Note: all growths at constant currency (cc), which excludes exchange rate fluctuations vs. PY

Bioscience Constrained by Plasma Supply While the Other Divisions Grew in 2021

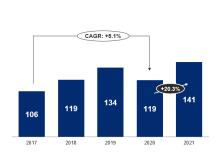




Diagnostic



Bio Supplies



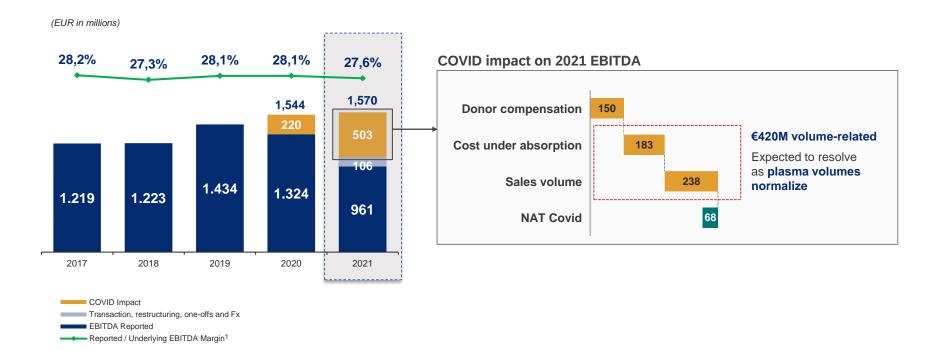
Hospital

- Underlying demand strong of key proteins
- Plasma supply constrained IG volume growth, especially in the U.S.
- Strong performance of new product launches: Xembify[®] gaining traction, VISTASEAL[™] and TAVLESSE[®]
- Growth driven by COVID-19 test sales in Spain and Hungary (+34% vs. 2020)
- Underlying NAT donor screening impacted by Zika testing mandate termination; whole-blood donations strong in the U.S. and Japan
- Blood Typing Solutions strong growth also contributed due to higher volume in the U.S.

- Bio Supplies Commercial, declined due to lower sales of blood cells and serum triggered by lower donations
- Plasma sales to third-parties increased based on pre-existing supply agreements
- Strong growth following normalization of hospital pharmacy investments
- Noteworthy sales of Pharmatech, and IV therapy
- Contract manufacturing growth stemmed from albumin bags and third-party sales



Underlying EBITDA Margin Revealed Strong Fundamentals



¹ Reported EBITDA for 2017-2019, and Underlying EBITDA for 2020 as per FY21 Results Presentation





Investments in plasma and innovation pipeline...

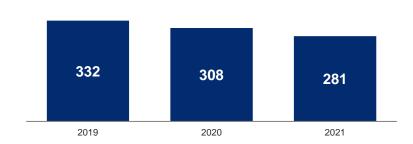


Strategically Invested to Support and Accelerate Growth

Enhancing industrial capacities

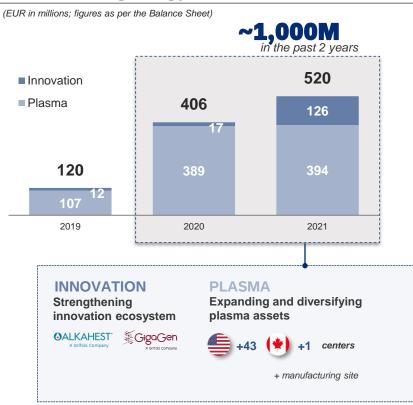
(EUR in millions)

EUR 900+ million CAPEX in 3 years



- New 6ML plasma fractionation plant in Clayton, North Carolina
- World's largest sterile purification and filling plant in Clayton
- · New albumin purification and sterile filling plant for flexible containers in Dublin
- · Expansion of fibrin adhesive and topical thrombin production plant in Barcelona
- Upgrade Grifols' facilities in Canada

Strengthening plasma and innovation

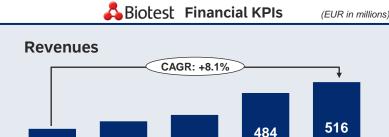


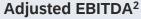
Biotest: A Transformational Investment

Complementary Business to Boost Performance

- Family business founded in 1946, HQ in Dreieich (Germany)
- Specialized on immunology and hematology
- Broad plasma pipeline to be launched in the short-term
- Direct commercial presence in 10 countries, marketed in 90+
- Manufacturing sites reaching 3mL plasma production capacity
- Strong presence in Europe: 31 plasma centers*

Closing Investment EUR 1.5Bn Capital: 70% Voting rights: 96% Preferred shares: 43%





378

2017

400

2018

Excludes mainly Next Level Project costs (production and R&D costs)

419

2019

2020



GRIFOLS

¹ As of June 30, 2022

147

2021

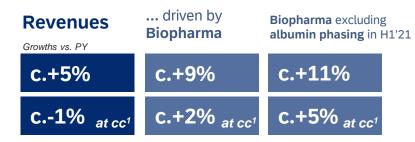


Investments in plasma and innovation pipeline...



Improving Performance in the First Half of 2022

Outlook H1 2022



Performance leveraging on:

- Accelerated plasma collections already reached pre-COVID weekly levels
- · Price increases on key proteins
- Robust underlying demand
- Product mix driven by subcutaneous immunoglobulin (SCIG)
- Higher alpha-1 patient base and strong albumin demand
- Partially impacted by albumin positive temporary effect in H1'21 due to commercial integration with SRAAS

EBITDA Margin

Reported



Significant sequential improvement vs. H2 2021

- Higher plasma volumes, price increases, product and country mix
- Financial discipline: cost savings and R&D prioritization
- Inflationary labor costs persist
- Donor compensation still high
- Fixed plasma costs decline as volume recovers

Constant currency (cc), which excludes exchange rate fluctuations vs. PY

Positive Performance for 2022 and Beyond

Expected Strong H2 Performance

Plasma

Collections expected to continue **accelerating** throughout 2022 driven by:

- · New and recently-acquired centers
- Larger volumes from regular centers
- Plasma collections upside if B1/B2 visa restrictions in the U.S. southern border are lifted
- Technological, digital and operational enhancements

Revenues

Sound revenue growth stemming from **Biopharma**, supported by:

- Strong underlying demand
- Global price improvements
- Product and geo mix

Margins

Sequential gradual expansion

- Triggering plasma volume
- Cost savings
- Still high donor compensation; inflationary pressures including wages

Despite Short-Term Challenges, Commitment to Deleveraging Remains Firm

Deleveraging path supported by...

Gradual

1 sequential EBITDA improvement

- · Plasma collection momentum
- · Strong underlying demand
- Global price improvements
- Product mix
- Lower cost per liter

Reinforcing our financial discipline

	Business optimization	Structural cost plan and R&D prioritization EUR 100M savings
	Cash dividends	No cash dividends until leverage ratio <4x
>	Divestments	 Hemostasis technology line (USD 25M) (Oct'21) Sale of VCN Biosciences (USD 75M¹) (Oct'21) Closure of the production of blood bags (Nov'21) Sale of Hospital Software BU (USD 100M) (already agreed)
	CAPEX	Lowering CAPEX in 2022 and 2023 as already well-invested
	M&A	Not pursuing any meaningful M&A

Prepared to Respond to the Current Macro-economic Context

Inflation

Inflationary pressures driving higher incentive to donate Impact on labor costs

No significant impact on energy costs (<1% of cost per liter)

Interest rates

Optimized financial structure. Low exposure to interest rate hikes, since c.65% of debt tied to a fixed interest rate

Exchange rate EUR/USD

FX tailwind in 2022

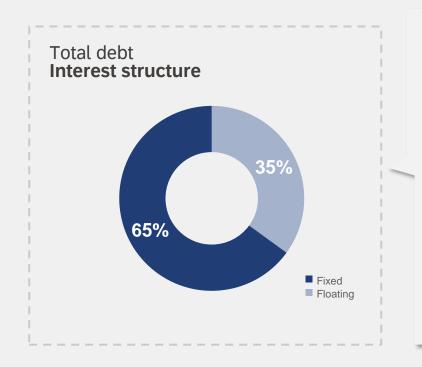
Ukraine-Russia

No direct commercial or industrial presence in Ukraine/Russia Representing <0.3% of total sales

Supply chain

Cross-licensing of our industrial facilities supporting efficient supply chain management

Low Exposure to Interest Rates Hike: 65% Fixed



EUR floating debt

13% of total debt outstanding

USD floating debt

22% of total debt outstanding

Investments in Plasma and Innovation Pipeline to Accelerate Growth and Profitability

During 2020 and 2021, Grifols invested to drive further revenue growth and margin expansion

Plasma collection momentum supports sequential performance improvement

Delivered on commitments: divestments and cost optimization

Focus on value creation assets to fuel cash flow and margins

Deleveraging remains a key priority, levering on EBITDA improvement and cash flow generation

Prepared to respond to the current macro-economic context

GRIFOLS 2022 Investor & Analyst Day

FINAL REMARKS

Delivering on Commitments Outlined in 2020 and 2021

Víctor Grifols Deu, co-CEO

Our Commitments



Innovation



Core Business



Organization

Accelerating innovation and optimizing our portfolio

Gaining focus on Biopharma, Plasma Procurement, Diagnostics and Bio Supplies



Optimizing our business:
Divestments and cost
optimization

Streamlined our organizational model

Leveraging sustainability leadership to enhance financial and non-Financial performance



Accelerating Innovation and Optimizing Our Portfolio...

Expanding our innovation footprint...

13 R&D sites and +1,100 employees Dreieich, Germany Biopharma & Biotest 4 Hub SF Area. CA **Dublin** Emeryville, Düdingen, Switzerland San Carlos Biopharma Diagnostic LA. San Diego & South San 4 Hub Europe 4 Hub RTP, NC-Francisco, CA Biopharma RTP. NC and Diagnostic Andorra **Biopharma Immunology** Barcelona, Bilbao and Zaragoza, Spain Biopharma and Diagnostic R&D sites Innovation Hub



R&D collaboration agreement

... while prioritizing on our R&D portfolio

Re-prioritized R&D projects

8

projects discontinued

~€30M

savings



Accelerated and focus on priority projects



Reinforced by Biotest R&D pipeline



... With Strong Pipeline Enhanced by Grifols' Scalability and Broader Commercial Reach

Accelerated priority projects... **Market potential GRIFOLS** Xembify® in SID (CLL) ~USD 1Bn **ATIII in Sepsis** ~ USD 1Bn **AMBAR-Next** USD 1Bn+ **GigaGen Platform** USD 1Bn+



CMV IG Hyper



Additional proteins to boost revenue per liter and expand margins

USD 200-300M

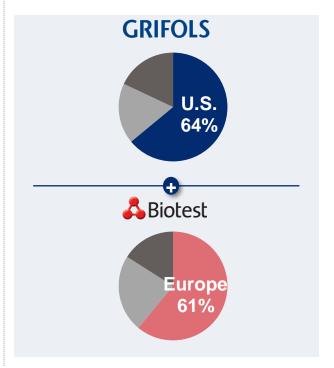


Biopharma | Strong Business Fundamentals Support Biotest Acquisition

Diversified donor center and manufacturing footprint

Europe North America Biotest 313 donor centers 89 donor centers +31 donor centers 3 manufacturing sites 3 manufacturing sites +1 manuf. site **MEA** 1 donor center (+19 planned for 2022 & 2023) 400+ donor centers and 1 manufacturing site 7 manufacturing sites Manufacturing site

Geographic fit



Note: number of plasma centers as of June 30, 2022



Diagnostic | Securing U.S. NAT Business While Untapping China Opportunity Through SRAAS Agreement

CTS agreement



Long-term agreement:



- CTS will operate Grifols testing labs
- Grifols will supply NAT instrumentation and testing



- √ Long-term NAT supply agreement
- ✓ Increasing efficiency of Grifols labs leveraging on CTS expertise
- ✓ Building a **long-term partnership** with largest lab in the world



Distribution agreement through SRAAS



- Integrated commercial model combining Grifols' heritage with SRAAS commercial expertise and broad reach
- Leverage SRAAS branding
- Fastest IVD market worldwide with untapped market potential
- Value chain consolidation with the streamlining of all operations from importation to distribution

Fuel immunohematology (IH) business line to achieve leadership position





Bio Supplies | High Growth and Profitability Business With the Full Integration of Access Biologicals

Bio Supplies

- Leverages discarded products from Biopharma
- Finds niche markets

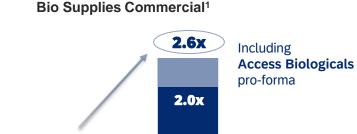
Access Biologicals

Vertical integration to obtain **higher margin** from Grifols products

Commercial knowledge to grow in cell culture market

Enhanced Bio Supplies portfolio

High growth engine for the future



2021

Key customers: Biotechnology research & manufacturing and Diagnostic industries

¹ Bio Supplies Commercial includes sales of biological products for non-therapeutic use

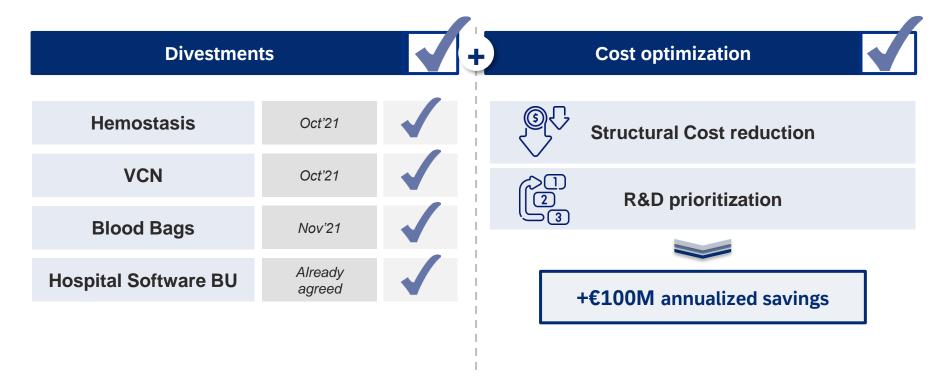
2018

EBIT 35-40%





Optimizing Our Business | Focusing On Value-Creation Businesses While Achieving Structural Cost Savings

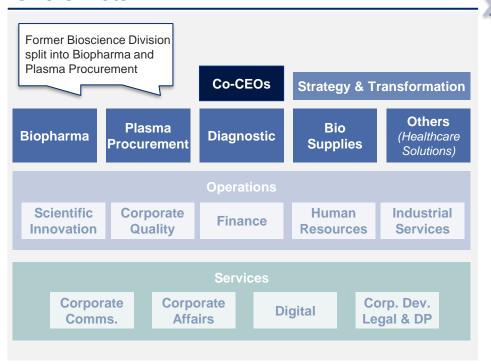




Evolving Our Organization to Reinforce Competitiveness

Streamlined Value-Driven Business Units to Increase Effectiveness and Efficiencies

One Grifols



Building Principles

- 1 Full-fledged Business Units
- 2 Streamlined value-driven organization
- 3 End-to-end business ownership
- Enhanced effectiveness and operational efficiencies
- ✓ Stronger governance model
- ✓ Accountability over execution
- Cutting down on the complexity
- Key support functions provide speed & agility
- ✓ Reduce time-to-market reaction

Delivering on Commitments Outlined in 2020 and 2021

Key Takeaways

Business value

Continuous growth and value creation

CAGR: +8.1% (at cc)

Stable Cost per Liter

while increasing plasma supply

EBITDA: 28%

Delivered on our commitments:

- Strengthened the core:
- Gained focus on Biopharma, Plasma Procurement, Diagnostics & Bio Supplies
- Divestments & Cost Optimization
- Prioritized & reinforced our innovation pipeline
- Streamlined organization

Further performance fueled by delivering on innovation backed by strong fundamentals

Sound business revenue **growth** & sequential gradual margin **expansion**

Supported by strong underlying demand, pricing environment and product/geographical mix coupled with plasma volume efficiencies and cost savings

Build on robust core: 7 manufacturing plants 400+ plasma centers

Harvest innovation efforts

New plasma proteins and indications as well as therapies beyond plasma





- Alpha-1 antitrypsin deficiency (AATD): Inherited disease characterized by low levels of, or no,alpha-1 antitrypsin (AAT) in the blood. This protein made in the liver, reaches other organs (such as the lungs), after being released into the blood stream, enabling its normal function
- Albumin: The most abundant protein found in plasma (approximately 60% of human plasma). Produced in the liver, it is important in regulating blood volume by maintaining the oncotic pressure of the blood compartment
- Alzheimer's disease: This is the most common form of dementia. This incurable, degenerative, and terminal disease was first described by German psychiatrist and neuropathologist Alois Alzheimer in 1906 and was named after him
- Anti-thymocyte globulin (ATG): blood serum that contains antibodies that bind with human T cells. It is given to the patient before a stem cell transplant to
 destroy T cells and decrease the risk of graft-versus-host disease
- ASFA: American Society for Apheresis An organization of physicians, scientists, and allied health professionals whose mission is to advance apheresis medicine for patients, donors, and professionals through education, evidence-based practice, research, and advocacy
- Autoimmune disease: Condition in which the immune system mistakenly attacks healthy cells
- Babesiosis/Babesia virus: Disease caused by microscopic parasites that infect red blood cells
- Beta-amyloid: Protein strongly implicated in Alzheimer's diseases. Beta-amyloid is the main component of certain deposits found in the brains of patients of Alzheimer's disease
- Bullous pemphigoid: is an autoimmune disease that appears when the immune system attacks the skin and causes blisters, more common in the elderly
- CIDP (Chronic Inflammatory Demyelinating Polyneuropathy) Neurological disorder which causes gradual weakness, numbness, pain in arms and legs and difficulty in walking
- Cirrhosis: Medical condition which is a result of advanced liver disease. It is characterized by there placement of liver tissue by fibrosis (scar tissue) and regenerative nodules (lumps that occur due to attempted repair of damaged tissue)
- · Cognitive impairment: Alterations in thinking, learning, memory, judgment, and decision making
- COVID-19: Infectious disease caused by a new strain of coronavirus. 'CO' stands for corona, 'VI' for virus, and 'D' for disease
- ELISA: Enzyme-linked immunosorbent assay
- EMA: European Medicines Agency



- Factor VIII or FVIII: This is an essential blood clotting factor also known as anti-hemophilic factor (AHF). In humans, Factor VIII is encoded by the F8 gene. Defects in this gene results in hemophilia A, a gender linked disease that occurs predominantly in males. FVIII concentrated from donated blood plasma, or alternatively recombinant FVIII, or rFVIII can be given to hemophiliacs to restore hemostasis
- Factor IX: This is an important blood clotting factor also known as Christmas factor or plasma thromboplastin component (PTC). It is one of the serine proteases of the coagulation system and belongs to the peptidase family S1. In humans, a deficiency of this protein causes hemophilia B, a sex-linked disease that occurs predominantly in males
- FDA: Food and Drug Administration. U.S. Health Authority
- Fibrin sealant: Surgical adhesive material derived from plasma
- Fibrinogen: Coagulation factor found in human plasma crucial for blood clot formation
- Fractionation: Process of separating plasma into its component parts, such as albumin, immunoglobulin, alpha-1 antitrypsin and coagulation factors
- · GMP: Good manufacturing practice
- · GPO: Group Purchasing Organization
- HAE (Hereditary Angioedema): Rare but serious genetic disorder characterized by recurrent episodes of severe swelling (angioedema), particularly of the face and airways, and abdominal cramping. It is caused by low levels or improper function of the C1- esterase inhibitor protein
- · HBV: Hepatitis B Virus
- HCV: Hepatitis C Virus
- · Hematocrit: value that is defined by the amount of blood volume occupied by red blood cells, with respect to that occupied by total blood
- Hematology: The study of blood, blood forming organs, and blood diseases
- · Hemoderivative: Proteins obtained by fractionation of human blood plasma. See plasma derived proteins
- Hemophilia: Genetic deficiency characterized by the lack of one of the clotting factors. It has two main variants:
 - Hemophilia A: genetic deficiency of coagulation Factor VIII, which causes increased bleeding (usually affects males)
 - Hemophilia B: genetic deficiency of coagulation Factor IX



- · Hemotherapy: Treatment of a disease using blood, blood components and its derivatives
- · HIV: Human Immunodeficiency Virus
- Hyperimmune globulins: type of immunoglobulins prepared in a manner similar to human normal immunoglobulin, except that the donor has high titers of antibodies against an organism or antigen in their plasma
- IA: Immunoassays. These are systems available in several formats that may be used to detect antibodies, recombinant proteins or a combination of the two
- Intravenous: Administration of drugs or fluids directly into a vein
- Immunohematology: A branch of hematology related to the study of recombinant proteins and antibodies and their effects on blood and the relationships between blood disorders and the immune system. Also referred to as Transfusional Medicine blood bank, its main activities include blood typing, compatibility tests and crossmatching and antibody identification
- **Immunology:** This is a branch of biomedical science that covers the study of all aspects of the immune system in organisms. It deals with the physiological functioning of the immune system in states of both health and disease; malfunctions (autoimmune diseases, hypersensitivities, immune deficiency, transplant rejection) and the physical, chemical and physiological characteristics of the components of the immune system in vitro, in situ, and in vivo
- Immunoglobulin (IgG): Also known as antibodies, are proteins derived from plasma. They control de body's immune response. They have multiple indications and some of their main uses are to treat: (i) immune deficiencies, (ii) inflammatory and autoimmune diseases and (iii) acute infections. IVIG is an immunoglobulin administered intravenously that contains IgG (immunoglobulin (antibody)G)
- ITP (Chronic immune thrombocytopenia): Autoimmune disorder in which patients produce antiplatelet autoantibodies and specialized white blood cells that destroy their blood platelets. This results in a low blood platelet count (thrombocytopenia) that may produce bruising or excessive bleeding
- IVD: In vitro Diagnostic
- IV solutions/Intravenous solution: Medicine or homogeneous mixture of a substance in liquid, enabling it to be infusedinto the circulatory system through a needle
- Lipemic plasma: plasma with a cloudy and/or milky appearance, caused by excess lipids (hyperlipidemia) mainly cholesterol and/or triglycerides in the blood, which in some cases becomes evident
- MRB: Marketing Research Bureau



- Molecular Diagnostic: Discipline that studies genomic (DNA) and proteomic (proteins)expression patterns and uses the information to distinguish between normal, precancerous, and canceroustissues at the molecular level
- Monoclonal antibody (mAb): Antibody produced by a single clone of cells typically used in immunotherapy (such as in the treatment of autoimmune or inflammatory disorders and cancer), diagnostic testing and cell identification and tracking. Monoclonal antibodies are a cornerstone of immunology and are increasingly coming into use as therapeutic agents
- Myasthenia Gravis (MG): Chronic autoimmune, neuromuscular disease that causes weakness in the skeletal muscles that worsens after periods of activity and improves after periods of rest. These muscles are responsible for functions involving breathing and moving parts of the body
- NAT: Nucleic Acid Amplification Testing
- Neurology: Science that deals with the anatomy, functions and organic disorders of nerves and the nervous system
- Northamerica: includes the U.S. and Canada
- · Ophthalmology: branch of medicine and surgery that deals with the diagnosis and treatment of eye diseases
- Pandemic: The worldwide spread of a new disease
- · Parkinson's Disease: complex neurodegenerative disorder in which each patient experiences a different combination of motor and non-motor symptoms
- PCR: Polymerase chain reaction is a method widely used to rapidly make millions to billions of copies of a specific DNA sample, allowing scientists to take a very small sample of DNA and amplify it to a large enough amount to study in detail
- pdFVIII: Plasma-derived Factor VIII
- Pharmacovigilance: Practice of monitoring the effects of medical drugs after they have been licensed for use, especially in order to identify and evaluate previously unreported adverse reactions
- Plasma: Yellow-coloured liquid part of the blood, consisting of a mix of a large number of proteins in solution
- Plasma-derived proteins: Purified plasma proteins with therapeutic properties that are obtained through the fractionation of human plasma. Albumin, immunoglobulins, factor VIII and alpha-1 antitrypsin are the main plasma proteins
- Plasma proteomic: describes the high throughput analysis of plasma biomarkers using very powerful, sensitive and specific instruments

- Plasmapheresis: Plasmapheresis is a technique which separates plasma from other blood components, such as red blood cells, platelets and other cells. These unused blood components are suspended in saline solution and immediately reinjected back into the donor. Because the donor is only providing plasma and not whole blood, the recovery process is faster and better tolerated and the donor is able to make donations more frequently. Plasmapheresis was developed by Jose Antonio Grifols Lucas in the year 1951. It is the only procedure that is capable of obtaining sufficient quantities of plasma to cover the manufacturing needs for the different plasma protein therapies
- Pneumology: is the specialty that takes care of the diagnosis and treatment of respiratory diseases Pulmunologists treat everything related to the respiratory system from asthma to tuberculosis
- PPTA: Plasma Protein Therapeutics Association
- Primary arthroplasty: Surgery performed to replace damaged joints for various reasons, such as hip fractures, osteoarthritis or other rheumatic diseases, by artificial joints called prostheses
- **Primary immunodeficiency (PI):** Inherited condition where there is an impaired immune response, weakening the immune system and allowing infections and other health problems to occur more easily. It may be in one or more aspects of the immune system
- **ProlastinR/ProlastinR -C:** This is a concentrated form of alpha-1 antitrypsin (AAT), derived from human plasma and approved only for chronic, or ongoing, replacement therapy in people with genetic AAT deficiency. Given as prescribed, Prolastin raises the levels of AAT in the blood and lungs. Raising the AAT level may help reduce the damage to the lungs caused by destructive enzymes
- **Proteome:** set of proteins that an organism synthesizes from the genes it contains to give the cell its individual character. This set of proteins determines what organisms are like, how their bodies work and how they behave
- Recombinant: Protein prepared by recombinant technology, coded by the manipulated gene. Procedures are used to join together segments in a cell-free system (an environment outside a cell organism). They are known as highly potent medicines that are safe from off-target side effects and take a shorter time to develop than small molecules
- Recovered plasma: plasma derived from whole blood collected in blood donations
- **rFVIII:** Recombinant Factor VIII is the antihemophilic factor A, obtained using recombinant DNA technology. With this technology, pure factor is synthesized in the laboratory instead of being extracted from blood plasma

- Rh (Rhesus) blood group system: Most important blood group system after ABO. The Rh blood group system consists of 50 defined blood-group recombinant proteins, among which the five recombinant proteins D,C, c, E and are the most important. The commonly used terms Rh factor, Rh positive and Rh negative refer to the D antigen only
- · ROW: Rest of the World
- SARS-CoV-2: Severe acute respiratory syndrome coronavirus 2 is the strain of coronavirus that causes coronavirus disease 2019 (COVID-19), the respiratory illness responsible for the COVID-19 pandemic
- Secondary immunodeficiency: Occurs when the immune system is compromised due to an environmental factor. Examples of these outside forces include HIV, chemotherapy, severe burns or malnutrition
- · SCIG: subcutaneous immunoglobulin
- Single-cell transcriptomics: technique to characterize cell identity
- SCIG/SubQ: Sub-cutaneous
- · Thrombin: Enzyme that presides over the conversion of a substance called fibrinogen to fibrin, which promotes blood clotting
- Transfusion medicine: Branch of medicine that encompasses among others, immunohematology, blood and plasma screening and blood typing
- West Nile Virus (WNV): Virus transmitted by mosquitoes. Humans are mainly infected through mosquito bites, but infection can occur through organ transplantation and blood
- Von Willebrand Disease (vWD): This is the most common hereditary coagulation abnormality described in humans, although it can also be acquired as a result of other medical conditions. It arises from a qualitative or quantitative deficiency of von Willebrand factor (vWF), a multimeric protein that is required for platelet adhesion
- · Zika virus: Infectious disease spread by the bite of an infected Aedes species mosquito