GRIFOLS

Half-Year 2025 Results

Grifols increases its revenues by 7% to EUR 3,677m and boosts its net profit to EUR 177m

- Revenues reached EUR 3,677 million up 7.0% at constant currency (cc¹), driven by the solid performance of the Biopharma segment, which recorded 8.2% growth at constant currency
- Adjusted EBITDA increased to EUR 876 million, which represents a 12.7% cc growth and 23.8% margin
- Group net profit increased to EUR 177 million, close to 4x the figure reported in H1 2024
- Free cash flow² before M&A improved by EUR 182 million year on year
- Leverage ratio³ decreased to 4.2x, with liquidity standing at EUR 1.4 billion⁴
- On track to launch Fibrinogen in Europe in the fourth quarter of 2025 and in the first half of 2026 in the U.S., following FDA approval
- Reinstated a dividend payment of EUR 0.15 per share, reflecting the company's commitment to shareholder returns
- Reaffirmed guidance for 2025⁵, with improved guidance for FCF pre-M&A to EUR 375-425 million

Barcelona, Spain – July 29, 2025 – Grifols (MCE:GRF, MCE:GRF.P, NASDAQ:GRFS), a global healthcare company and leader in plasma-derived medicines, today announced results for the first half of 2025, driven by a second quarter marked by continued improvements across key operational and financial metrics. These results reflect the ongoing execution of Grifols' Value Creation Plan.

Revenues for the first half of the year grew by 7.0% cc to EUR 3,677 million, driven by the performance of the Biopharma business, which increased by 8.2% cc. Adjusted EBITDA reached EUR 876 million, up by 12.7% cc year-over-year, representing a 23.8% margin, supported by product mix, continuous improvement initiatives and operational leverage. Net profit surged to EUR 177 million, reflecting a 387.6% increase compared to the same period of 2024.

Free cash flow pre-M&A significantly improved to positive EUR 30 million in the second quarter, resulting in minus EUR 14 million for the first half of the year. This represents a EUR 182 million year-over-year improvement, mainly driven by EBITDA growth, working capital management and reduced interest costs.

Grifols further strengthened its financial position, reducing its leverage ratio to 4.2x, down from 4.5x in the previous quarter and 5.5x in the first half of 2024, with a liquidity position of EUR 1.4 billion. The company remains focused on continuing to improve its credit profile.

¹ Operating or constant currency (cc) excludes changes rate variations reported in the period.

² Free Cash Flow includes cash from operating activities + cash flow from investing activities, both as per International Financial Reporting Standards (IFRS), and excludes lease payments.

³ Defined as per the Credit Agreement.

⁴ Cash and cash equivalents of €559m + unused credit facilities €1,251m - unused RCF facilities maturing in Nov 2025 c€396m.

⁵ Please refer to 2025 Guidance on page 38 of the Capital Markets Day Presentation (27 Feb 2025).

As part of its capital allocation framework, Grifols successfully completed the delisting of Biotest from the Frankfurt Stock Exchange and increased its equity stake to 80.32%. This transaction, with a total cost of EUR 108 million, was fully funded through available financial resources.

The company declared a EUR 0.15 per share dividend payment supported by continued underlying earnings and free cash flow generation momentum. This reflects the company's strong commitment to shareholder returns.

Nacho Abia, CEO of Grifols, said: "The company's strong performance in the first six months of 2025 reflects the solid execution of our Value Creation Plan. The momentum of the business is clear: in a context of strong underlying demand, we continue to capitalize on the strength of our Biopharma business unit while advancing on key priorities."

Nacho Abia added: "While the value creation will ultimately benefit our shareholders, we continue to be fully committed to developing solutions that address patients' needs - a priority that has defined Grifols for more than 116 years."

Rahul Srinivasan, CFO of Grifols, said: "The company's strong first-half performance underscores both the attractive fundamentals and resilience of our business. We remain resolutely focused on leveraging the strengths of our business model and disciplined execution, capitalizing on our underlying momentum and operational focus to deliver on our deleveraging and free cash-flow generation priorities, whilst reinforcing the strong confidence in the company's long-term value."

On the exposure and potential impact of trade tariffs, the company is well-prepared, thanks to a locally operated and vertically integrated business model. Since its international expansion over 30 years ago, Grifols has invested consistently in a global network of donation, processing, and distribution centers for plasma-derived medicines, allowing it to operate locally in the United States, Europe, Egypt, and Canada. The integration minimizes the need for imports and/or exports within the U.S. market and also strengthens its flexible and resilient structure in the face of regulatory changes or new tariffs.

Biopharma drives strong growth in the first half of 2025

Biopharma revenue grew by 8.2% cc, led by continued momentum in the immunoglobulin (IG) franchise, while Alpha-1 continued to gain traction. Specifically, IG revenues rose by 12.5% cc, with strong growth in both its intravenous form (IVIG), which outpaced market growth, and subcutaneous form (SCIG), which delivered 66% cc in the first part of 2025. Grifols continues to consolidate its leadership in key indications such as primary and secondary immunodeficiencies and CIDP.

Albumin posted sequential improvement in Q2 2025, following license renewal in China. The 1.3% cc growth in this quarter resulted in a first half performance of minus 3.7% cc.

Revenues from Alpha-1 and Specialty proteins continued to perform positively, growing 4.8% cc in the first half of the year. Alpha-1 continues to benefit from the company's leading 70% global market share. As part of this protein's commercial growth strategy, the SPARTA study and the subcutaneous formulation trial for Alpha-1 are progressing as planned.

Grifols maintains its plan to launch fibrinogen in Europe in the fourth quarter of 2025 and in the United States in the first half of 2026, following FDA approval.



Diagnostic business maintains positive momentum

Diagnostic revenues reached EUR 332 million, an increase of 2.8% cc, driven by Molecular Donor Screening (MDS) across EMEA and Asia-Pacific, as well as solid performance in Blood Typing Solutions (BTS) in key countries. Noteworthy is the FDA approval to begin manufacturing Gel Cards and reagent Red Blood Cells at Grifols' San Diego facility.

Grifols reaffirms 2025 guidance and improves the Free Cash Flow pre-M&A guidance

As Grifols enters the second half of 2025 with strong underlying momentum across its core businesses, the company reaffirms its 2025 guidance shared during its Capital Markets Day on February 27, 2025, and improves the guidance for Free Cash Flow pre-M&A guidance to EUR 375-425 million.

While the recent depreciation of the U.S. dollar presents a headwind to reported Revenue and EBITDA in the second half, the impact on Group Profit, leverage and Free Cash Flow pre-M&A is expected to be broadly neutral. The Group expects to largely mitigate this headwind through a combination of the benefit of underlying business momentum and targeted cost levers.

	1		Reported	d		Like for Like ¹	H1 2024
(in million EUR except	: %)	Q1'25	Q2'25	H1'25	Var vs. PY	Var vs. PY	Reported
NET REVENUE		1,786m	1,891m	3,677m	7.0% cc	10.1% cc	3,444m
GROSS MARGIN		695m	744m	1,438m	11.6% cc	19.8% cc	1,301m
▶ Margin		38.9%	39.3%	39.1%	+30bps	+300bps	37.8%
EBITDA ADJ.		400m	475m	876m	12.7% cc	20.1% сс	791m
▶ Margin		22.4%	25.1%	23.8%	+80bps	+171bps	23.0%
PROFIT BEFORE TAX	(115m	191m	306m	168.1%		114m
GROUP PROFIT		60m	117m	177m	387.6%		36m
FREE CASH FLOW p	re-M&A²	-44m	30m	-14m	182m		-196m
LEVERAGE RATIO ³	Total net LR	4.5x		4.2x	-1.3x		5.5x
LEVERAGE RATIO	Net secured LR	2.7x		2.7x	-0.8x		3.4x
LIQUIDITY		1,675m		1,414m ⁴	499m		915m ⁵

Note: All figures are presented on a consolidated basis (including Biotest). When specified, figures presented at currency (cc), excluding exchange rate fluctuations over the period. See Annex for reconciliations.

2 FCF definition and reconciliation to the Cash Flow Statement in slide 34 in the Annex.

3 Everage ratio defined as per the Credit Agreement in slide 31 in the Annex.

4 FCF H 2025, cash and cash equivalents of 6559m + unused credit facilities 61.251m - unused RCF facilities maturing in Nov 2025 c6396m.

5 Liquidity position excluding EUR 1.6bn SRAAS proceeds

Alternative Performance Measures (APMs)

This document contains the following Alternative Performance Measures (APMs): Consolidated EBITDA Reported, Consolidated EBITDA Adjusted, Leverage Ratio as per the Credit Facility, Net Debt as per the Credit Facility, Free Cash Flow, Working Capital, and non-recurring items. For further details on the definition, explanation on the use, and reconciliation of APMs, please see the Appendix of the Presentation as well as the "Alternative Performance Measures" document from Grifols website www.grifols.com/en/investors.

CONFERENCE CALL

Grifols will host a conference call today, July 29, 2025, at 6:30pm CET/12:30pm EST to provide review of the company's business results for the second quarter of 2025. To view and listen to the webcast and view the presentation, click on Q2 2025 Results or visit the website www.grifols.com/en/investors. Participants are advised to register in advance of the conference call.

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About Grifols

Grifols is a global healthcare company founded in Barcelona in 1909 committed to improving the health and well-being of people around the world. A leader in essential plasma-derived medicines and transfusion medicine, the company develops, produces, and provides innovative healthcare services and solutions in more than 110 countries.

Patient needs and Grifols' ever-growing knowledge of many chronic, rare and prevalent conditions, at times life-threatening, drive the company's innovation in both plasma and other biopharmaceuticals to enhance quality of life. Grifols is focused on treating conditions across four main therapeutic areas: immunology, infectious diseases, pulmonology and critical care.

A pioneer in the plasma industry, Grifols continues to grow its network of donation centers, the world's largest with close to 400 across North America, Europe, Africa and the Middle East, and China.

As a recognized leader in transfusion medicine, Grifols offers a comprehensive portfolio of solutions designed to enhance safety from donation to transfusion, in addition to clinical diagnostic technologies. It provides high-quality biological supplies for life-science research, clinical trials, and for manufacturing pharmaceutical and diagnostic products. The company also supplies tools, information and services that enable hospitals, pharmacies and healthcare professionals to efficiently deliver expert medical care.

Grifols, with more than 23,800 employees in more than 30 countries and regions, is committed to a sustainable business model that sets the standard for continuous innovation, quality, safety, and ethical leadership.

In 2023, Grifols' economic impact in its core countries of operation was EUR 9.6 billion. The company also generated 193,000 jobs, including indirect and induced.

The company's class A shares are listed on the Spanish Stock Exchange, where they are part of the Ibex-35 (MCE:GRF). Grifols non-voting class B shares are listed on the Mercado Continuo (MCE:GRF.P) and on the U.S. NASDAQ through ADRs (NASDAQ:GRFS). For more information about Grifols, please visit www.grifols.com.

Legal Disclaimer

The facts and figures contained in this report that do not refer to historical data are 'projections and future hypotheses'. Words and expressions such as 'believe', 'expect', 'anticipate', 'predict', 'hope', 'intend', 'should', 'will try to achieve', 'is estimated', 'future' and similar expressions, insofar as they refer to the Grifols group, are used to identify future projections and hypotheses. These expressions reflect the assumptions, hypotheses, expectations and predictions of the management team at the time of writing this report, and these are subject to a series of factors that mean that the real results may be materially different. The future results of the Grifols group could be affected by events related to its own activities, such as shortages of supplies of raw materials for the manufacture of its products, the appearance on the market of competing products, or changes in the regulatory framework of the markets in which it operates, among others. At the date of preparation of this report, the Grifols group has adopted the necessary

GRIFOLS

measures to mitigate the potential impact of these events. Grifols, S.A. assumes no obligation to publicly report, revise or update the projections or future hypotheses to adapt them to facts or circumstances after the date of writing of this report, except when expressly required by applicable legislation. This document does not constitute an offer or invitation to purchase or subscribe shares in accordance with the provisions of Law 6/2023, of 17 March, on the Securities Markets and Investment Services, and any regulations implementing said legislation. Furthermore, this document does not constitute an offer to purchase, sell or exchange, or a solicitation of an offer to purchase, sell or exchange any securities, or a solicitation of any vote or approval in any other jurisdiction. The information contained in this document has not been verified or revised by the external auditors of the Grifols group.



Legal Disclaimer

Important Information

This presentation does not constitute an offer or invitation to purchase or subscribe shares, in accordance with the provisions of the Regulation (EU) 2017/1129 of the European Parliament and of the Council of 14 June 2017 on the prospectus to be published when securities are offered to the public or admitted to trading on a regulated market, and repealing Directive 2003/71/EC, the Spanish Securities Market and Investment Services Law (Law 6/2023, of 17 March, as amended and restated from time to time) and its implementing regulations. In addition, this document does not constitute an offer of purchase, sale or exchange, nor a request for any vote or approval in any other jurisdiction. This information has not been audited.

Forward-Looking Statements

This presentation 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.

Alternative Performance Measures (APMs)

This document and any related conference call or webcast (including a Q&A session) contain, in addition to the financial information prepared in accordance with IFRS, alternative performance measures ('APMs') as defined in the guidelines issued by the European Securities and Markets Authority ('ESMA') on October 5, 2015. APMs are used by Grifols' management to evaluate the group's financial performance, cash flows or financial position in making operational and strategic decisions for the group and therefore are useful information for investors and other stakeholders. Certain key APMs form part of executive directors, management and employees' remuneration targets.

APMs are prepared on a consistent basis for the periods presented in this document. They should be considered in addition to IFRS measurements, may differ to definitions given by regulatory bodies relevant to the group and to similarly titled measures presented by other companies. They have not been audited, reviewed or verified by the external auditor of Grifols. For further details on the definition, explanation on the use, and reconciliation of APMs, please see the appendix as well as the "Alternative performance measures" document from our website www.grifols.com/en/investors.

Agenda

01 Business Review

02 Financials

03 Final Remarks

04 Annex







Rahul Srinivasan
Chief Financial Officer
(CFO)

Q2'25 Results

Strong Performance Driven by A Clear Focus on Execution

Nacho Abia

Chief Executive Officer (CEO)

Continued Execution in Q2 Supporting Strong First-Half Performance

H1'25

€3,677m

Revenue

€876m EBITDA Adj.

-€14m

Free Cash Flow (pre-M&A)³

4.2X Leverage ratio⁴

H1'25 vs H1'24

+7.0%cc¹ (+10.1%cc LFL²)

+12.7%cc (+20.1%cc LFL)

+€182m

-0.4x

- Executing on our Value Creation Plan driving strong performance across all key financial metrics
- Strong business momentum on the back of robust underlying demand, even after absorbing the IRA impact.
 Structurally well-positioned to navigate uncertainty from potential tariffs impact
- ► Continuous improvement initiatives supporting margin expansion
- Generating free cash flow and deleveraging
- Delivering on innovation milestones
- Executing on capital allocation framework including the reinstatement of dividends with €0.15 per share interim dividend

All figures are presented on a consolidated basis

¹ Constant currency (cc), excluding exchange rate fluctuations over the period. See Annex for reconciliations.;

² Like For Like (LFL) excludes the impact of Inflation Reduction Act (IRA) and Fee-For-Service / GPO reclassification.

³ FCF definition and reconciliation to the Cash Flow Statement in slide 34 in the Annex.

⁴ Leverage ratio defined as per the Credit Agreement in slide 31 in the Annex.

Value Creation Plan Execution Delivering Results

Commercial growth

Levers

Leveraging strong momentum of key proteins

+8.2% cc

Biopharma growth (H1'25 vs H1'24)

Margin expansion (even despite IRA impact)

Driving efficiencies through plasma and mfg. yield improvements

+80bps (**+171bps LFL**)
EBITDA Adj margin (H1'25 vs H1'24)

Pipeline execution

Delivering on our innovation milestones

Q4'25

Fibrinogen EU launch

Plasma supply & industrial footprint

Enablers

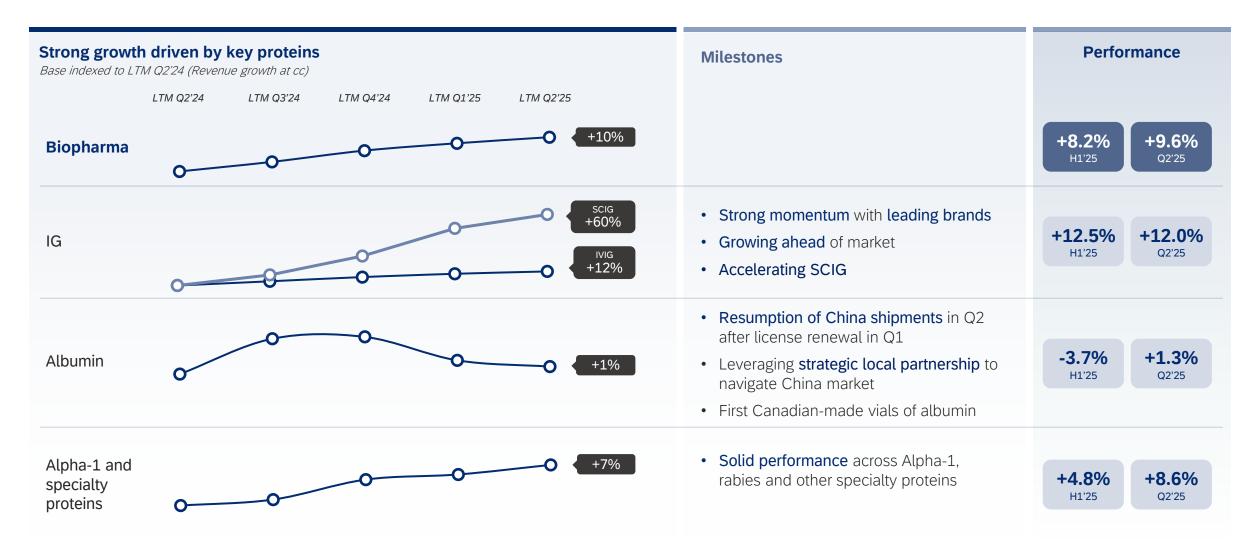
Vertically integrated value chain in the US and Europe, complemented by strategic hubs and partnerships ex-US, providing valuable optionality and demonstrating resilience amidst a dynamic market backdrop

Innovation

Driving clinical trials traction across life cycle management (LCM) and new products and indications for core proteins



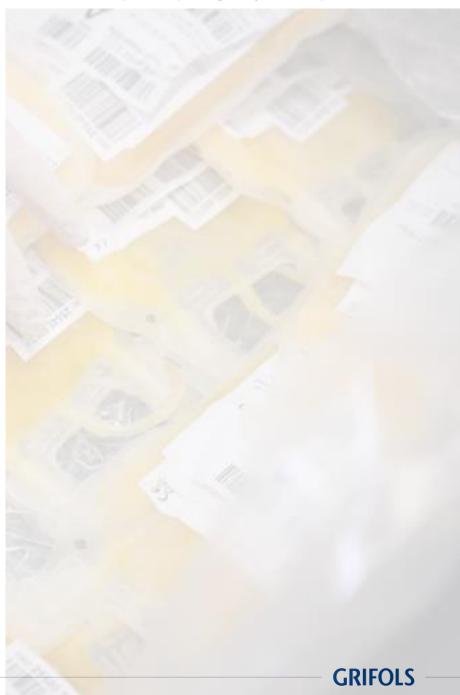
Commercial Execution Fuels Biopharma Performance Offsetting IRA



Note: All figures are presented on a consolidated basis, and at constant currency (cc), excluding exchange rate fluctuations over the period.



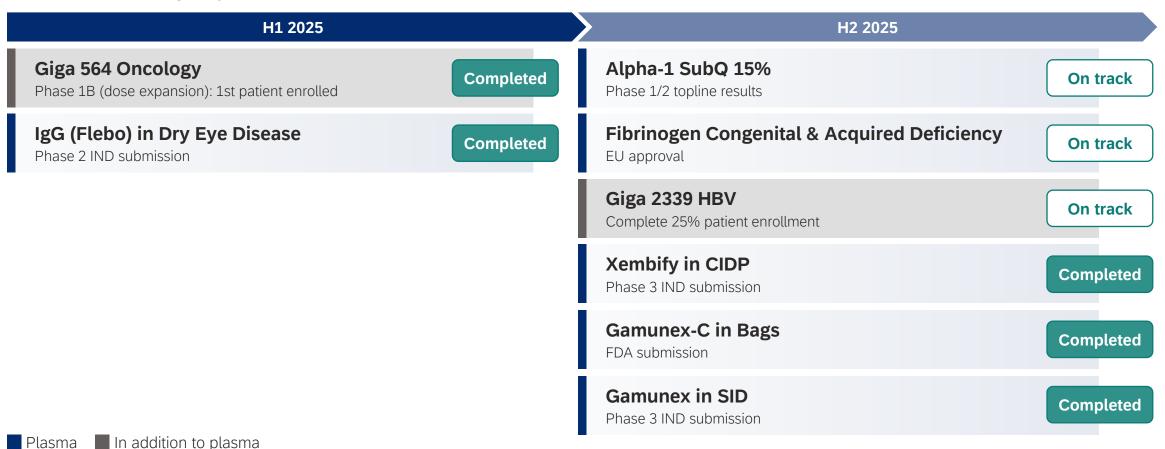
- Achieved 60% individualized nomogram implementation in U.S. centers
- ► Kicked-off 2nd nomogram wave to reach 100% adoption by 2026
- ► On track to deliver double-digit growth in volume per center
- ► Higher IVIG manufacturing yield from operational excellence
- Ongoing efficiency improvements resulting in reduction in donation time
- Continuous optimization of plasma supply



Q2 2025 Results - 8 -

Delivering on our Pipeline Milestones with New Indications and Progress in Life Cycle Management

Innovation milestone targets by ...





Advancing Diagnostic to Reinforce Market Position and Profitability

Strong growth driven by k	key business segments	Milestones	Performance (H1'25)
Diagnostic		 High-margin complementary business Significant EBITDA contribution High cash flow generation 	+2.8% cc ¹
Blood Typing Solutions	 Strengthening presence in core markets Focus on increasing profitability 	 FDA approval to start GelCards and Red Blood Cells' production in the USA San Diego site Absorb continued growth and respond to needs Strategically broaden and diversify footprint 	+7.1% cc
Donor Screening (Molecular)	 Sustaining leadership position Developing Next Generation MDS 	 Strategic alliance with Inpeco Focused on providing full automation solutions Lead-in to develop the "lab of the future" 	+2.2% cc
Donor Screening (Immunoassay)	 Developing new immunoassay technology for blood and plasma screening 	 On track with the ISARD platform development to be launched by 2029-30 	+8.1% cc ²

All figures are presented on a consolidated basis

GRIFOLS - 10 -

Q2 2025 Results

¹ Constant currency (cc), excluding exchange rate fluctuations over the period. See Annex for reconciliations.; ² Positively impacted by the timing of true up of royalties

Strong Q2 Supporting a Record H1 Performance

Rahul Srinivasan

Chief Financial Officer (CFO)

Record H1'25, Continued Strong Momentum

			Reported	d		Like for Like ¹	H1 2024
(in million EUR excep	t %)	Q1'25	Q2'25	H1'25	Var vs. PY	Var vs. PY	Reported
NET REVENUE		1,786m	1,891m	3,677m	7.0% cc	10.1% cc	3,444m
GROSS MARGIN		695m	744m	1,438m	11.6% cc	19.8% cc	1,301m
▶ Margin		38.9%	39.3%	39.1%	+30bps	+300bps	37.8%
EBITDA ADJ.		400m	475m	876m	12.7% cc	20.1% cc	791m
▶ Margin		22.4%	25.1%	23.8%	+80bps	+171bps	23.0%
PROFIT BEFORE TAX	X	115m	191m	306m	168.1%		114m
GROUP PROFIT		60m	117m	177m	387.6%		36m
	_						
FREE CASH FLOW p	re-M&A ²	-44m	30m	-14m	182m		-196m
LEVERAGE RATIO ³	Total net LR	4.5x		4.2x	-1.3x		5.5x
LEVERAGE RATIO	Net secured LR	2.7x		2.7x	-0.8x		3.4x
LIQUIDITY		1,675m		1,414m ⁴	499m		915m ⁵

Note: All figures are presented on a consolidated basis (including Biotest). When specified, figures presented at currency (cc), excluding exchange rate fluctuations over the period. See Annex for reconciliations.



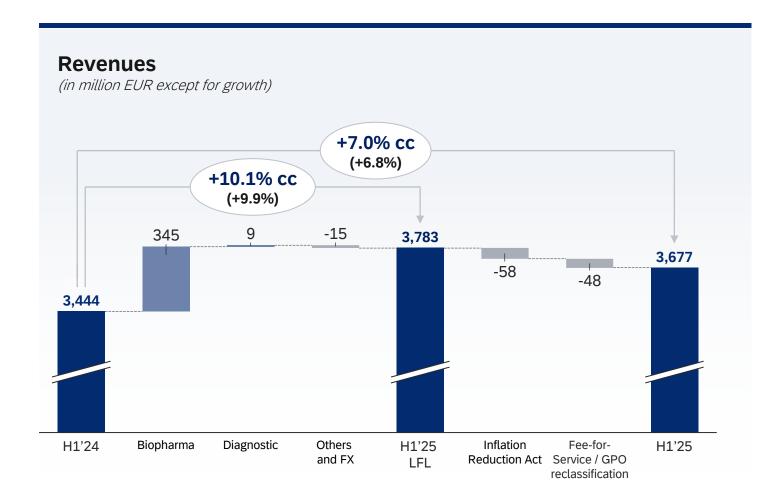
¹ Like For Like (LFL) excludes the impact of IRA and Fee-For-Service / GPO reclassification. ² FCF definition and reconciliation to the Cash Flow Statement in slide 34 in the Annex.

³ Leverage ratio defined as per the Credit Agreement in slide 31 in the Annex.

⁴ For H1 2025, cash and cash equivalents of €559m + unused credit facilities €1,221m - unused RCF facilities maturing in Nov 2025 c€366m.

⁵ Liquidity position excluding EUR 1.6bn SRAAS proceeds

Biopharma Drives +7.0% cc Revenue Growth



Biopharma I +8.2% cc | +11.8% cc LFL¹

 Solid performance backed by strong underlying demand led by IG and Alpha-1's further traction

Diagnostic I +2.8% cc

 Driven by MDS² backed by strong EMEA and APAC sales and BTS² in key countries

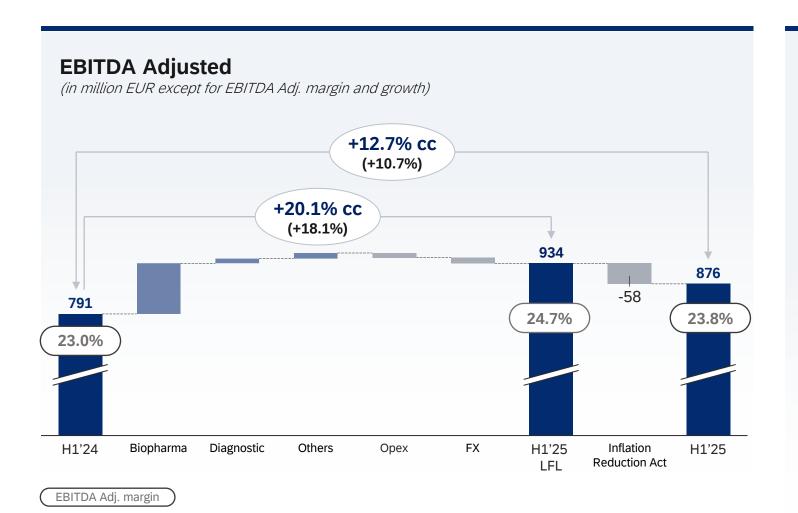
Note: All figures are presented on a consolidated basis (including Biotest). Percentage rates are presented at constant currency (cc), excluding exchange rate fluctuations over the period. See Annex for reconciliations. ¹Like For Like (LFL) excludes the impact of IRA (EUR 58 million) and Fee-For-Service / GPO reclassification (EUR 48 million). See Annex for reconciliations.

² Molecular Donor Screening (MDS); Blood Typing Solutions (BTS).



Q2 2025 Results - 13 -

Significant EBITDA Growth and Margin Expansion Despite IRA Impact



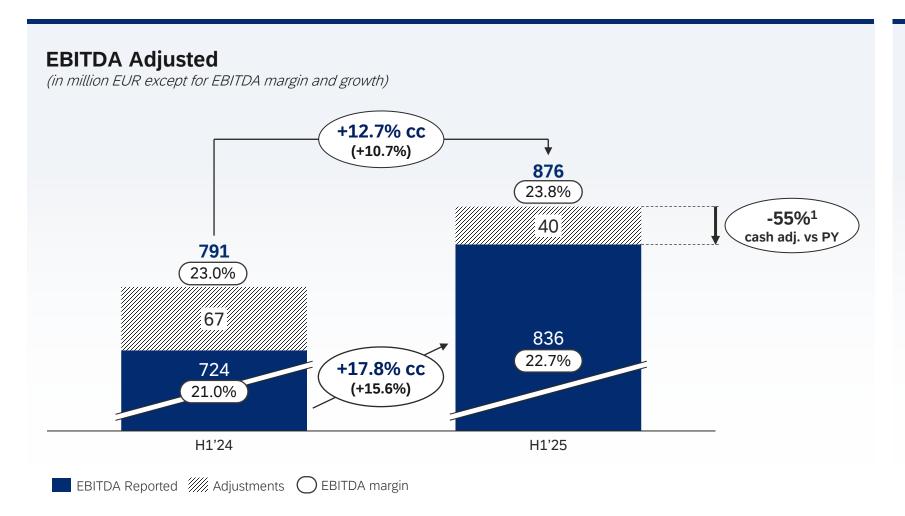
- **Strong EBITDA** growth led by **Biopharma**:
 - Volume growth
 - Cost Per Liter optimization
 - Yield improvements
 - Operational leverage
 - Cost discipline
- IRA impact consistent with expectations
- Rather muted FX impact in H1

Note: All figures are presented on a consolidated basis (including Biotest).

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Q2 2025 Results - 14 -

Lower Cash Adjustments Accelerating EBITDA Convergence



 Continued focus on reducing cash adjustments between EBITDA Adjusted and Reported

- Drivers of reduced cash adjustments:
 - Lower transaction costs
 - Lower restructuring costs



Q2 2025 Results - 15 - -

¹ Cash adjustments include transaction costs, restructuring costs and other non-recurring items as reflected in the reconciliation of slide 30 in the Annex Note: All figures are presented on a consolidated basis (including Biotest)

On Track for Further Improvement in Free Cash Flow Generation

EBITDA Adjusted to Free Cash Flow reconciliation

(in million EUR)

	Q1'25	Q2'25	H1'25	Var vs H1'24
EBITDA Adjusted	400	475	876	84
Inventories	(61)	(30)	(91)	41
Receivables	(93)	(25)	(118)	(28)
Payables	26	25	51	4
Net working capital	(128)	(30)	(158)	17
CAPEX	(128)	(83)	(211)	11
IT and R&D	(39)	(34)	(73)	(10)
Taxes	(3)	(45)	(48)	15
Interests	(55)	(235)	(290)	49
Others	(91)	(19)	(110)	15
Free Cash Flow pre-M&A ¹	(44)	30	(14)	182

- Reduction in cash adjustments to EBITDA Adjusted
- Working capital management
- Deleveraging benefit and lower RCF utilization reducing cash interest vs. H1'24
- ▶ FCF outperformance is despite a more punitive phasing in H1'25 with respect to Capex, IT and R&D and Interest

Note: All figures are presented on a consolidated basis (including Biotest)

GRIFOLS Q2 2025 Results - 16 -

¹ FCF definition and reconciliation to the Cash Flow Statement in slide 34 in the Annex

Capital Allocation Discipline

Balance Sheet Strength

- Continued deleveraging with leverage ratio² at 4.2x vs. 4.6x in Q4'24
- Significant secured capacity (LR at 2.7x)
- No meaningful maturities until Q4'27
- Strong €1.4bn³ liquidity
- Focused on continued credit re-rating progress

Organic Business

- Continued planned investments
- Improvement in Free Cash Flow generation pre-M&A¹
- Confident H2'25 FCF outlook

Inorganic Efforts (Corporate Simplification)

- Successful delisting of Biotest from Frankfurt Stock Exchange
- Biotest progressing as planned

Shareholder Returns

- Consistent with capital allocation framework
- Continued underlying earnings and FCF generation momentum
- Reinstatement of dividends:
 - 0.15 EUR/share interim dividend on August 13th, 2025
- First dividend since June 2021

Note: All figures are presented on a consolidated basis (including Biotest). When specified, figures presented at currency (cc), excluding exchange rate fluctuations over the period. See Annex for reconciliations. ¹ FCF definition and reconciliation to the Cash Flow Statement in slide 34 in the Annex.

Q2 2025 Results GRIFOLS

² Leverage ratio defined as per the Credit Agreement in slide 31 in the Annex.

³ For H1 2025, cash and cash equivalents of €559m + unused credit facilities €1,221m - unused RCF facilities maturing in Nov 2025 c€366m.

FY25 Outlook

- Strong underlying momentum
- Impact of a depreciating USD
 - A headwind for Revenue and EBITDA
 - EBITDA benefiting partially from natural hedge
 - For reference: on average, every 0.01 USD depreciation against EUR for the full year represents a full year EBITDA headwind of approx. EUR 7 million
 - But broadly neutral impact on Group Profit, Leverage and FCF
- Expectation of continued strong underlying momentum and cost levers to help mitigate impact of USD depreciation
- Reaffirmed 2025 guidance, with improved guidance for FCF pre-M&A of €375 - €425m





Final Remarks

Nacho Abia
Chief Evecutive Of

Chief Executive Officer (CEO)

GRIFOLS

A Clear Focus on Execution

- Executing effectively on our Value Creation Plan with proven results
- ▶ Building on targeted strong momentum to reach FY25 guidance
- Expanding margins on the back of efficiency initiatives progress
- Pipeline execution to unlock long-term strategic value
- Improving FCF generation and deleveraging
- Well equipped to manage global market dynamics







Revenue | Q2 2025

	Q2 2025	Q2 2024	% vs F	Υ
In thousands of euros			Reported	At cc*
Revenue by Business Unit	1,891,132	1,817,908	4.0%	6.6%
Biopharma	1,632,629	1,527,652	6.9%	9.6%
Diagnostic	161,588	164,261	(1.6%)	0.4%
Bio Supplies	36,403	73,719	(50.6%)	(49.3%)
Others	60,512	52,276	15.8%	17.2%
	1			1
Revenue by Country	1,891,132	1,817,908	4.0%	6.6%
US + CANADA	1,067,726	1,037,872	2.9%	5.7%
EU	401,252	418,882	(4.2%)	(4.2%)
ROW	422,154	361,154	16.9%	21.4%

^{*} Constant currency (cc) excludes exchange rate fluctuations over the period.



Business Performance

Revenue | H1 2025

	H1 2025	H1 2024	% vs F	PΥ
In thousands of euros			Reported	At cc*
Revenue by Business Unit	3,676,940	3,443,613	6.8%	7.0%
Biopharma	3,153,789	2,922,355	7.9%	8.2%
Diagnostic	331,632	322,544	2.8%	2.8%
Bio Supplies	68,958	100,731	(31.5%)	(31.8%)
Others	122,561	97,983	25.1%	25.0%
Revenue by Country	3,676,940	3,443,613	6.8%	7.0%
US + CANADA	2,092,916	1,963,198	6.6%	6.1%
EU	791,526	750,309	5.5%	5.4%
ROW	792,498	730,106	8.5%	11.0%

^{*} Constant currency (cc) excludes exchange rate fluctuations over the period.



Q2 2025 Results

P&L | Q2 2025

		Q2 2025			Q2 2024		% vs	S PY
In thousands of euros	Reported	One-offs	Reported excl. One-offs	Reported	One-offs	Reported excl. One-offs	Reported	Reported excl. One-offs
Net Revenue	1,891,131	-	1,891,131	1,817,908	-	1,817,908	4.0%	4.0%
Cost of Sales	(1,147,337)	18,883	(1,128,454)	(1,147,464)	15,731	(1,131,733)	0.0%	0.3%
Gross Margin	743,794	18,883	762,677	670,444	15,731	686,175	10.9%	11.1%
% Net revenue	39.3%	-	40.3%	36.9%	-	37.7%	-	-
R&D	(96,193)	-	(96,193)	(90,695)	1,492	(89,203)	(6.1%)	(7.8%)
SG&A	(298,061)	5,025	(293,036)	(327,336)	24,751	(302,585)	8.9%	3.2%
Operating Expenses	(394,254)	5,025	(389,229)	(418,031)	26,243	(391,788)	5.7%	0.7%
Other Income	-	-	-	-	-	-	-	-
Share of Results of Equity Accounted Investees - Core Activities	(686)	-	(686)	46,909	(5,618)	41,291	(101.5%)	(101.7%)
OPERATING RESULT (EBIT)	348,854	23,908	372,762	299,322	36,356	335,678	16.5%	11.0%
% Net revenue	18.4%	-	19.7%	16.5%	-	18.5%	-	-
Financial Result	(158,346)	-	(158,346)	(232,489)	46,763	(185,726)	31.9%	14.7%
Share of Results of Equity Accounted Investees	-	-	-	145	-	145	(100.0%)	(100.0%)
PROFIT BEFORE TAX	190,508	23,908	214,416	66,978	83,119	150,097	184.4%	42.9%
% Net revenue	10.1%	-	11.3%	3.7%	-	8.3%	-	-
Income Tax Expense	(52,128)	(6,466)	(58,594)	(42,214)	9,915	(32,299)	(23.5%)	(81.4%)
% of pre-tax income	27.4%	-	27.3%	63.0%	-	21.5%	-	
CONSOLIDATED PROFIT	138,380	17,442	155,822	24,764	93,034	117,798	458.8%	32.3%
Results Attributable to Non-Controlling Interests	(21,318)	(1,830)	(23,148)	(9,923)	(3,056)	(12,979)	(114.8%)	(78.3%)
GROUP PROFIT	117,062	15,612	132,674	14,841	89,978	104,819	688.8%	26.6%
% Net revenue	6.2%	-	7.0%	0.8%	-	5.8%		

Q2 2025 Results



P&L | H1 2025

		H1 2025			H1 2024		% vs	; PY
In thousands of euros	Reported	One-offs	Reported excl. One-offs	Reported	One-offs	Reported excl. One-offs	Reported	Reported excl. One-offs
Net Revenue	3,676,940	-	3,676,940	3,443,613	-	3,443,613	6.8%	6.8%
Cost of Sales	(2,238,479)	28,852	(2,209,627)	(2,142,324)	33,293	(2,109,031)	(4.5%)	(4.8%)
Gross Margin	1,438,461	28,852	1,467,313	1,301,289	33,293	1,334,581	10.5%	9.9%
% Net revenue	39.1%	-	39.9%	37.8%	-	38.8%		
R&D	(192,239)	-	(192,239)	(181,157)	1,601	(179,556)	(6.1%)	(7.1%)
SG&A	(622,713)	14,308	(608,405)	(666,387)	47,662	(618,725)	6.6%	1.7%
Operating Expenses	(814,952)	14,308	(800,644)	(847,544)	49,263	(798,281)	3.8%	(0.3%)
Other Income	-	-	-	-	-	-	-	-
Share of Results of Equity Accounted Investees - Core Activities	(5,797)	3,850	(1,947)	49,379	(5,618)	43,761	(111.7%)	(104.4%)
OPERATING RESULT (EBIT)	617,712	47,010	664,722	503,123	76,938	580,061	22.8%	14.6%
% Net revenue	16.8%	-	18.1%	14.6%	-	16.8%		
Financial Result	(311,939)	-	(311,939)	(389,089)	46,763	(342,326)	19.8%	8.9%
Share of Results of Equity Accounted Investees	-	-	-	-	-		-	-
PROFIT BEFORE TAX	305,773	47,010	352,783	114,034	123,701	237,736	168.1%	48.4%
% Net revenue	8.3%	-	9.6%	3.3%	-	6.9%		
Income Tax Expense	(74,970)	(22,049)	(97,019)	(66,993)	(902)	(67,895)	(11.9%)	(42.9%)
% of pre-tax income	24.5%	-	27.5%	21.9%	-	28.6%		
CONSOLIDATED PROFIT	230,803	24,961	255,764	47,041	122,799	169,841	390.6%	50.6%
Results Attributable to Non-Controlling Interests	(54,016)	(2,076)	(56,092)	(10,782)	(6,612)	(17,394)	(401.0%)	(222.5%)
GROUP PROFIT	176,787	22,885	199,672	36,259	116,187	152,447	387.6%	31.0%
% Net revenue	4.8%	-	5.4%	1.1%	-	4.4%		

Q2 2025 Results - 25 - GRIFOLS

Cash Flow | Q2 2025

In thousands of euros (on a reported basis)	Q2 2025	Q2 2024	% vs PY
Reported Group Profit	117,062	14,841	689%
Depreciation and Amortization	107,035	113,786	(6%)
Net Provisions	17,320	40,220	(57%)
Other Adjustments and Other Changes in Working Capital	(55,979)	(80,290)	30%
Change in Operating Working Capital	(30,170)	164,304	(118%)
Changes in Inventories	(30,602)	(2,772)	(1004%)
Change in Trade Receivables	(24,810)	64,011	(139%)
Change in Trade Payables	25,242	103,065	(76%)
Net Cash Flow From Operating Activities	155,267	252,861	(39%)
Business Combinations and Investments in Group Companies	(23,174)	1,445,351	(102%)
CAPEX	(60,249)	(43,066)	(40%)
R&D/Other Intangible Assets	(34,374)	(41,247)	17%
Other Cash Inflow / (Outflow)	(7,853)	(11,709)	33%
Net Cash Flow From Investing Activities	(125,651)	1,349,329	(109%)
Free Cash Flow	29,617	1,602,190	(98%)
Issue / (Repayment) of Debt	(70,070)	49,861	(241%)
Capital Grants	(2,802)	2,004	(240%)
Other Cash Flows From / (Used in) Financing Activities	(90,534)	7,574	(1295%)
Net Cash Flow From Financing Activities	(163,407)	59,439	(375%)
Total Cash Flow	(133,791)	1,661,629	(108%)
Cash and Cash Equivalents at the Beginning of the Period	752,818	448,545	68%
Effect of Exchange Rate Changes in Cash and Cash Equivalents	(60,336)	3,046	(2081%)
Cash and Cash Equivalents at the End of the Period	558,691	2,113,220	(74%)



Q2 2025 Results - 26 -

Cash Flow | H1 2025

In thousands of euros (on a reported basis)	2025 YTD	2024 YTD	% vs PY
Reported Group Profit	176,787	36,259	388%
Depreciation and Amortization	218,785	219,402	0%
Net Provisions	27,861	54,886	(49%)
Other Adjustments and Other Changes in Working Capital	25,555	(47,656)	154%
Change in Operating Working Capital	(158,252)	(174,952)	10%
Changes in Inventories	(91,428)	(132,882)	31%
Change in Trade Receivables	(118,155)	(89,669)	(32%)
Change in Trade Payables	51,331	47,599	8%
Net Cash Flow From Operating Activities	290,735	87,939	231%
Business Combinations and Investments in Group Companies	(102,172)	1,425,131	(107%)
CAPEX	(109,395)	(81,116)	(35%)
R&D/Other Intangible Assets	(72,879)	(63,251)	(15%)
Other Cash Inflow / (Outflow)	(20,549)	(19,387)	(6%)
Net Cash Flow From Investing Activities	(304,995)	1,261,377	(124%)
Free Cash Flow	(14,260)	1,349,316	(101%)
Issue / (Repayment) of Debt	(223,744)	203,586	(210%)
Capital Grants	3,463	5,699	(39%)
Other Cash Flows From / (Used in) Financing Activities	(94,779)	15,610	(707%)
Net Cash Flow From Financing Activities	(315,060)	224,895	(240%)
Total Cash Flow	(329,320)	1,574,211	(121%)
Cash and Cash Equivalents at the Beginning of the Period	979,780	529,577	85%
Effect of Exchange Rate Changes in Cash and Cash Equivalents	(91,769)	9,432	(1073%)
Cash and Cash Equivalents at the End of the Period	558,691	2,113,220	(74%)



Q2 2025 Results - 27 -

Balance Sheet | 2025

In thousands of euros

Assets

	jun-25	dic-24
Non-Current Assets	14,520,953	15,677,699
Goodwill and Other Intangible Assets	10,538,461	11,297,492
Property Plant & Equipment	3,096,578	3,341,846
Investments in Equity Accounted Investees	77,507	68,996
Non-Current Financial Assets	423,495	490,492
Other Non-Current Assets	384,911	478,873
Current Assets	5,246,208	5,727,543
Non-Current Contract Assets Held for Sale	-	-
Inventories	3,346,864	3,560,098
Current Contract Assets	88,012	35,978
Trade and Other Receivables	933,125	836,015
Other Current Financial Assets	263,377	243,156
Other Current Assets	56,139	72,515
Cash and Cash Equivalents	558,691	979,780
Total Assets	19,767,159	21,405,241

Liabilities

	jun-25	dic-24	
Equity	7,598,153	8,607,025	
Capital	119,604	119,604	
Share Premium	910,728	910,728	
Reserves	4,185,197	4,054,505	
Treasury Stock	(132,168)	(134,448)	
Current Year Earnings	176,786	156,920	
Other Comprehensive Income	(91,067)	776,418	
Non-Controllling Interests	2,429,073	2,723,298	
No-Current Liabilities	10,178,855	10,642,070	
Non-Current Financial Liabilities	9,117,632	9,490,644	
Other Non-Current Liabilities	1,061,223	1,151,426	
Current Liabilities	1,990,152	2,156,146	
Current Financial Liabilities	522,103	676,087	
Other Current Liabilities	1,468,049	1,480,059	
Total Equity and Liabilities	19,767,159	21,405,241	



Q2 2025 Results - 28 -

Like-for-Like (LFL) Reconciliation

In millions of euros	Q2'25	Q1'25	H1'25
Revenue Reported	1,891	1,786	3,677
Fee-for-Service / GPO Reclassification	33	15	48
Inflation Reduction Act (IRA)	30	28	58
Revenue Like-for-Like	1,954	1,829	3,783
In millions of euros	Q2'25	Q1'25	H1'25
<u>In millions of euros</u> Operating Results (EBIT)	Q2'25 349	Q1'25 269	H1'25
Operating Results (EBIT)	349	269	618
Operating Results (EBIT) Depreciation & Amortization	349 107	269 112	618 219
Operating Results (EBIT) Depreciation & Amortization Reported EBITDA	349 107 456	269 112 381	618 219 836
Operating Results (EBIT) Depreciation & Amortization Reported EBITDA Total adjustments	349 107 456 19	269 112 381 20	618 219 836 40
Operating Results (EBIT) Depreciation & Amortization Reported EBITDA Total adjustments EBITDA Adjusted	349 107 456 19 475	269 112 381 20 400	618 219 836 40 876



Q2 2025 Results - 29 -

EBIT to EBITDA and EBITDA Adjusted

In thousand of euros	Q2 2025	Q1 2025	Q4 2024	Q3 2024	Q2 2025 LTM	Q2 2024
OPERATING RESULT (EBIT)	348,854	268,857	371,859	317,034	1,306,605	299,321
Depreciation & Amortization	(107,035)	(111,750)	(110,130)	(108,364)	(437,280)	(114,310)
Reported EBITDA	455,889	380,607	481,990	425,398	1,743,884	413,631
% Net revenue	24.1%	21.3%	24.4%	23.7%	23.4%	22.8%
			1 000	24.672	22.502	10.005
Restructuring costs	-	-	1,889	21,673	23,562	10,095
Transaction costs	3,842	7,466	9,306	7,882	28,495	16,145
Impairments	-	3,850	24,265	787	28,902	-
Biotest Next Level Project	5,481	6,738	7,340	5,113	24,672	4,922
SRAAS One-off	-	-	-	-	-	(5,618)
Other non-recurring items	9,873	1,817	1,155	1,245	14,089	1,613
Total adjustments	19,195	19,872	43,954	36,700	119,720	27,157
Adjusted EBITDA	475,084	400,479	525,944	462,098	1,863,605	440,788
% Net revenue	25.1%	22.4%	26.6%	25.8%	25.0%	24.2%



Q2 2025 Results - 30 - -

Leverage Ratio as per Credit Agreement

In millions of euros except ratio.	Q2'25	Q1'25	Q4'24	Q3'24	Q2'24
Non-Current Financial Liabilities	9,118	9,390	9,491	8,836	8,752
Non-recurrent Lease Liabilities (IFRS16)	(978)	(1,026)	(1,025)	(969)	(1,025)
Current Financial Liabilities	522	657	676	1,017	2,757
Recurrent Lease Liabilities (IFRS16)	(112)	(119)	(117)	(111)	(109)
Cash and Cash Equivalents	(559)	(753)	(980)	(645)	(2,113)
Net Financial Debt as per Credit Agreement	7,992	8,149	8,046	8,128	8,262

In millions of euros except ratio.	LTM Q2'25	LTM Q1'25	LTM Q4'24	LTM Q3'24	LTM Q2'24
OPERATING RESULT (EBIT)	1,307	1,257	1,192	1,075	1,005
Depreciation & Amortization	(437)	(445)	(439)	(443)	(444)
Reported EBITDA	1,744	1,702	1,631	1,518	1,450
IFRS 16	(118)	(117)	(113)	(113)	(110)
Restructuring costs	52	63	55	57	34
Transaction costs	28	41	49	59	65
Cost savings, operating improvements and synergies on a "run rate"	173	165	159	146	136
Other one-offs	23	(34)	(28)	(62)	(75)
Total adjustments	159	119	122	87	50
Adjusted EBITDA LTM as per Credit Agreement	1,902	1,819	1,753	1,605	1,500
Leverage Ratio as per Credit Agreeement	4.2x	4.5x	4.6x	5.1x	5.5x



Q2 2025 Results - 31 - -

Leverage Ratio as per Consolidated EBITDA and Net Debt as per Balance Sheet

In millions of euros except ratio.	Q2'25	Q1'25	Q4'24	Q3'24	Q2'24
Non-Current Financial Liabilities	9,118	9,390	9,491	8,836	8,752
Current Financial Liabilities	522	657	676	1,017	2,757
Cash and Cash Equivalents	(559)	(753)	(980)	(645)	(2,113)
Net Financial Debt	9,081	9,294	9,187	9,208	9,396

In millions of euros except ratio.	LTM Q2'25	LTM Q1'25	LTM Q4'24	LTM Q3'24	LTM Q2'24
OPERATING RESULT (EBIT)	1,307	1,257	1,192	1,075	1,005
Depreciation & Amortization	(437)	(445)	(439)	(443)	(444)
Reported EBITDA	1,744	1,702	1,631	1,518	1,450
Leverage Ratio Reported	5.2x	5.5x	5.6x	6.1x	6.5x

GRIFOLS

Q2 2025 Results - 32 - -

NCI Contributions

LTM Q2 2025

In thousand of euros	GDS	Biotest	BPC	Haema
Profit after tax from continuing operations	139,858	(108,513)	46,846	15,634
Income tax expense	(38,234)	30,048	(13,336)	(10,710)
Financial result	78,678	(31,383)	(2,073)	8,093
Amortisation and depreciation	(47,060)	(51,138)	(7,739)	(8,639)
Consolidated EBITDA	146,473	(56,040)	69,995	26,890
Impact IFRS16- Finance Leases (leases of plasma donation centre properties)	(2,512)	(8,309)	(5,759)	(4,738)
Restructuring costs	861	1,050	-	230
Share of profits assoc core activit	-	19,154	-	-
Impairment	-	4,388	-	-
Consolidated EBITDA under Credit Agreement	144,823	(39,758)	64,236	22,382
% of non-controlling interest	45%	20%	100%	100%
Consolidated EBITDA according to Credit Agreement non-controlling interest	65,170	(7,824)	64,236	22,382
Cash and cash equivalents	(993)	(57,683)	(5,491)	(21,569)
Financial assets/liabilities with Grifols	(1,050,599)	607,353	· -	-
Leasing liabilities (leases of real estate of plasma donation centres)	11,463	58,270	51,777	20,958
Loans and other financial liabilities	1,446	66,260	121	-
Total Balance Sheet Net Debt	(1,038,682)	674,200	46,407	(611)
Impact IFRS16- Finance Leases (leases of plasma donation centre properties)	(11,463)	(58,270)	(51,777)	(20,958)
impact if Noto Finance Leases (leases of plasma donation centre properties)	(11,403)	(30,270)	(31,777)	(20,930)
Total Net Financial Debt according to Credit Agreement	(1,050,146)	615,930	(5,370)	(21,569)

Note: Last Twelve Months figures (LTM).



FCF pre-M&A Reconciliation to Cash Flow Statement

Free Cash Flow pre-M&A = EBITDA Adjusted- Net Working Capital - CAPEX (including capitalized IT and R&D, and extraordinary growth CAPEX) - Others - Interest - Taxes. In the Consolidated Annual Accounts, this reconciles to Cash flow generation from operating and investing activities excluding impact from M&A and associated costs and expenses. Excludes lease payments, consistent with prior disclosed guidance.

In million Euros	Q2'25	Q2'24
EBITDA Adjusted	475	441
Changes in working capital	(30)	164
CAPEX	(60)	(43)
R&D and IT	(34)	(41)
Taxes	(45)	(59)
Interests	(235)	(233)
Others	(12)	-32
FCF Before Extraordinary Items	59	196
Extraordinary Growth CAPEX	(23)	(119)
Restructuring and transaction costs	(6)	(20)
Free Cash Flow	30	57

In million Furos

H1'25	H1'24
876	791
(158)	(175)
(109)	(81)
(73)	(63)
(48)	(63)
(290)	(340)
(98)	-98
99	(29)
(102)	(141)
(11)	(25)
(14)	(196)

In million Euros	Q2'25	H1'25
Net Cash Flow From Operating Activities ¹	155	291
Net Cash Flow From Investing Activities ¹	(126)	(305)
Free Cash Flow	30	(14)

¹ Statement of Cash Flow According IFRS-EU



Q2 2025 Results - 34 -

Net Revenue Reconciliation at cc | Q2 2025

In thousands of euros	Q2 2025	Q2 2024	% Var
Reported Net Revenues	1,891,132	1,817,908	4.0%
Variation due to Exchange Rate Effects	46,001		
Net Revenues at Constant Currency	1,937,133	1,817,908	6.6%
In thousands of euros	Q2 2025	Q2 2024	% Var
	·		
Reported Biopharma Net Revenues	1,632,629	1,527,652	6.9%
Variation due to Exchange Rate Effects	40,932		
Reported Biopharma Net Revenues at Constant Currency	1,673,561	1,527,652	9.6%
In thousands of euros	Q2 2025	Q2 2024	% Var
Reported Diagnostic Net Revenues	161,588	164,261	(1.6%)
Variation due to Exchange Rate Effects	3,398		
Reported Diagnostic Net Revenues at Constant Currency	164,986	164,261	0.4%
In thousands of euros	Q2 2025	Q2 2024	% Var
Reported Bio Supplies Net Revenues	36,403	73,719	(50.6%)
Variation due to Exchange Rate Effects	936		
Reported Bio Supplies Net Revenues at Constant Currency	37,339	73,719	(49.3%)
In thousands of euros	Q2 2025	Q2 2024	% Var
Reported Others & Intersegments Net Revenues	60,512	52,276	15.8%
Variation due to Exchange Rate Effects	735		
Reported Other & Intersegments Net Revenues at Constant Currency	61,247	52,276	17.2%

In thousands of euros	Q2 2025	Q2 2024	% Var
Reported U.S. + Canada Net Revenues	1,067,726	1,037,872	2.9%
Variation due to Exchange Rate Effects	29,538		
Reported U.S. + Canada Net Revenues at Constant Currency	1,097,264	1,037,872	5.7%
In thousands of euros	Q2 2025	Q2 2024	% Var
Reported EU Net Revenues	401,252	418,882	(4.2%)
Variation due to Exchange Rate Effects	130		
Reported EU Net Revenues at Constant Currency	401,382	418,882	(4.2%)
In thousands of euros	Q2 2025	Q2 2024	% Var
Reported ROW Net Revenues	422,154	361,154	16.9%
Variation due to Exchange Rate Effects	16,334		
Reported ROW Net Revenues at Constant Currency	438,488	361,154	21.4%



Q2 2025 Results - 35 - -

Net Revenue Reconciliation at cc | H1 2025

In thousands of euros	H1 2025	H1 2024	% Var
Reported Net Revenues	3,676,940	3,443,613	6.8%
Variation due to Exchange Rate Effects	6,565		
Net Revenues at Constant Currency	3,683,505	3,443,613	7.0%
In thousands of euros	H1 2025	H1 2024	% Var
Reported Biopharma Net Revenues	3,153,789	2,922,355	7.9%
Variation due to Exchange Rate Effects	7,009		
Reported Biopharma Net Revenues at Constant Currency	3,160,798	2,922,355	8.2%
In thousands of euros	H1 2025	H1 2024	% Var
Reported Diagnostic Net Revenues	331,632	322,544	2.8%
Variation due to Exchange Rate Effects	(53)		
Reported Diagnostic Net Revenues at Constant Currency	331,579	322,544	2.8%
In thousands of euros	H1 2025	H1 2024	% Var
Reported Bio Supplies Net Revenues	68,958	100,731	(31.5%)
Variation due to Exchange Rate Effects	(306)	100,731	(31.3%)
_	` ′	100 721	(24.00()
Reported Bio Supplies Net Revenues at Constant Currency	68,652	100,731	(31.8%)
In thousands of euros	H1 2025	H1 2024	% Var
Reported Others & Intersegments Net Revenues	122,561	97,983	25.1%
Variation due to Exchange Rate Effects	(85)		
Reported Other & Intersegments Net Revenues at Constant Currency	122,476	97,983	25.0%

In thousands of euros	H1 2025	H1 2024	% Var
Reported U.S. + Canada Net Revenues	2,092,916	1,963,198	6.6%
Variation due to Exchange Rate Effects	(10,778)		
Reported U.S. + Canada Net Revenues at Constant Currency	2,082,138	1,963,198	6.1%
In thousands of euros	H1 2025	H1 2024	% Var
Reported EU Net Revenues	791,526	750,309	5.5%
Variation due to Exchange Rate Effects	(547)		
Reported EU Net Revenues at Constant Currency	790,979	750,309	5.4%
In thousands of euros	H1 2025	H1 2024	% Var
In allousands of Euros			
Reported ROW Net Revenues	792,498	730,106	8.5%
Variation due to Exchange Rate Effects	17,889		
Reported ROW Net Revenues at Constant Currency	810,387	730,106	11.0%



Q2 2025 Results - 36 -

EBITDA Adjusted Reconciliation at cc | Q2 2025

EBITDA Adjusted Q2'25:

In thousands of euros	Q2 2025	Q2 2024	% Var
EBITDA Adjusted	475,086	440,788	7.8%
Variation due to Exchange Rate Effects	16,443		
EBITDA Adjusted at Constant Currency	491,529	440,788	11.5%

EBITDA Adjusted Like-for-Like Q2'25:

In thousands of euros	Q2 2025	Q2 2024	% Var
EBITDA Adjusted Like for Like	505,870	440,788	14.8%
Variation due to Exchange Rate Effects	17,430		
EBITDA Adjusted Like for Like at Constant Currency	523,300	440,788	18.7%



Q2 2025 Results - 37

EBITDA Adjusted Reconciliation at cc | H1 2025

EBITDA Adjusted H1'25:

In thousands of euros	H1 2025	H1 2024	% Var
EBITDA Adjusted	875,565	791,190	10.7%
Variation due to Exchange Rate Effects	15,952		
EBITDA Adjusted at Constant Currency	891,517	791,190	12.7%

EBITDA Adjusted Like-for-Like H1'25:

In thousands of euros	H1 2025	H1 2024	% Var
EBITDA Adjusted Like for Like	934,040	791,190	18.1%
Variation due to Exchange Rate Effects	15,832		
EBITDA Adjusted Like for Like at Constant Currency	949,872	791,190	20.1%



Q2 2025 Results - 38 -

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