

First Half 2018 Results

Grifols increases net profits by 15% to EUR 319 million, with sustained operational revenue growth of 7%

- **Revenues grow to EUR 2,120 million, a 7.1% cc¹ increase driven by growth in all divisions and global markets**
- **Bioscience Division sales grow by 6.6%² cc to EUR 1,690 million, with a significant upturn in sales of the main plasma proteins**
- **Grifols reinforces its corporate strategy to increase and diversify its plasma supply and consolidates its leadership position with 225 centers, 190 in the U.S. and 35 in Europe following the acquisition of Haema**
- **Diagnostic Division sales reach EUR 339 million (2.2%² cc) and new approvals broaden its portfolio of transfusion medicine and specialty diagnostic solutions**
- **Hospital Division sales expand by 20.7%² cc to EUR 59 million, spurred by sales of physiological saline solution in the U.S. and growth of Pharmatech**
- **Bio Supplies reaches EUR 40 million in sales, a 40.9%² cc increase**
- **EBITDA reaches EUR 614 million and EBITDA margin remains stable at 29.0%**
- **Financial results and lower taxes in the U.S. contribute to a 15% increase in net profit to EUR 319 million**
- **Grifols allocates EUR 265 million to dividends in 2017, a 21.5% year-on-year increase, after paying EUR 142 million for the final dividend in June 2018**

Barcelona, July 27, 2018.- Grifols (MCE: GRF, MCE: GRF.P, NASDAQ: GRFS) reported EUR 2,120.1 million in revenues for the first half of 2018, a 7.1% increase at constant currency (cc) and a 3.3% decrease when taking into account exchange rate fluctuations, particularly the euro-dollar. The company consolidated its growth in all divisions and regions where it operates.

Demand for the main plasma proteins remains strong, as evidenced by higher sales of immunoglobulin, albumin and alpha-1 antitrypsin. The **Bioscience Division** reported sales of EUR 1,689.9 million, a 6.6%² cc increase and 4.0% decline taking exchange rate variations into account.

¹ Constant currency (cc) excludes exchange rate variations.

² Comparable revenues considering inter-segment sales.

The **Diagnostic Division's** revenues reached EUR 339.4 million, a 2.2%² increase and 7.0% decline due to foreign currency exchange fluctuations. Sales of the division's NAT technology donor-screening solutions (Procleix[®] NAT Solutions) and blood typing business lines were the main engines of growth.

The **Hospital Division** reached EUR 58.7 million in revenues, an increase of 20.7%² cc and 16.1% when factoring in the exchange rate. This upward trend was driven primarily by higher U.S. sales of Grifols' IV solutions, manufactured in the group's Murcia (Spain) plant, and the international expansion of the division's Pharmatech line, comprised by hospital pharmacy systems and equipment.

The **Bio Supplies Division** recorded revenues of EUR 40.1 million for the first six months of 2018, an uptick of 40.9%² cc and 25.1% taking into account exchange rate variations.

EBITDA totaled EUR 614.2 million and the EBITDA margin remains stable at 29.0%. The group continues to note the impact of higher plasma costs associated with its strategic long-term investment plan to increase and diversify its plasma supply. In alignment with this plan, Grifols aims to satisfy the projected growing demand for plasma proteins and remain on its path of sustainable growth.

Grifols' investment efforts have reinforced its leadership position in plasma donation centers. The company currently owns 225 centers: 190 in the U.S. and 35 in Europe following the acquisition of the German firm Haema. In addition, the execution of a call option for the remaining 51% of Interstate Blood Bank Inc. (IBBI), exercisable in 2019, will expand the group's network by 26 centers.

Net **R+D+i** investments totaled EUR 141.3 million, including both in-house and external projects. This figure represents a 9.3% increase compared to the same period last year.

As part of its integrated R+D+i strategy, Grifols continuously assesses the suitability of its diverse projects. To this end, the company decided to divest in TiGenix and tender its shares in a takeover bid by Takeda, resulting in a cash influx of EUR 70.1 million and gain of EUR 32.0 million. The transaction improved the **financial result** by 30.1% to EUR -103.2 million, compared to EUR -147.6 million for the same period in 2017.

The **effective tax rate** remains at 20% following the U.S. tax reform approved in December 2017.

Net profit increased by 14.8% during the first half of 2018 to EUR 319.0 million, which represents 15.0% of total revenues.

At the end of June 2018, Grifols' **net financial debt** totaled EUR 5,560.3 million, including EUR 668.5 million in cash and taking into account, among other transactions, the EUR 220.0 million acquisition of 100% of Haema's capital and EUR 142.1 million payout for the final 2017 dividend, approved in the General Ordinary Shareholders Meeting.

Total dividend allocations in 2017, including the final dividend paid in June 2018 (EUR 0.20 gross per share) and the interim dividend paid in December 2017 (EUR 0.18 gross per share), amounted to EUR 265.1 million. This record figure denotes a 21.5% increase compared to the previous year and confirms Grifols' commitment to generating and delivering shareholder value.

The net financial debt-to-EBITDA ratio was 4.43x (4.34x cc). Standard & Poor's (S&P) improved Grifols' credit scores by raising the rating on its senior secured debt to BB+. The corporate rating remains at BB and its outlook is "stable".

As of June 30, 2018, undrawn lines of credit totaled EUR 400 million and Grifols' liquidity position was roughly EUR 1,100 million.

Grifols' cash flow generation remains high and provides the necessary solvency to meet growth investments. Unlevered operating cash flow reached EUR 348.1 million in the first half of 2018, bearing in mind higher inventory levels stemming from greater sales volume and new plasma centers.

Key Financial Metrics for the First Half of 2018:

<i>In millions of euros except % and EPS</i>	1H 2018	1H 2017	% Var
NET REVENUE (NR)	2,120.1	2,192.4	(3.3%)
GROSS MARGIN	47.5%	50.3%	
EBITDA	614.2	644.4	(4.7%)
% NR	29.0%	29.4%	
ADJUSTED EBITDA⁽¹⁾	614.2	663.9	(7.5%)
% NR	29.0%	30.3%	
EBIT	506.2	537.8	(5.9%)
% NR	23.9%	24.5%	
REPORTED GROUP PROFIT	319.0	277.9	14.8%
% NR	15.0%	12.7%	
ADJUSTED⁽²⁾ GROUP PROFIT	355.9	330.2	7.8%
% NR	16.8%	15.1%	
CAPEX	102.1	135.3	(24.5%)
R&D NET INVESTMENT	141.3	129.3	9.3%
EARNINGS PER SHARE (EPS) REPORTED	0.47	0.41	14.8%
	June 2018	December 2017	% Var
TOTAL ASSETS	11,433.6	10,920.3	4.7%
TOTAL EQUITY	3,971.2	3,634.0	9.3%
CASH & CASH EQUIVALENTS	668.5	886.5	(24.6%)
LEVERAGE RATIO	4.43./(4.34 cc) ⁽³⁾	3.96/(4.34 cc) ⁽³⁾	

⁽¹⁾ Excludes non-recurring items and associated with recent acquisitions

⁽²⁾ Excludes non-recurring items and associated with recent acquisitions, amortization of deferred expenses associated to the refinancing and amortization of intangible assets related to acquisitions

⁽³⁾ Constant currency (cc) excludes the impact of exchange rate movements

PERFORMANCE BY DIVISION

Grifols drives “One Grifols” strategy among its main divisions

Bioscience Division: 6.6% solid operating growth and strategic investments to meet rising market demand

Demand in the hemoderivatives sector is solid and maintains its upward trend. Grifols further consolidated the Bioscience Division’s sales growth and continues to make inroads to increase and diversify its access to plasma.

The division recorded a 6.6%² cc increase in revenue growth to EUR 1,689.9 million over the first half of the year. Higher sales volumes of the main plasma proteins (immunoglobulin, albumin and alpha-1 antitrypsin) and the favorable price impact in some markets offset the decline in factor VIII sales.

The euro-dollar exchange rate exerted a negative effect on the division’s overall performance, resulting in a 4% decline compared to the same period last year.

Sales of **immunoglobulin** were the primary drivers of growth during this period. Demand for this plasma protein continues to grow, especially in the U.S. and European Union countries.

Grifols is the global leader in immunoglobulin sales. It boasts a solid position in the treatment of primary immunodeficiencies (PID) and leads the neurology area to treat diseases such as chronic inflammatory demyelinating polyneuropathy (CIDP).

Sales of **alpha-1 antitrypsin** grew significantly in the U.S. and European countries as a result of higher rates of diagnosis.

Grifols maintains its leadership position in alpha-1 antitrypsin sales and expanded its product portfolio. The new liquid alpha-1 formulation (Prolastin[®]-C Liquid) was approved by the U.S. Food and Drug Administration (FDA) and is scheduled for launch in the second half of 2018. This new formulation, along with the new FDA-approved genetic diagnostic test developed by the Diagnostic Division, will contribute to improving the diagnosis and treatment of alpha-1 antitrypsin deficiency.

Albumin sales notably increased, especially in China, the U.S. and European countries.

Plasma-derived Factor VIII sales followed the same trend as the first quarter of 2018. Demand has dropped significantly as a result of declining use to treat patients with inhibitors in immune tolerance induction (ITI) therapy.

Despite this downturn, the company continues to advocate plasma-derived factor VIII as the best treatment option to eradicate inhibitors, which affect an estimated 35% of hemophilia A patients³. On the other hand, Grifols continues to reinforce its position to treat previously untreated patients (PUPs) with severe hemophilia A, especially in the United States.

³ Source: Oldengurg J, et al. *Haematologica* 2015; 100(2):149-156

Grifols remains committed to expanding its line of [specialty proteins](#), which allow the company to build a differential product portfolio for patients, as well as optimize production capacity and raw materials costs.

In the hyperimmunoglobulins segment, Grifols expanded its product portfolio with the development of two new formulations: intramuscular immunoglobulin (GamaSTAN[®]) to treat patients exposed with the hepatitis A virus and measles, scheduled to launch in the second half of the year; and anti-rabies immunoglobulin (HyperRAB[®]), introduced last May in the U.S. to treat patients for rabies exposure. Both earned FDA approval in the first half of 2018.

Diagnostic Division: continuous innovation leads to a broader product portfolio, with 5 new FDA-approved products

Diagnostic Division revenues reached EUR 339.4 million, representing a 2.2%² cc increase and a 7.0% decline taking into account exchange rate variations.

[Transfusion medicine](#) continues to be the main driver of growth. Sales of NAT technology for plasma and blood donation screening ([Procleix[®] NAT Solutions](#)) remained robust.

The company continues its efforts to enhance its product portfolio with the development of new reagents. In the second quarter of the year, the FDA approved two new Procleix[®] Panther diagnostic tests, one that simultaneously detects two types of the human immunodeficiency virus (HIV-1 and HIV-2) and hepatitis B and C, and another that detects the West Nile virus. The market launch is scheduled for the second half of 2018.

Sales of the division's blood typing line rose significantly, especially analyzers (Wadiana[®], Erytra[®] and Erytra Eflexys[®]) and reagents (DG-Gel[®] cards). A solid sales strategy in the U.S. and Europe fueled this strong performance. A year following its launch in Europe, Middle East and Africa more than 100 units of Erytra Eflexys[®] have been sold in the region of which 60% are conversions from competitors.

The company began marketing its new line of conventional antisera in the U.S., used to determine blood types and carry out manual pre-transfusion blood compatibility tests, after earning FDA approval in the second quarter of 2018. This FDA approval marks an important milestone since it allows the division to expand and complement its blood-typing product portfolio.

In addition to the positive strides in transfusion medicine, Grifols fortified its position in [specialty diagnostics](#) after earning new approvals that widen its product portfolio. In the first half of 2018, the FDA approved two new diagnostic tests to detect autoimmune diseases. These diagnostics utilize the HELIOS system developed by Aesku and distributed by Grifols.

Moreover, in May 2018, Grifols' Immunohematology Center in San Marcos, Texas (U.S.) enhanced its catalogue of transfusion tests with a blood compatibility test used for certain cases for multiple myeloma patients.

Hospital Division: global expansion boosts growth by more than 20% cc

The Hospital Division increased its revenues by 20.7%² cc (16.1%) to EUR 58.7 million. The division reported higher sales in all of its business lines, most notably **IV solutions** following the distribution of Grifols' physiological saline solution in the U.S., as well as their usage in Grifols' network of plasma donation centers to restore circulatory volume. This will increase the vertical integration of the process and its quality and regular supply. Grifols' IV solutions are manufactured in the Murcia (Spain) plant.

Sales of the **Pharmatech** line, comprised by hospital pharmacy solutions and reinforced with the acquisition of MedKeeper, grew considerably in the U.S. and in certain Latin American markets. Regulatory changes in hospital pharmacy and compounding operations in the U.S. represent a significant market opportunity for Grifols, a recognized supplier of integrated solutions that enhance the efficiency and control of hospital pharmacy services.

Bio Supplies Division

This division focuses mainly on sales of biological products for non-therapeutic uses and overseeing manufacturing agreements with Kedrion, which led to an increase in sales to EUR 40.1 million compared to EUR 32.1 million reported in the same period in 2017.

Revenues by division and region:

<i>In thousands of euros</i>	1H 2018	% of Net Revenues	1H 2017**	% of Net Revenues	% Var	% Var cc*
BIOSCIENCE	1,689,875	79.7%	1,759,852	80.3%	(4.0%)	6.6%
DIAGNOSTIC	339,432	16.0%	365,014	16.6%	(7.0%)	2.2%
HOSPITAL	58,734	2.8%	50,610	2.3%	16.1%	20.7%
BIO SUPPLIES	40,124	1.9%	32,073	1.5%	25.1%	40.9%
OTHERS	11,578	0.5%	1,606	0.1%	620.9%	701.8%
INTERSEGMENTOS	(19,625)	(0.9%)	(16,708)	(0.8%)	17.5%	31.1%
TOTAL	2,120,118	100.0%	2,192,447	100.0%	(3.3%)	7.1%

* Constant currency (cc) excludes the impact of exchange rate movements

** Comparable revenues considering intersegment sales

<i>In thousands of euros</i>	1H 2018	% of Net Revenues	1H 2017	% of Net Revenues	% Var	% Var cc*
US + CANADA	1,412,542	66.6%	1,494,131	68.2%	(5.5%)	7.0%
EU	369,207	17.4%	338,288	15.4%	9.1%	9.5%
ROW	338,369	16.0%	360,028	16.4%	(6.0%)	5.1%
TOTAL	2,120,118	100.0%	2,192,447	100.0%	(3.3%)	7.1%

* Constant currency (cc) excludes the impact of exchange rate movements

SECOND QUARTER 2018

Operating growth in Grifols' main divisions and geographic regions

Grifols' revenues reached EUR 1,097.1 million in the second quarter of 2018, a 6.7%² cc increase and 3.0% decline taking into account exchange rate variations. Sales grew in the main divisions and all regions where the company operates. Sales in the principal European Union markets (+8.0% cc and +7.7%), as well as in the U.S. and Canada (+7.3% cc and -4.3%) were especially strong.

The Bioscience Division led overall sales, with a 7.4%² cc increase in revenues to EUR 882.3 million. Of note was the continued solid demand for immunoglobulin and alpha-1 antitrypsin in the U.S. and European countries, and higher sales of albumin in China, the U.S. and European countries.

Revenue growth of the Diagnostic Division moderated in the second quarter of 2018, reaching EUR 174.5 million. The division's sales were led by growth in NAT technology systems and blood typing solutions.

The Hospital Division grew by 23.0%² cc (17.6%) to EUR 31.4 million, proof of its solid internationalization strategy and clear focus on the U.S. market.

Revenues by division and region:

<i>In thousands of euros</i>	2Q 2018	% of Net Revenues	2Q 2017**	% of Net Revenues	% Var	% Var cc*
BIOSCIENCE	882,334	80.4%	906,213	80.1%	(2.6%)	7.4%
DIAGNOSTIC	174,501	15.9%	189,880	16.8%	(8.1%)	0.5%
HOSPITAL	31,419	2.9%	26,709	2.4%	17.6%	23.0%
BIO SUPPLIES	13,968	1.3%	17,671	1.6%	(21.0%)	(12.1%)
OTHERS	7,133	0.7%	1,573	0.1%	353.5%	400.5%
INTERSEGMENTS	(12,249)	(1.2%)	(11,279)	(1.0%)	8.6%	20.7%
TOTAL	1,097,106	100.0%	1,130,767	100.0%	(3.0%)	6.7%

* Constant currency (cc) excludes the impact of exchange rate movements

** Comparable revenues considering intersegment sales

<i>In thousands of euros</i>	2Q 2018	% of Net Revenues	2Q 2017	% of Net Revenues	% Var	% Var cc*
US + CANADA	732,929	66.8%	765,561	67.7%	(4.3%)	7.3%
EU	190,103	17.3%	176,541	15.6%	7.7%	8.0%
ROW	174,074	15.9%	188,665	16.7%	(7.7%)	3.1%
TOTAL	1,097,106	100.0%	1,130,767	100.0%	(3.0%)	6.7%

* Constant currency (cc) excludes the impact of exchange rate movements

INVESTMENT ACTIVITIES: ACQUISITIONS, CAPEX AND R+D+i

Haema acquisition

In alignment with Grifols' corporate strategy to expand and diversify its access to plasma, the company announced the acquisition of 100% share capital of Haema, the leading independent network of donation centers in Germany and largest transfusion service in the country. After fulfilling the conditions set for the transaction, Grifols acquired the firm for EUR 220 million.

The transaction includes the Haema business; 35 donation centers in nine states and three more under construction; a 24,000-square-meter building in Leipzig (Germany), home to the company's headquarters; and a main laboratory in Berlin (Germany). Haema collected approximately 800,000 liters of plasma in 2017.

Agreement with Boya Bio-Pharmaceutical

Grifols has entered into an agreement with Boya Bio-Pharmaceutical, a leading Chinese producer of plasma-derived medicines, to build and manage plasma donation centers in China.

The project investment totals EUR 50 million and Grifols will control 50% of the political and economic rights.

The plasma donation centers will be built and managed in adherence to the guidelines established by the China's National Health and Family Planning Commission, the FDA and the European Medicines Agency (EMA), among others. Grifols will bring its experience and know-how to ensure that the construction and management of the centers meet the same high standards of quality as the rest of its global network.

In accordance with current Chinese legislation, the plasma collected in these centers will be supplied to Boya Bio-Pharmaceutical, although Grifols reserves the right to access up to 50% of the total volume when the applicable legislation allows.

Capital investments (CAPEX)

Grifols invested EUR 102.1 million over the first six months of the year as part of its on-going efforts to enhance and expand the production facilities of its four divisions. Capital investments progress as outlined in the 2016-2020 Capital Investment Plan, endowed with EUR 1,200 million to guarantee the company's long-term sustainable growth.

More than EUR 140 million in R+D+i investments in the first half

The company allocated EUR 141.3 million for R+D+i activities in the first half of 2018, taking into account net internal and external investments. This figure represents a 9.3% increase compared to the same period last year.

In terms of clinical trials, Grifols continues to research the potential benefits of albumin in the treatment of cirrhosis. Also of note is the completion of the AMBAR (Alzheimer Management By Albumin Replacement) phase IIb/III clinical trial for the treatment of Alzheimer's disease. The company plans to publish AMBAR's results in the fourth quarter of 2018.

CORPORATE RESPONSIBILITY

Talent: greater job creation, training and professional development

The Grifols' team grew to 18,664 employees over the first half of the year, a 2% increase compared to the same period in 2017. For administrative purposes, these figures do not include the approx. 1,100 Haema employees who now form part of Grifols following the acquisition agreement. The most significant growth was in Spain, where Grifols' expanded its workforce by 4.2% to 3,798 people. The talent pool grew in North America by 1.4% to 13,861 employees and by 2.6% in ROW (rest of the world) to 1,005 employees.

The average seniority of Grifols' personnel is 5.9 years and the average age is 37.8; more than 58% of employees are younger than 40. In terms of gender, women make up 58% of the workforce, while men comprise 42%.

Grifols continues its efforts to attract and retain talent. Occupational health and safety, and continuous training and development were the main areas of focus for the first half of 2018. Training and development initiatives centered on technical training programs, onboarding initiatives for new employees, performance reviews and leadership development.

Safety initiatives included a behavior-based management program that aims to interweave safety issues organization-wide, as the company works toward standardizing safety and health programs throughout the group.

Environmental management

Environmental management is one of the main pillars of the group's corporate responsibility actions.

Grifols continues to make significant progress on its 2017-2019 Environmental Plan, whose principal objectives include reducing the consumption of electricity, natural gas and water, in addition to improving waste management and recovery.

In the first half of 2018, external audits based on the ISO 14001 standard were carried out in the Diagnostic Division's facilities in Emeryville, California (U.S.) with satisfactory results. More than 75% of Grifols' production plants are ISO-14001-certified. In addition, the plants in Spain and the United States have adopted the new 2015 version of this international environmental management standard.

Worth highlighting are two recent environmental accolades awarded to Grifols' manufacturing complex in Clayton, North Carolina (U.S.). These facilities earned the highest distinction possible in the Environmental Stewardship Initiative, which promotes the development and implementation of innovative solutions that reduce the impact on the environment beyond mere legal compliance. The Clayton complex's office building also became the first in Johnston County to receive the Leadership in Energy and Environmental Design (LEED) in the Silver category for its socially responsible design.

Transparency: Grifols voluntarily discloses transfers of value to health professionals and healthcare organizations

In 2015, Grifols voluntarily adopted the Code of Conduct on Industry Interactions with Healthcare Professionals and Healthcare Organizations of the European Federation of Pharmaceutical Industries and Associations (EFPIA) in alignment with its commitment to transparency. For the third consecutive year, the company disclosed all payments and other transfers of value to health professionals and health sector organizations in 33 European countries, including Spain.

In Europe, Grifols' transfer of value totaled EUR 11.7 million in 2017, compared to EUR 11.8 million in 2016. The group's R+D+i activities in Spain accounted for 60% of total transfers of value in Europe.

Although EFPIA applies to medicines, Grifols voluntarily expanded its scope to include transfers unrelated to medications and those made by its three main divisions. Grifols applies this policy of transparency in the United States as stipulated by the regulatory body (Centers for Medicaid and Medicare Services, or CMS).

Committed to patients: more than 25 million international units of clotting factors donated to the World Federation of Hemophilia (WFH)

Grifols has collaborated with the World Federation of Hemophilia (WFH) for more than a decade, supporting its efforts to improve access to treatment for bleeding disorders around the world.

The donation forms part of Grifols' 2014 commitment to donate at least 200M IU of factor VIII to the WFH Humanitarian Aid Program over a span of eight years. To date, this initiative has improved access to care and treatment for patients with bleeding disorders in 47 developing countries.

Investor and Analyst annual meeting

The company hosted its annual investor and analyst meeting in Barcelona in June 2018. Grifols executives summarized results of the different divisions and outlined the group's capex plans, primary research projects and financial performance.

Grifols included in the FTSE4Good index

Grifols was selected for inclusion in the FTSE4Good sustainability index, specifically, the FTSE4Good Global, FTSE4Good Europe and FTSE4Good Ibex indices.

The sustainability indices or ESG indices rate companies on their environmental, social and corporate governance (ESG) performance, in addition to their financial indicators.

REGARDING FINANCIAL INFORMATION: The financial information corresponding to the first half of 2018 included in this document forms part of the information provided by the company.

REGARDING NON-FINANCIAL INFORMATION: The 2017 Corporate Responsibility Report offers an overview of the most relevant economic, environmental and social impacts in Grifols' value chain and their influence on stakeholder decisions in accordance with Global Reporting Initiative (GRI) information requirements and recommendations. Published in May 2018, the report is available on Grifols' corporate website: www.grifols.com.

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About Grifols

Grifols is a global healthcare company with more than 75 years of legacy dedicated to improving the health and well-being of people around the world. Grifols produces essential plasma-derived medicines for patients and provides hospitals and healthcare professionals with the tools, information and services they need to help them deliver expert medical care.

Grifols' three main divisions -Bioscience, Diagnostic and Hospital- develop, produce and market innovative products and services that are available in more than 100 countries.

With a network of 225 plasma donation centers, Grifols is a leading producer of plasma-derived medicines used to treat rare, chronic and, at times, life-threatening conditions. As a recognized leader in transfusion medicine, Grifols offers a comprehensive portfolio of diagnostic products designed to support safety from donation through transfusion. The Hospital Division provides intravenous (IV) therapies, clinical nutrition products and hospital pharmacy systems, including systems that automate drug compounding and control drug inventory.

Grifols is headquartered in Barcelona, Spain and has 18,300 employees in 30 countries.

In 2017, sales exceeded 4,300 million euros. Grifols demonstrates its strong commitment to advancing healthcare by allocating a significant portion of its annual income to research, development and innovation.

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 US NASDAQ via ADRs (NASDAQ:GRFS).

For more information, visit www.grifols.com

PROFIT AND LOSS ACCOUNT

<i>In thousands of euros</i>	1H 2018	1H 2017	% Var
NET REVENUE (NR)	2,120,118	2,192,447	(3.3%)
COST OF SALES	(1,113,858)	(1,089,246)	2.3%
GROSS MARGIN	1,006,260	1,103,201	(8.8%)
% NR	47.5%	50.3%	
R&D	(112,247)	(121,575)	(7.7%)
SG&A	(387,771)	(443,789)	(12.6%)
<i>OPERATING EXPENSES</i>	<i>(500,018)</i>	<i>(565,364)</i>	<i>(11.6%)</i>
OPERATING RESULT (EBIT)	506,242	537,837	(5.9%)
% NR	23.9%	24.5%	
<i>FINANCIAL RESULT</i>	<i>(103,188)</i>	<i>(147,583)</i>	<i>(30.1%)</i>
SHARE OF RESULTS OF EQUITY ACCOUNTED INVESTEEES	(5,729)	(10,295)	(44.4%)
PROFIT BEFORE TAX	397,325	379,959	4.6%
% NR	18.7%	17.3%	
INCOME TAX EXPENSE	(79,442)	(102,589)	(22.6%)
<i>% OF PRE-TAX INCOME</i>	<i>20.0%</i>	<i>27.0%</i>	
CONSOLIDATED PROFIT FOR THE YEAR	317,883	277,370	14.6%
RESULT ATTRIBUTABLE TO NON-CONTROLLING INTERESTS	(1,096)	(491)	123.2%
GROUP PROFIT FOR THE PERIOD	318,979	277,861	14.8%
% NR	15.0%	12.7%	

GROUP PROFIT RECONCILIATION

<i>In millions of euros</i>	1H 2018	1H 2017	% Var
REPORTED GROUP PROFIT	319.0	277.9	14.8%
% NR	15.0%	12.7%	
Amortization of deferred financial expenses	27.1	33.5	(19.1%)
Amortization of intangible assets acquired in business combinations	19.0	18.7	1.7%
Non-recurring items and associated with recent acquisitions	-	19.5	
Tax impacts of amortization adjustments	(9.2)	(19.4)	(52.5%)
ADJUSTED⁽¹⁾ GROUP NET PROFIT	355.9	330.2	7.8%
% NR	16.8%	15.1%	

⁽¹⁾ Excludes non-recurring items and associated with recent acquisitions, amortization of deferred expenses associated to the refinancing and amortization of intangible assets related to acquisitions

CASH FLOW

In thousands of euros

	1H 2018	1H 2017
REPORTED GROUP PROFIT	318,979	277,861
DEPRECIATION AND AMORTIZATION	107,958	106,549
NET PROVISIONS	(24,464)	(279)
OTHER ADJUSTMENTS AND OTHER CHANGES IN WORKING CAPITAL	9,310	38,774
CHANGES IN INVENTORIES	(139,046)	(64,217)
CHANGES IN TRADE RECEIVABLES	(19,391)	59,135
CHANGES IN TRADE PAYABLES	(4,161)	(39,260)
<i>CHANGE IN OPERATING WORKING CAPITAL</i>	<i>(162,598)</i>	<i>(44,342)</i>
NET CASH FLOW FROM OPERATING ACTIVITIES	249,185	378,563
BUSINESS COMBINATIONS AND INVESTMENTS IN GROUP COMPANIES	(255,406)	(1,813,163)
CAPEX	(102,080)	(135,269)
R&D/OTHER INTANGIBLE ASSETS	(28,754)	(10,887)
OTHER CASH INFLOW / (OUTFLOW)	56,790	20,467
NET CASH FLOW FROM INVESTING ACTIVITIES	(329,450)	(1,938,852)
FREE CASH FLOW	(80,265)	(1,560,289)
ISSUE / (REPAYMENT) OF DEBT	(19,790)	1,723,945
DIVIDENDS (PAID) / RECEIVED	(140,168)	(95,274)
OTHER CASH FLOWS FROM/(USED IN) FINANCING ACTIVITIES	(1,110)	(151,374)
NET CASH FLOW FROM FINANCING ACTIVITIES	(161,068)	1,477,297
TOTAL CASH FLOW	(241,333)	(82,992)
CASH AND CASH EQUIVALENTS AT THE BEGINNING OF THE YEAR	886,521	895,009
EFFECT OF EXCHANGE RATE CHANGES IN CASH AND CASH EQUIVALENTS	23,311	(61,799)
CASH AND CASH EQUIVALENTS AT THE END OF THE PERIOD	668,499	750,218

BALANCE SHEET

ASSETS

<i>In thousands of euros</i>	June 2018	December 2017
NON-CURRENT ASSETS	8,486,246	7,974,948
GOODWILL AND OTHER INTANGIBLE ASSETS	6,335,669	5,859,840
PROPERTY PLANT & EQUIPMENT	1,819,289	1,760,053
INVESTMENTS IN EQUITY ACCOUNTED INVESTEEES	225,781	219,009
NON-CURRENT FINANCIAL ASSETS	38,448	69,889
OTHER NON-CURRENT ASSETS	67,059	66,157
CURRENT ASSETS	2,947,373	2,945,316
INVENTORIES	1,806,765	1,629,293
TRADE AND OTHER RECEIVABLES	415,744	386,410
OTHER CURRENT FINANCIAL ASSETS	21,059	10,738
OTHER CURRENT ASSETS	35,306	32,354
CASH AND CASH EQUIVALENTS	668,499	886,521
TOTAL ASSETS	11,433,619	10,920,264

EQUITY AND LIABILITIES

<i>In thousands of euros</i>	June 2018	December 2017
EQUITY	3,971,244	3,633,965
CAPITAL	119,604	119,604
SHARE PREMIUM	910,728	910,728
RESERVES	2,452,375	2,027,648
TREASURY STOCK	(55,441)	(62,422)
INTERIM DIVIDENDS	0	(122,986)
CURRENT YEAR EARNINGS	318,979	662,700
OTHER COMPREHENSIVE INCOME	221,155	93,807
NON-CONTROLLING INTERESTS	3,844	4,886
NON-CURRENT LIABILITIES	6,446,685	6,308,312
NON-CURRENT FINANCIAL LIABILITIES	6,023,747	5,901,815
OTHER NON-CURRENT LIABILITIES	422,938	406,497
CURRENT LIABILITIES	1,015,690	977,987
CURRENT FINANCIAL LIABILITIES	205,095	155,070
OTHER CURRENT LIABILITIES	810,595	822,917
TOTAL EQUITY AND LIABILITIES	11,433,619	10,920,264

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