Investors and Analysts Meeting

Barcelona, March 5, 2009



Forward Looking Statements

This presentation contains forward-looking statements based on current assumptions and forecasts made by Grifols Group Management

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 Madrid Stock Exchange.

The company assumes no liability whatsoever to update these forward-looking statements or to conform them to future events or developments.



Presentation Guidelines

- ➡ Following the Spanish Stock Exchange (CNMV) Guidelines for investor meetings the information included in this presentation has been already filed in the CNMV
- → The Q&A session must be focused on the content of this presentation, including explanations and/or clarifications
- Questions related to relevant information not included in this presentation can not be addressed
- ➡ It is Grifols' investor relation policy not to provide with financial guidance in addition to the information contained in this presentation



2008 Financials



2008 Main events

- Purchase of land and buildings for the industrial expansion in Parets del Vallés, where the most important industrial assets of the Group in Spain are located
- ➤ A plot of land (5,100 m²) adjacent to our industrial site in Parets has been acquired, for the future fractionation capacity expansion.
- Agreement with Stough Development for building new plasma centers.
- Acquisition of a plasma center located in San Diego (USA)
- Launch and Sale of "Q"®, a new coagulometer instrument.
- → "Club Deal" agreement, with 24 banks, amounting to €350 million to refinance
 the existing syndicated loan and to fund the expansion Plan for the period
 2008-2012.
- FDA approves new facilities, in building 325, for sterile filling and lyophilization of coagulation factors. Consequently, the South Filling Area is no longer operational
- ➤ The PID Clinical trial of Flebogamma DIF® 10% is concluded. A second clinical trial to include a new indication for the treatment of ITP has started.



2008 Main events

- In the AGM, dated June 13th, a dividend increase was approved moving the pay-out from 28% up to 40% of net income.
- ➤ With the aim of achieving excellence in the operation of our donor centers, the company has opened the Grifols Academy of Plasmapheresis (Glendale, AZ) in January 2009.
- Fibrin Glue clinical trial ready to begin.
- ➤ Grifols PediGri® On Line available now in US for all Grifols hemoderivatives. PediGri® offers full traceability from plasma donation to final product.
- Acquisition of a new corporate offices building, currently under construction, that will strengthen corporate culture and better management alignment.
- ➤ LA City Hall granted permits to build the new Flebogamma DIF® purification plant, which is expected to be operating by 2013. Construction started in Oct 2008.
- First sale of DG Gel® cards in France after the expiration of the Diamed European patent.
- Obtained the Registration Approval of Flebogamma DIF® in Australia. Q4 2008

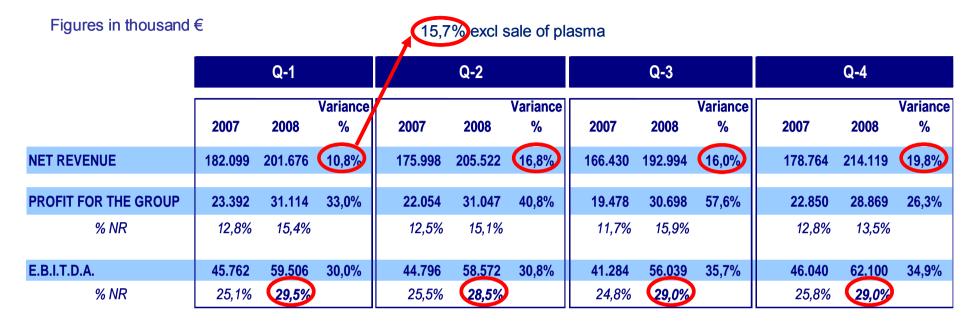


Strong 2008 earnings delivery

EUR '000	2007 Actual	2008 Actual	Variance vs Tot		Constant Rate 19,9%
NET REVENUES	703.291	814.311	<u>111.020</u>	15,8%	
COST OF SALES	387.632	416.127	28.495	7,4%	
GROSS MARGIN % NR	315.659 44,9% +400	398.184 48,9%	<u>82.525</u>	26,1%	
R & D - Technical area S.G.&A.	28.725 140.580	28.494 166.729	(231) 26.149	(0,8%) 18,6%	
OPERATING EXPENSES % NR	<u>169.305</u> 24,1%	195.223 24,0%	<u>25.918</u>	15,3%	
E.B.I.T. % NR	146.354 20,8%	202.961 24,9%	<u>56.607</u>	38,7%	
Financial expenses (income) Interest receivables Exchange (Gain) / Loss	20.798 (2.630) 4.618	29.870 (1.979) 2.825	9.071 652 (1.793)	43,6% (24,8%) (38,8%)	
FINANCIAL RESULT	<u>22.786</u>	<u>30.716</u>	<u>7.930</u>	34,8%	
PROFIT BEFORE TAXES	123.587	<u>172.269</u>	<u>48.682</u>	39,4%	
Tax expenses % tax rate	35.239 28,5%	50.153 29,1%	14.914	42,3%	
PROFIT FOR THE GROUP % NR	87.774 12,5%	121.728 14,9%	33.954	38,7%	
E.B.I.T.D.A. Profit % NR	177.882 25,3%	236.217 29,0%	<u>58.335</u>	32,8%	
FO1.6					7



Consistent 2008 quarterly results

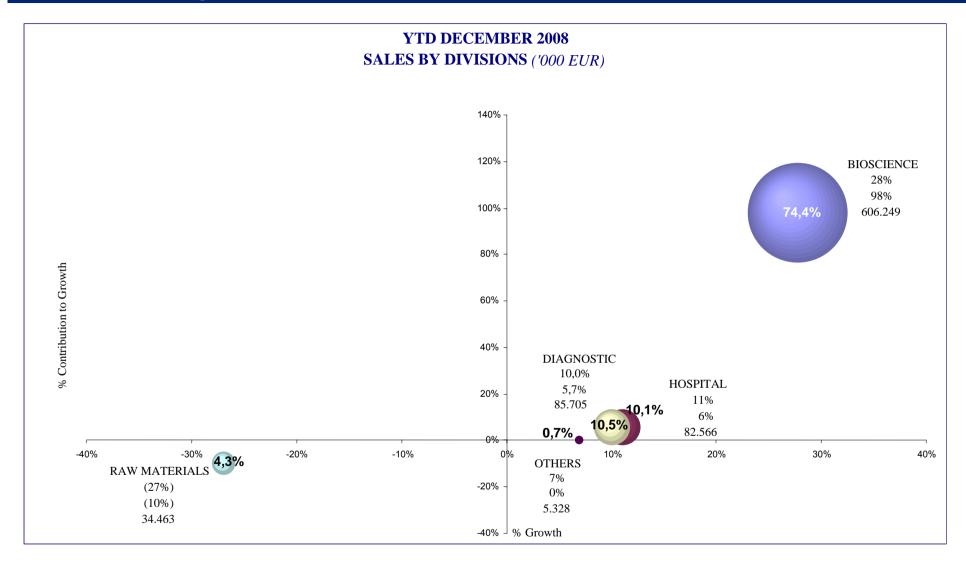


Figures in thousand

	YTD			
	2007	2008	Variance %	
NET REVENUE	703.291	814.311	15,8%	
PROFIT FOR THE GROUP	87.774	121.728	38,7%	
% NR	12,5%	14,9%		
E.B.I.T.D.A.	177.882	236.217	32,8%	
% NR	25,3%	29,0%		



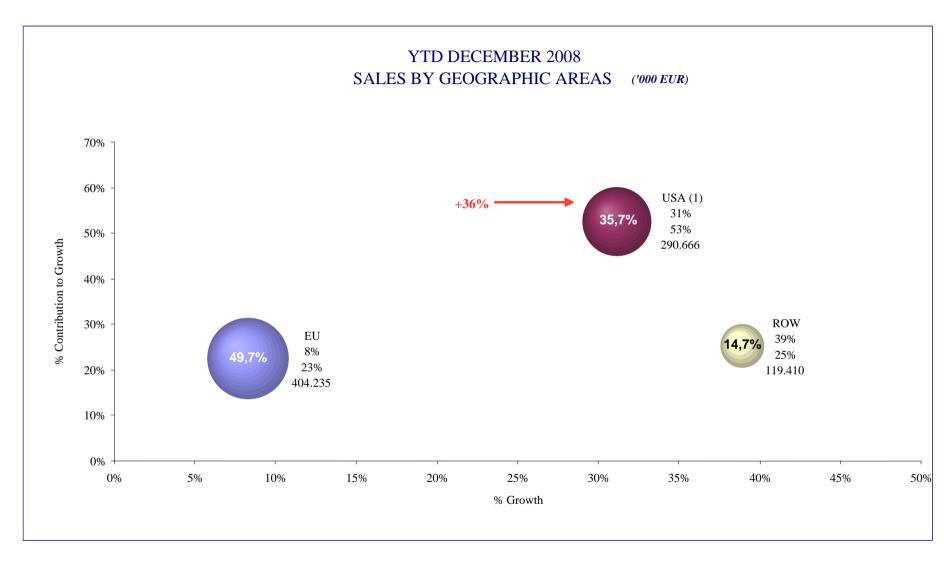
Robust sales growth across divisions



Growth and Contrib. at constant FX.



USA and ROW leading geographical growth

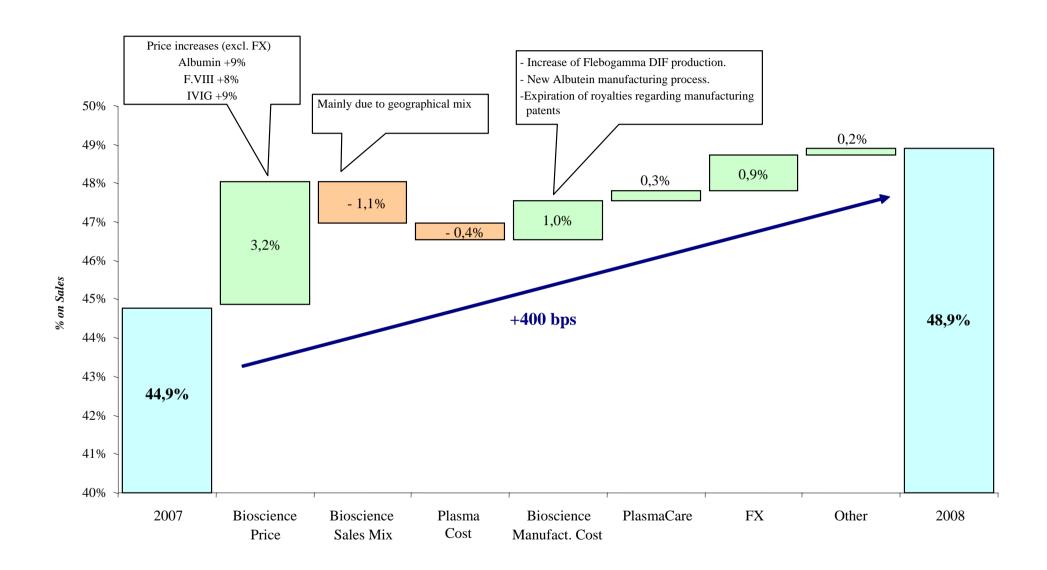


Growth and Contrib. at constant FX.

(1) Non-recurrent plasma sales in 2007 of €8,1 MM.

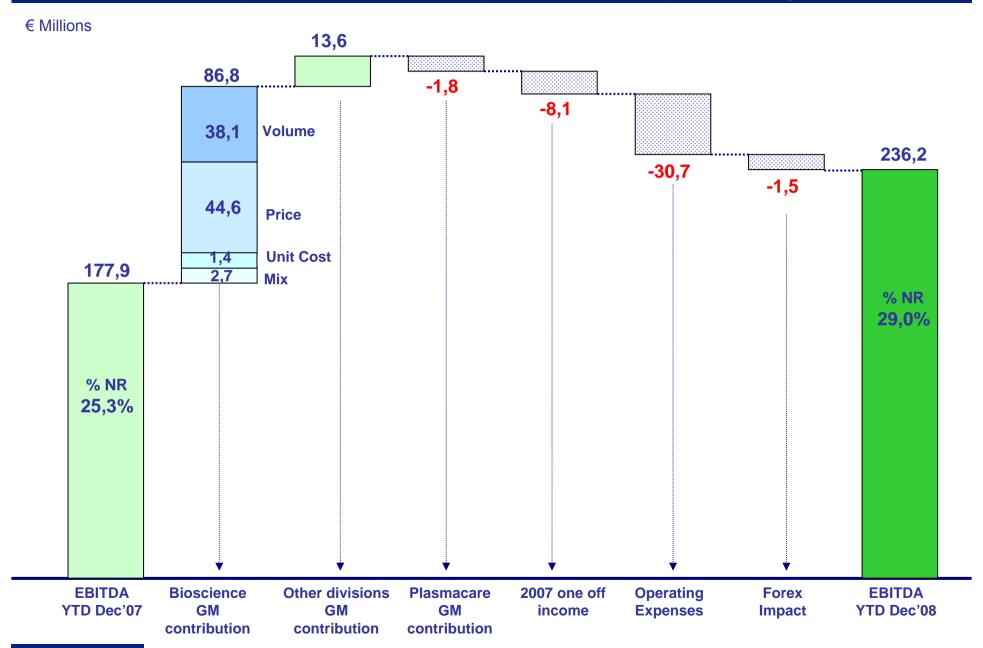


Price increase as a key driver of Gross Margin expansion





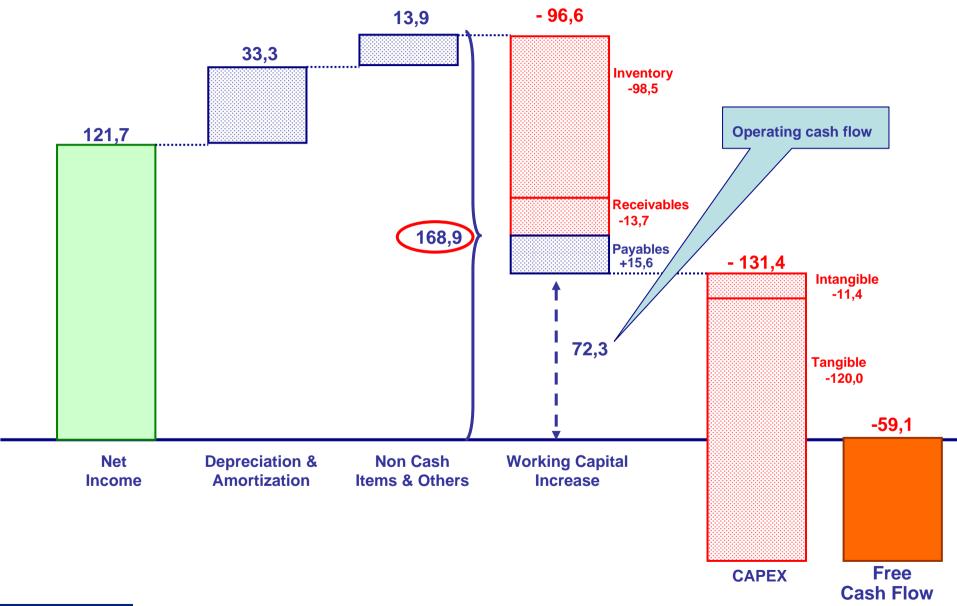
Immaterial FX impact in EBITDA improvement vs previous year





Solid cash flow generation pre - Inventory and Capex







Capex and Dividends as main uses of funds

€ Million

SOURCES			USES	
Net IncomeDepreciation	121,7 33,3		- CAPEX	131,4
Others Working Capital Increase	13,9 -96,6	.	- Treasury stock	4,2
- Operating Cash Flow	72,3		- Dividends	34,8
- Net Debt Increase	102,9	,	- Others	4,8
Total	175,2			Total 175,2
	=====			=====



Strong balance sheet with reasonable leverage

December 07	December 08
343,2	446,0
1,9	1,9
0,89	0,86
7,8	7,7
	343,2 1,9 0,89

V	Vorking Capital Ratios		
		December 07	December 08
S	tocks Turnover (Days)	255	327
D	ays sales outstanding (DSO)	90	83
D	ays payable outstanding (DPO)	66	65



GRIFOLS stock performance vs IBEX 35



Var. IBEX LTM: -43,9%

Var. IBEX 2009: -23,9%

•March 3rd, 2009

CORPORATE ACTIVITY



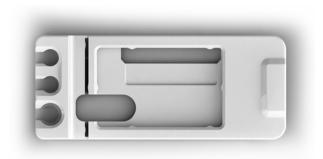
THE RATIONALE FOR GRIFOLS ACQUISITION OF THE HOLDING.

- Expand our product range in IM by adding a different and revolutionary technology.
- Gain access to all segments of the IM market including the "High throughput donors segment".
- Build the more complete and advanced range of products for Blood Typing and Pre-transfusion Diagnostics available in the market.
- Advance and accelerate Grifols entry into the USA IM Market.
- Establish a strong direct presence in Australia and New Zealand and increase Grifols visibility in the Asia Pacific region.

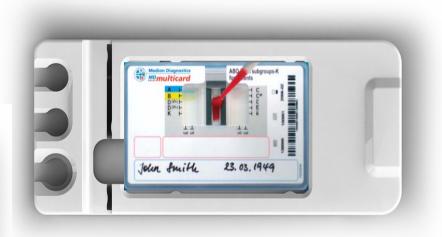






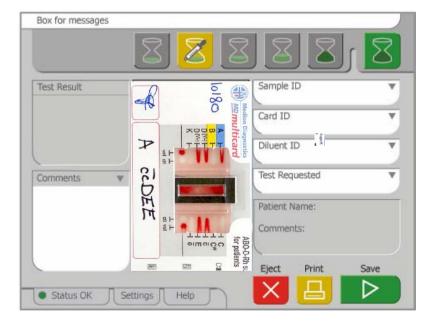








































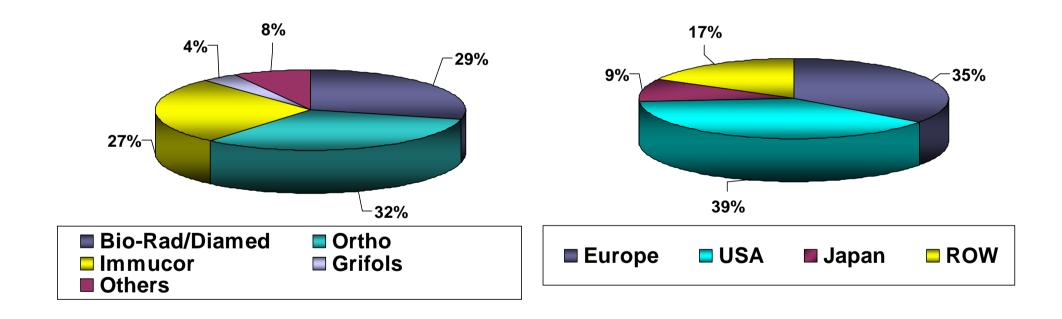
CONTRIBUTION OF THE NEW ACQUISITION TO THE GRIFOLS BUSINESSES

- The turnover of the new entity will represent close to 20 Million € (first 12 month of sales) of new businesses.
- In the first three years we will be looking at doubling these sales.
- As soon as the new Technology can be automated, the market potential for it will grow dramatically.
- The combination of Grifols and Medion current products together with the future joint developments will represent the most complete and advanced offer to approach the whole market of Blood Typing Worldwide.



IMMUNOHEMATOLOGY WORLDWIDE MARKET

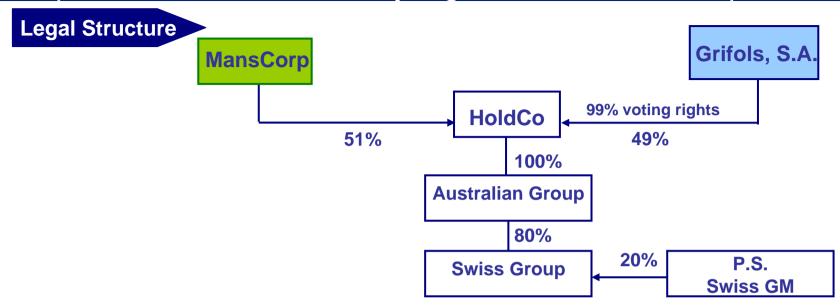
2008 Total Market size: 1 Billion USD



Source Grifols Data



Antipodes Transaction – Key legal and Financial aspects



Sell Option

- ✓ Any party shall have the option to sell their shares to the other shareholder.
- ✓ Option notice shall include price, date and place.
- ✓ Exercise periods: 5 / 10 years (stand still period of 5 years).

Management ✓ Grifols will have majority on the board.

Distribution

✓ Grifols Australia-Swiss distribution and Medion global distribution agreement.

Investment

- ✓ Grifols shares subscription in HoldCo for € 25 Million.
- ✓ € 10 Million will remain in cash to be used to support growth.



FUTURE OUTLOOK

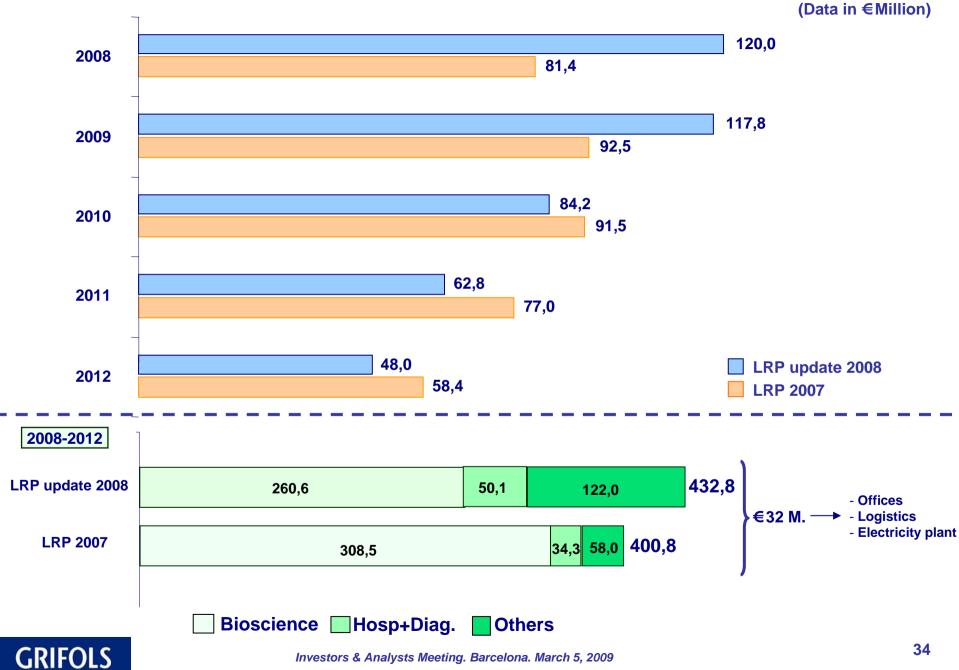
Short / Long term Outlook Wrap-up

- ✓ Product availability along with a continuous growth demand in developed and emerging countries, will drive <u>sales increase and market share gain</u>.
- ✓ In a very complicated macroeconomic environment <u>maintaining</u> or minor increasing in <u>selling prices</u> will be a positive achievement.
- ✓ <u>Plasma supply</u> evolution is closely monitored to properly manage any potential excess or shortage of plasma.
- ✓ Full commitment to <u>increase R&D budget</u> to support future developments.
- ✓ As a result of gross margin slowdown, EBITDA will keep at similar levels as previous year.
- ✓ Although <u>currency fluctuations</u> on the long run have no relevant impact on earnings, quarterly performance may be influenced.
- ✓ Financial gearing will be maintained at a very reasonable level.



LRP CAPEX UPDATE

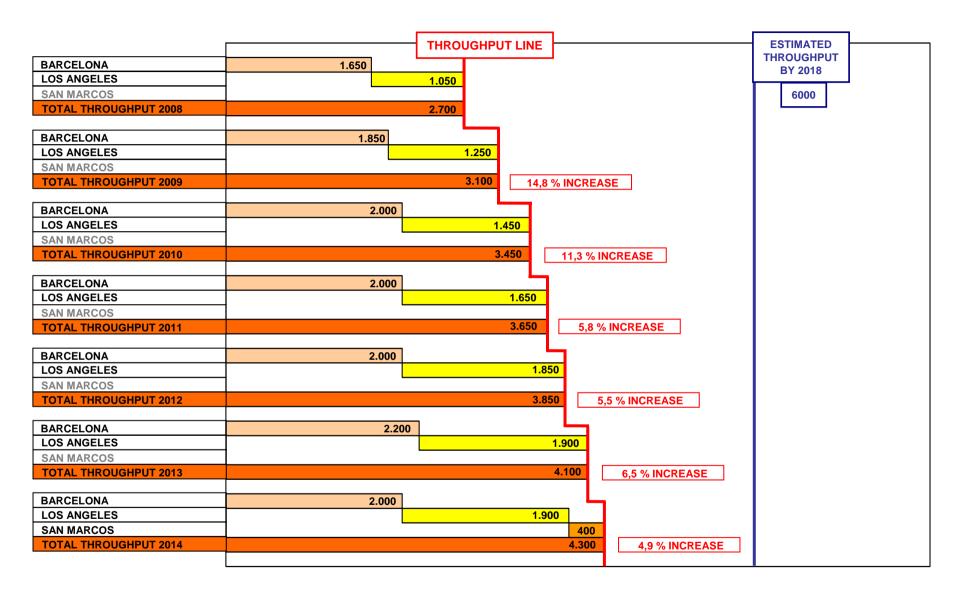
LRP Capex Update: Progress on track



Bioscience investment rationale

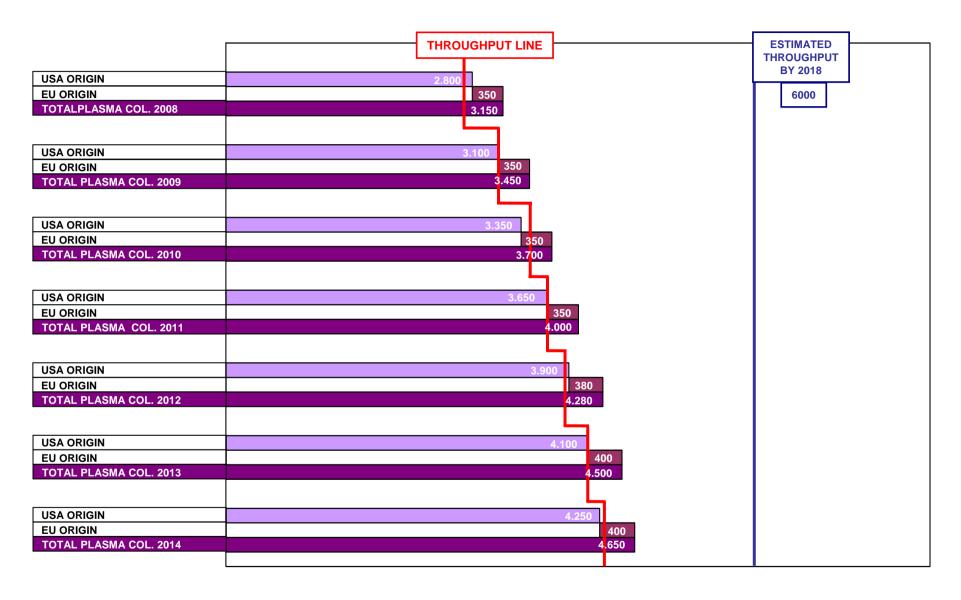


PLASMA THROUGHPUT EVOLUTION (2008-2014)



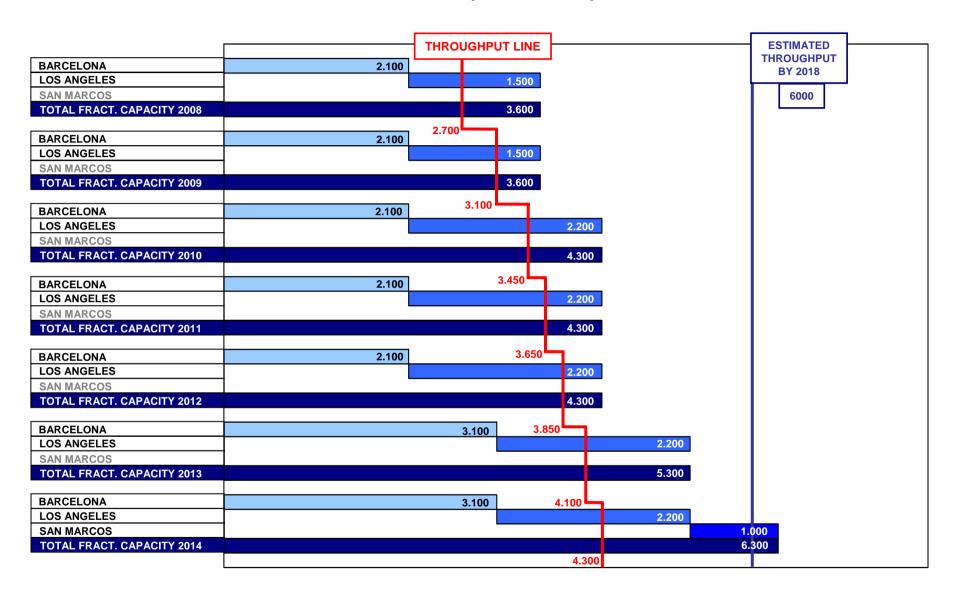


PLASMA COLLECTIONS EVOLUTION (2008-2014)



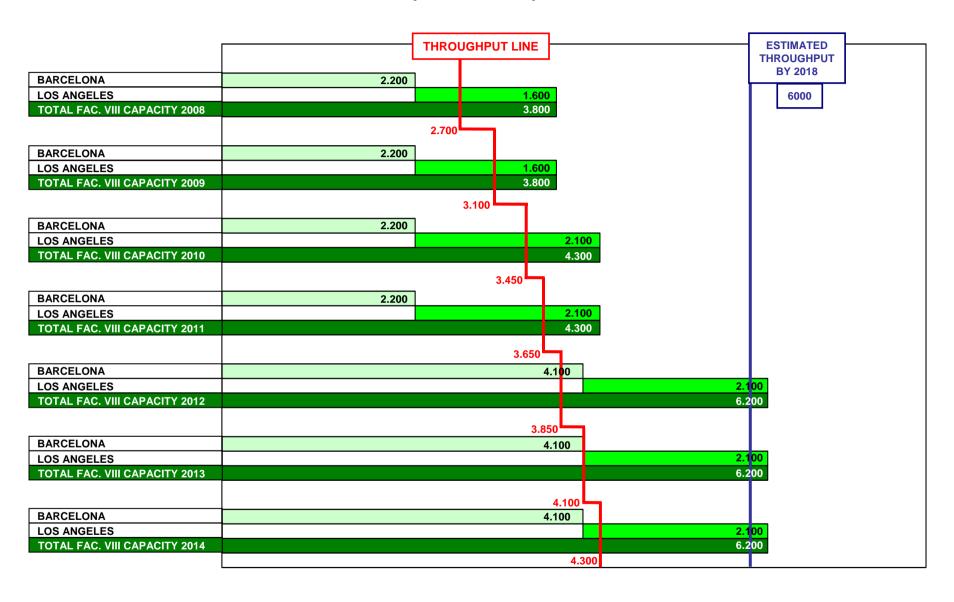


FRACTIONATION CAPACITY EVOLUTION (2008-2014)



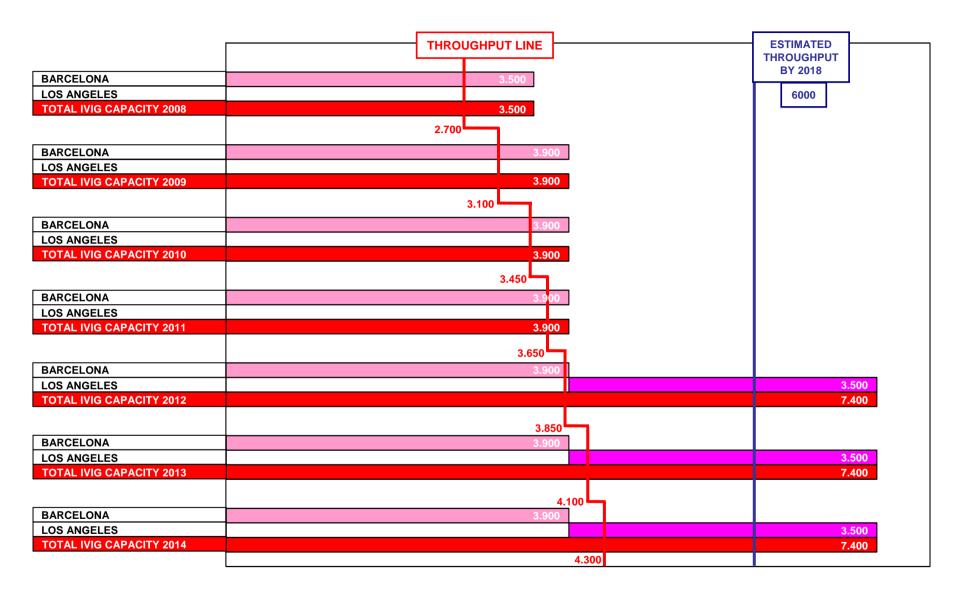


FACTOR VIII CAPACITY EVOLUTION (2008-2014)



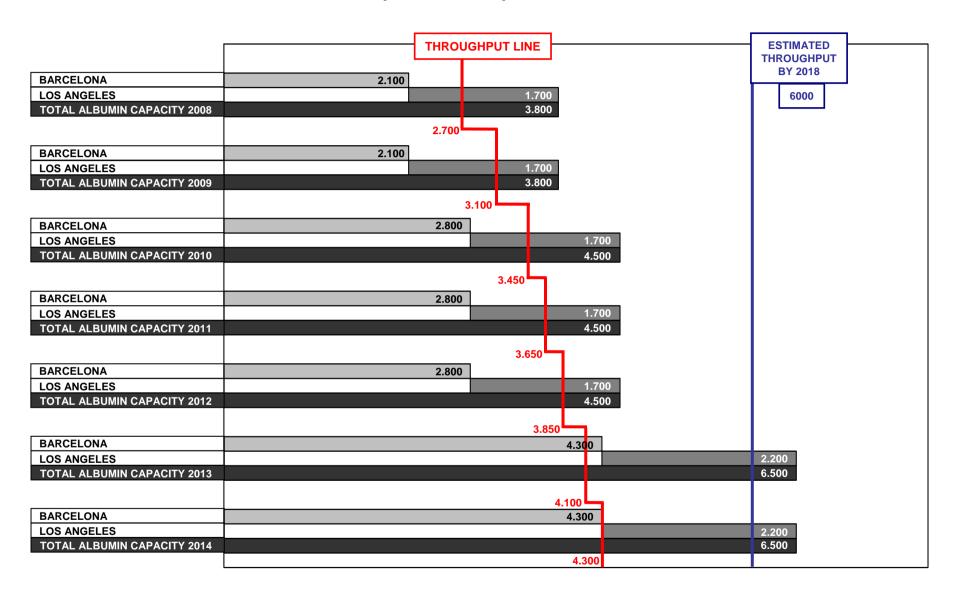


IVIG CAPACITY EVOLUTION (2008-2014)





ALBUMIN CAPACITY EVOLUTION (2008-2014)

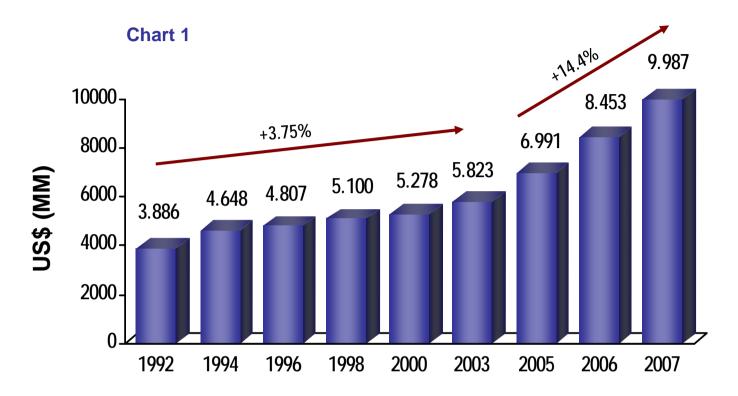




Market Dynamics & Trends



MARKET DATA. THE WORLDWIDE PLASMA DERIVATIVES MARKET



Sources: The Worldwide Plasma Fractions Market 2006, MRB - data 1992 to 2005. Grifols' estimate - data 2006 and 2007.

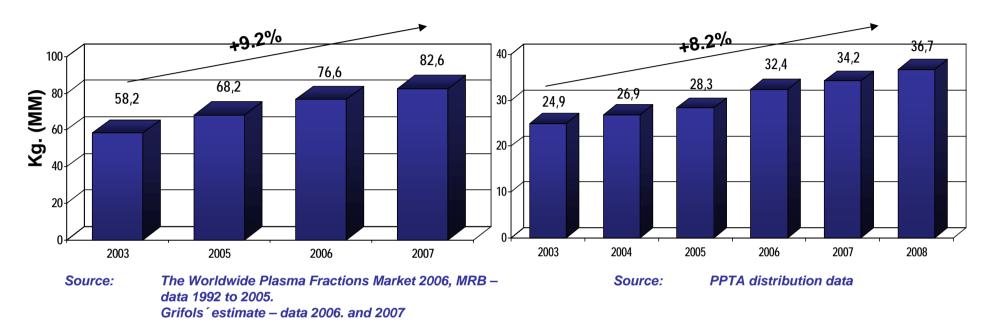
The world plasma derivatives market has grown consistently since early 1990s. Since 2003, worldwide sales growth has accelerated

Market growth continues at an accelerated rate.



SUPPLY AND DEMAND OF PLASMADERIVATIVES. IVIG

Chart 2. Evolution of the Worldwide market of IVIG Chart 3. Evolution of the U.S.A. market of IVIG



- Demand of IVIG continues to growth consistently in and outside the US.
- Growth drivers do not change:
 - New indications.
 - Emerging markets.

Supply-Demand of IVIG remains in balance



CURRENT MARKET CONDITIONS (I)

PRICING

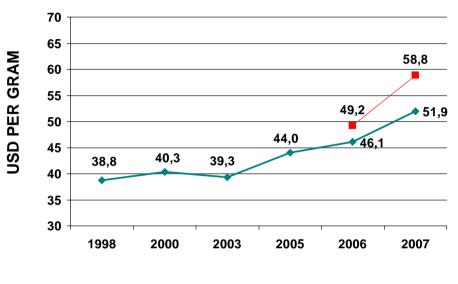
- Historical data for 2007 confirm positive price evolution for all products in all markets.
- Grifols continues to be positioned with ASP's equal or above the average.
- Grifols has continued leading the Albumin price recovery, especially in the US.
- Grifols prices in 2008 have continued to go up with increases ranging from 5% to 15% depending on product and geography.
- Market prices to our knowledge have performed similarly during last year.

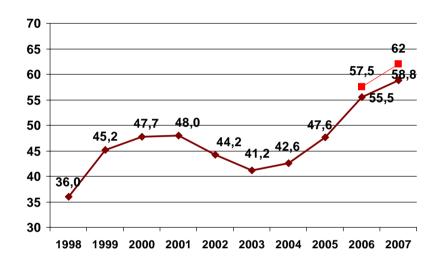


CURRENT MARKET CONDITIONS (II). Pricing IVIG

Chart 4. IVIG. Evolution of Average Sales Price (USD) per gram. Worldwide

Chart 5. IVIG. Evolution of Average Sales Price (USD) per gram. U.S.A.





Grifols prices

Grifols prices

Source: The Worldwide Plasma Fractions

Market 2006, MRB – data 1998 to

2005.

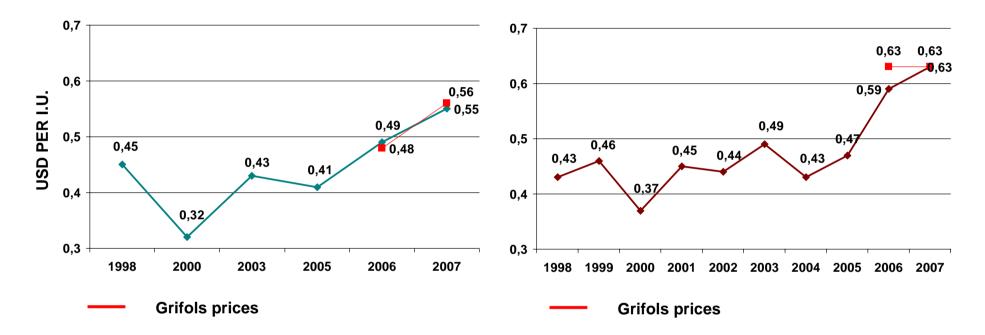
Grifols' estimate - data 2006 and 2007.

Source: The Plasma Fractions Market in United States 2007, MRB.



CURRENT MARKET CONDITIONS (III). Pricing pd Factor VIII

Chart 6. Factor VIII. Evolution of Average Sales Price Chart 7. Factor VIII. Evolution of Average (USD) per IU. Worldwide Sales Price (USD) per IU. U.S.A.



Source: The Worldwide Plasma Fractions

Market 2006, MRB – data 1998 to

2005.

Grifols´estimate – data 2006 and 2007.

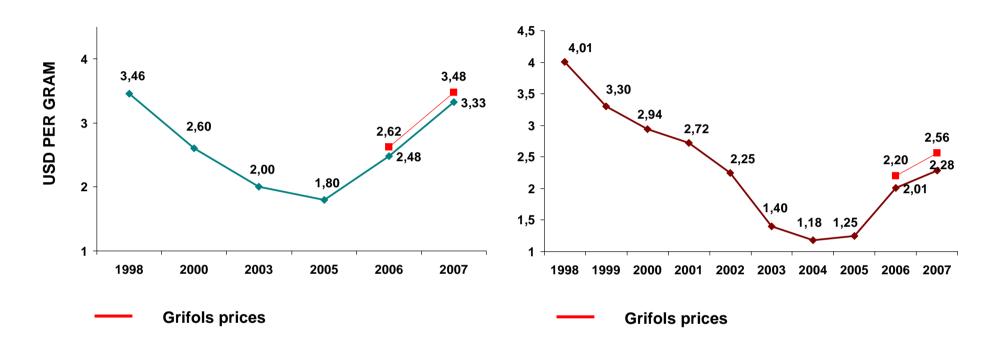
Source: The Plasma Fractions Market in United States 2007, MRB.



CURRENT MARKET CONDITIONS (IV). Pricing Albumin

Chart 8. Albumin. Evolution of Average Sales Price (USD) per gram. Worldwide

Chart 9. Albumin. Evolution of Average Sales Price (USD) per gram. U.S.A.



Source: The Worldwide Plasma Fractions

Market 2006, MRB – data 1998 to

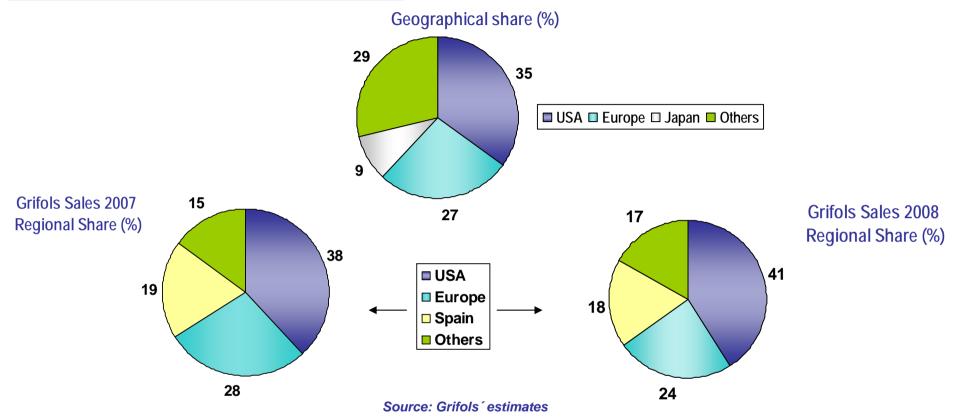
2005.

Grifols estimate - data 2006 and 2007.

Source: The Plasma Fractions Market in United States 2007, MRB.



MARKET DATA. THE WORLDWIDE PLASMA DERIVATIVES MARKET. World Total Sales 2007 US\$ 9,987 (MM)

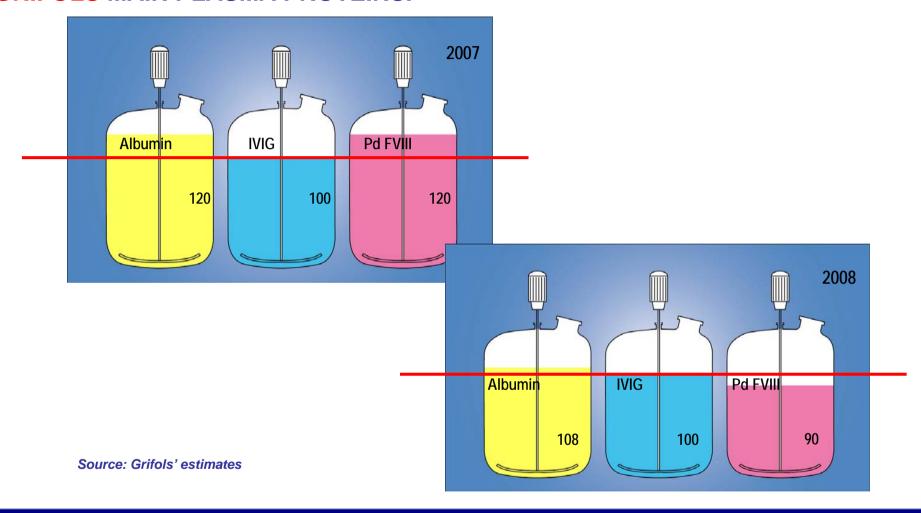


- Grifols geographical sales distribution has reached certain balance. Current share of US is already sufficient due to recent focus.
- Starting in 2009 Europe and ROW markets should obtain better product allocation.
- Opportunities in Emerging Markets will provide substantial contribution to the 2009 growth.

Grifols geographical focus to change to a more balanced growth profile



COMPARISON OF AMOUNT OF PLASMA NEEDED TO MANUFACTURE GRIFOLS MAIN PLASMA PROTEINS.

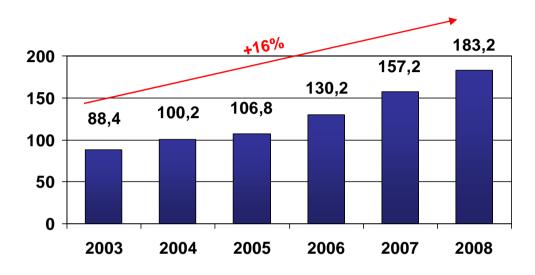


Grifols is now prepared to access new growing markets with an improved Factor VIII availability

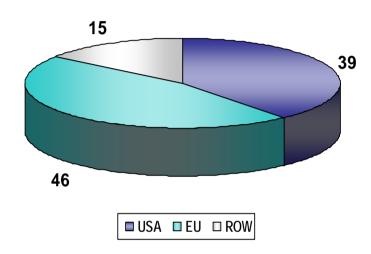


SUPPLY AND DEMAND OF PLASMA DERIVATIVES. pd FACTOR VIII

Grifols worldwide Factor VIII Sales (MM USD)



Geographical Distribution of Grifols Factor VIII Sales 2008 (I.U.)



Inhibitor eradication and von Willebrand treatment continue to be the drivers of the demand for pdFVIII in developed markets like the USA and Europe

Rest of the world represents 46 % of the world pd Factor VIII market in volume

Increasing Plasma throughput and current availability will allow Grifols to access the fast growing Hemophilia markets in emerging countries.



FUTURE MARKET CONDITIONS / GRIFOLS FUTURE OUTLOOK (I)

IVIG

- Global demand expected to grow over 7% per annum.
- Grifols will continue to convert to Flebogamma DIF® with special focus in Europe and Australia during 2009.
- Increased plasma throughput will also provide more product to market.
- Grifols increased availability should be absorbed by increasing demand.
- Grifols will continue to grow market share gradually in all markets.



FUTURE MARKET CONDITIONS / GRIFOLS FUTURE OUTLOOK (II)

PLASMA DERIVED FACTOR VIII

- Inhibitor treatment and VonWillebrand indication will continue to be the focus in developed markets like US and Europe.
- Improved product availability will allow us to address the fast growing emerging markets.

ALBUMIN

- Grifols may continue on allocation for Albumin for all 2009
- Publications about the use of Albumin in cirrhotic patients, particularly a publication expected in June 09 in the UK may stimulate the demand in several countries.
- Consequently, we anticipate prices to remain strong.



PROJECTS INVOLVING NON-CORE HEMODERIVATIVES

- Fibrin Sealant. Clinical Trial started 4Q 2008. FDA and European submission forecasted by 2011.
- AT New indications and New Markets. Clinical Trial Cardiac Surgery Phase II Start Date (Q1 2009). FDA and European submissions by 2013.
- PTC. New formulation including nanofiltration under development.

Increasing the income per liter of plasma as a way to improve profitability



NEW PRODUCTS TO BE LAUNCHED OVER THE NEXT TWO YEARS

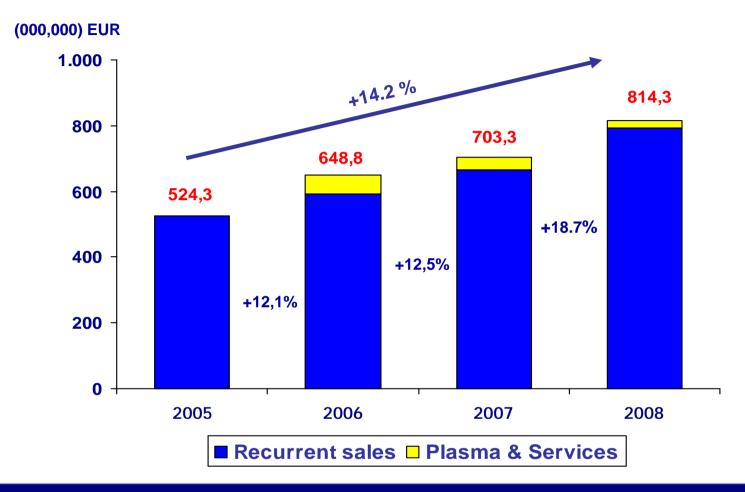
Bioscience

- Niuliva® (Hepatitis B IV Immunoglobulin) Indication, Liver Transplant. Launch date Q3 2009. Initially Italy and Spain, Latin-America one year later. Market potential 50 M €. Market share target 10% in 3 years.
- Flebogamma DIF® 5% Launches in several markets (Phase 1 in Australia, Portugal, Germany, UK Ireland, Holland, Phase 2 in Spain, and Italy and Phase 3 in other Markets: Latin America, Asia, etc).
- Flebogamma DIF® 10% Launch in the US Market Q1 2010. Launch in Europe Q1 2011.

New products contributing to the short term growth of Bioscience Division



TOTAL GRIFOLS SALES EVOLUTION



Three consecutive years of remarkable growth Sales at constant currency increased by 20 % in 2008



2009 SALES GROWTH PATTERN MAY BE DIFFERENT FROM PREVIOUS YEARS

- Sales growth in 2008 and 2007 was strongly based on increased selling prices.
 Volume in 2008 started to play a significant role especially for IVIG.
- Sales growth in 2009 will come mainly from volume, new markets for Bioscience products and new businesses development such as Diagnostics. Prices will play a very minor role.
- While price increases apply in many markets since January 1st, volume growth and new businesses progress more gradually.
- As a result of all the above we should expect a quarterly sales distribution pattern for 2009 slightly different from the one of previous years.

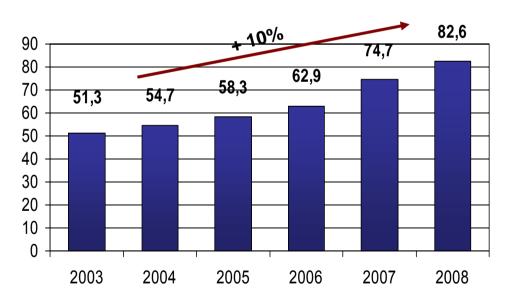
Sales growth in 2009 will take place in a progressive way giving a different quarterly distribution pattern



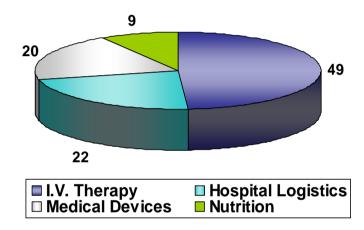
HOSPITAL DIVISION

- I.V. Therapy fluids and sterile compounding devices.
- Hospital Logistics: Automated drugs dispensing Systems, Hospital warehousing and software applications..
- Enteral and Parenteral Nutrition.
- Medical Devices.

Domestic business mainly



Hospital Division sales growth evolution



Hospital Division sales by business segment 2007



HOSPITAL DIVISION

New Products and New Businesses Opportunities for the next two years

- Enteral and Parenteral Nutrition product line extension
- New Grifols Engineering products and technologies applied to Hospital Logistic product range.
- Hospital logistics sales may be influenced in the next few years by budget constraints of health care systems.
- Grifols Partnership, third party manufacturing program to bring new growth opportunities.
- Distribution businesses in Spain continues to be very strong in the Medical Devices area with new products and opportunities.

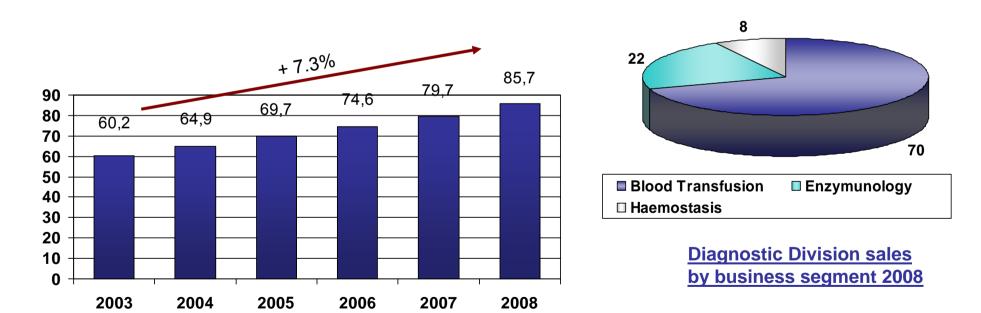
Hospital Division continues betting for a sustained growth



DIAGNOSTIC DIVISION

- Grifols activity is focused on the areas of Blood Transfusion, haemostasis and enzymoimmunoanalysis.
- Strong Instrumentation R&D and manufacturing capabilities.
- Grifols is the worldwide reference for Gel Cards technology automation.
- Reagents development in immunohematology, haemostasis and Elisa.

We position ourselves in market niches where we can become global market players with "world class" product ranges.



Diagnostic Division sales growth evolution



DIAGNOSTIC DIVISION. NEW PRODUCTS AND NEW BUSINESSES OPPORTUNITIES FOR THE NEXT TWO YEARS.

- DG Gel Gradual businesses expansion in territories where former patent expired in 2008. (France, UK Italy, Germany, etc.)
- Erytra® Immunohematology Analyzer. Launch 2010, Extension of Immunohematology Instrumentation Product line. Gaining access to higher throughput customers
- Q® Coagulometer introduction in different markets together with Haemostasis Reagents range.
- Swiss-Australian acquisition to support and accelerate immunohematology growth.

The basis for a continued growth of Diagnostic Division is established



Research and Development



BIOSCIENCE

SAFETY NEW PRODUCTS YIELD & EFFICIENCY REGULATORY

DIAGNOSTIC

INSTRUMENTS REAGENTS REGULATORY

GRIFOLS R & D

HOSPITAL

NEW PRODUCTS
CONTAINERS & PLASTICS
DEVICES & EQUIPMENTS

ENGINEERING

FACILITIES
PROCESSES
EQUIPMENT



BIOSCIENCE

SAFETY NEW PRODUCTS YIELD & EFFICIENCY REGULATORY

DIAGNOSTIC

INSTRUMENTS REAGENTS REGULATORY

GRIFOLS R & D

HOSPITAL

NEW PRODUCTS
CONTAINERS & PLASTICS
DEVICES & EQUIPMENTS

ENGINEERING

FACILITIES
PROCESSES
EQUIPMENT

R & D BIOSCIENCE SAFETY (I)

> NANOFILTRATION (PATHOGEN REMOVAL)

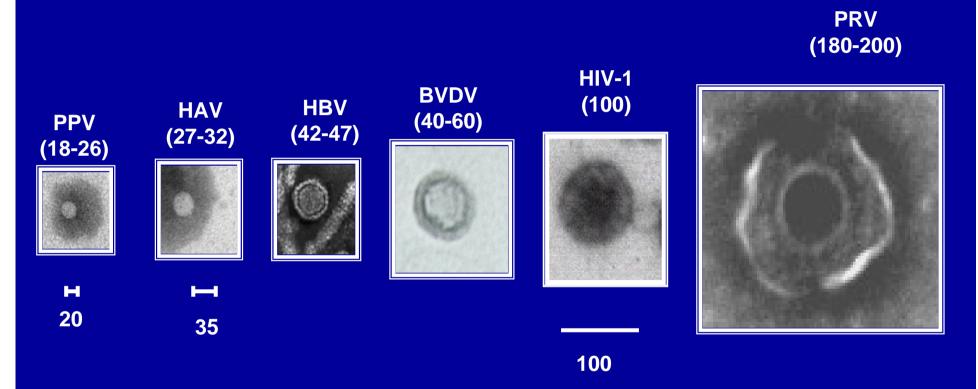
 ALBUMIN 20 nm WORK IN PROCESS FACTOR VIII VWF 20 nm WORK IN PROCESS FACTOR IX 15 nm OK PROTHROMBIN COMPLEX 15 nm WORK IN PROCESS 15 nm OK ANTITHROMBIN FIBRINGGEN 20 nm OK THROMBIN 15 nm OK • IVIG (FLEBOGAMMA DIF®) 20 nm OK A1 ANTITRYPSINE 15 nm OK

> INACTIVATION

• KNOWN PATHOGENS : CONTINUED STUDIES
• EMERGING PATHOGENS : WNV, AVIAN FLU, DENGUE, SARS



Relative sizes of viruses (in nm). Images from electron microscopy



Original PPV, HAV and PRV, EM pictures from Drs X. Abad and A. Bosch, University of Barcelona, Spain

PPV: Porcine Parvovirus.,**HAV:** Hepatitis A virus.,**HBV**: Hepatitis B virus., **BVDV**: Bovine Viral Diarrhoea virus., **HIV-1**: Human Immunodeficiency virus type 1., **PRV**: Pseudorabies virus. Sizes from Fields in Virology, 4th edition, 2001. Ed.: J. Pine, A. Snyder and R.E. Cook. Lippincott Williams & Wilkins.



R & D BIOSCIENCE SAFETY (II)

> POTENTIAL TSE RISK REDUCTION/ELIMINATION

DURING DONOR SELECTION

• DURING FRACTIONATION: STUDIES COMPLETED AND

SUBMITTED. INFECTIVITY

REDUCTION AFTER DIFFERENT

MANUFACTURING STEPS.

• DURING PURIFICATION: STUDIES COMPLETED AND

SUBMITTED

• DURING NANOFILTRATION: THIRD PARTIES STUDIES

PUBLISHED



R & D BIOSCIENCE. SAFETY (III)

> PLASMA

DONOR QUALIFICATION

• TESTING : REDUNDANT VOLUNTARY NAT

TESTING BEYOND REGULATIONS

(PCR, HEP A AND C, PARVO)

• INVENTORY HOLD AND LOOK BACK: PPTA GUIDELINE, VOLUNTARY

COMPLIANCE.

• HANDLING : CHANGING REMOVAL OF

POSITIVE UNITS AT EACH

CENTER TO REMOVAL AT

CENTRALIZED TEMPLE FACILITY,

UPON FDA APPROVAL OF NEW

SOFTWARE)



R & D BIOSCIENCE. SAFETY (IV)

> PEDIGRI®

• SAMPLE LIBRARY : OK IN EUROPE (1987)

BEING IMPLEMENTED IN USA (Q3

2009)

• PLASMA TRACEABILITY: OK

• FRACTION TRACEABILITY: OK

• LOT AND NUMBER IDENTIFICATION IN

EACH BOTTLE: OK

• ONLINE PUBLIC INFORMATION FOR OK IN EUROPE

PROFESSIONALS ONLY: OK IN USA (SEPT 2008)



BIOSCIENCE

SAFETY NEW PRODUCTS YIELD & EFFICIENCY REGULATORY

DIAGNOSTIC

INSTRUMENTS REAGENTS REGULATORY

GRIFOLS R & D

HOSPITAL

NEW PRODUCTS
CONTAINERS & PLASTICS
DEVICES & EQUIPMENTS

ENGINEERING

FACILITIES
PROCESSES
EQUIPMENT

R & D BIOSCIENCE. NEW PRODUCTS (I)

> POTENTIAL NEW INDICATIONS FOR EXISTING PRODUCTS (I)

• FVIII/VWF VON WILLEBRAND OK IN USA & EUROPE

(ITALY & UK)

• FVIII/VWF INHIBITOR ERRADICATION

• HEP B IVIG LIVER TRANSPLANTATION

(IMMUNOTOLERANCE)

FIRST LICENSE EXPECTED

DURING 2008.

LICENSE IN ITALY GRANTED Q4 2008

STUDY ONGOING

• A1-PI FIBROMIALGIA CLINICAL STUDY ONGOING

IN 2008.

FINISHED, LACK OF

EFFICACY.

• A1-PI CHRONIC FATIGUE PHASE II CLINICAL STUDY

ONGOING



R & D BIOSCIENCE. NEW PRODUCTS (II)

> POTENTIAL NEW INDICATIONS FOR EXISTING PRODUCTS (II)

• ALBUMIN ALZHEIMER CLINICAL STUDY

ONGOING.

SUBMISSION OF INTERIM RESULTS IN JULY 2009.

• ALBUMIN CIRRHOSIS CLINICAL STUDY ONGOING

• IVIG ALZHEIMER CLINICAL STUDY PENDING

SUBMISSION.

CLINICAL STUDY

ONGOING (JAN 09)

• AT CARDIOPULMONARY CLINICAL STUDY ONGOING



R & D BIOSCIENCE. NEW PRODUCTS (III)

> NEW FORMULATIONS & PROCESSES FOR EXISTING PRODUCTS

ALBUMIN NEW FORMULATION FOR STEM CELL CULTURE

• FVIII / FIX ALTERNATIVE FORMULATIONS, LONGER SHELL

LIFE

ALBUMIN NEW CONTAINER, Q4 2009

• FLEBOGAMMA DIF® 10 % CLINICAL TRIAL CONCLUDED, DATA SUBMITTED.

EXPECTED APPROVAL Q4 2009.

> NEW REGISTRATIONS FOR EXISTING PRODUCTS.

• AT IN USA CLINICAL TRIAL ONGOING

• FVIII VWF VON WILLEBRAND, REST OF THE WORLD

ONGOING

• A1-PI REST OF EUROPE, CLINICAL TRIAL SUBMISSION

Q4 2009

• FLEBOGAMMA DIF® 5% IN REST OF THE WORLD ONGOING



R & D BIOSCIENCE. NEW PRODUCTS (IV)

	NEW	/ PRO	TEINS
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• HEP B IVIG FIRST LICENSE EXPECTED IN 08

(ITALY AND SPAIN GRANTED IN DEC 08)

• FIBRIN SEALANT CLINICAL TRIALS TO START IN 2008 (ONGOING)

• FIBRINGEN CLINICAL TRIALS TO START IN 2009

• THROMBIN CLINICAL TRIALS TO START IN 2009



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R & D BIOSCIENCE. YIELD AND EFFICIENCY IMPROVEMENTS (I)

> PRODUCTION

• HARMONIZATION THE TWO FACILITIES TO USE THE SAME

METHODS

• YIELD IMPROVEMENT ALL PRODUCTS PRODUCED IN BOTH

FACILITIES. IT'S A "NEVER ENDING" PROCESS.



R & D BIOSCIENCE. YIELD AND EFFICIENCY IMPROVEMENTS (II)

> FACILITIES (I)

• ALBUTEIN L.A. NEW FACILITY (BLDG 317) TO PRODUCE

FRAC V BY "BARCELONA" METHOD.

SUBMISSION JUNE 09. APPROVAL EXPECTED

Q4 2009.

• ALBUTEIN L.A. NEW FACILITY (BLDG 325) FOR STERILE

FILLING.

SUBMISSION JUNE 09. APPROVAL EXPECTED

Q4 2009.

COAGULATION FACTORS

 L.A. NEW FACILITY FOR STERILE FILLING.

FDA APPROVED JULY 08 AND EMEA APPROVAL

EXPECTED MARCH 09.

NEW PURIFICATION BLDG 325 2ND PHASE ONGOING

FACTORS IN LOS ANGELES.

FACILITIES FOR COAG.



R & D BIOSCIENCE. YIELD AND EFFICIENCY IMPROVEMENTS (III)

> FACILITIES (II)

- SECOND PRODUCTION CONSTRUCTION STARTED OCT PLANT FOR FLEBOGAMMA 2008.COMPLETION EXPECTED Q2 2010 LOS ANGELES .
- NEW AREAS FOR ASCEPTIC COMPLETION EXPECTED Q4 2009 FILLING AND FREEZE DRYING IN BARCELONA.
- MINIFRAC.LOS ANGELES READY FOR SUBMISSION (Q2 2009, EXPECTED FRACTIONATION EXPANSION APPROVAL Q4 2009).



R & D BIOSCIENCE. YIELD AND EFFICIENCY IMPROVEMENTS (IV)

ENGINEERING

• PLASMA BOTTLES OPENING 2ND GENERATION, HIGHER EFFICIENCY AND

YIELD. ONGOING.

PLASMA HANDLING
 AUTOMATION OF THE WHOLE PROCESS

FROM RECEPTION TO CLASSIFICATION AND

SHIPPING.

BARCELONA: IN WORKING CONDITIONS

SINCE 2003

TEMPLE: IN WORKING CONDITIONS

SINCE JUNE 08



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R & D BIOSCIENCE. REGULATORY & PATENTS

> REGULATORY

ALTHOUGH NOT CONSIDERED AS AN R & D AREA, REGULATORY AFFAIRS DEPARTMENT IS RESPONSIBLE FOR THE PRODUCTION, FILE AND SUBMISSION OF ALL THE DOCUMENTATION GENERATED BY R & D: LICENSES OF NEW INDICATIONS, IN EXISTING OR NEW MARKETS, EXTENDED INDICATIONS OF EXISTING PRODUCTS IN EXISTING OR NEW MARKETS, SUBMISSION OF NEW FORMULATIONS OR EQUIPMENT CHANGES AMONG OTHERS.

REGULATORY AFFAIRS IS BASED AND CENTRALIZED IN BARCELONA WITH 30 EMPLOYEES. LOS ANGELES R.A. DEPARTMENT CONSISTS OF 5 EMPLOYEES, AND EACH OF OUR AFFILITATES HAS AT LEAST ONE PERSON DEVOTED TO REGULATORY AND REGULATORY COMPLIANCE.

> PATENTS

PATENTS REFLECT SOMEHOW THE EFFICIENCY AND SUCCESS OF R & D PROJECTS, SOME OF WHICH FAIL AND SOME SUCCEED.

PATENTS ARE DRAWN BY THE TEAM RESPONSIBLE FOR THE GIVEN PROJECT, BE IT EXTRACTION, OBTENTION OR PURIFICATION METHODS OF A PROTEIN OR THE DESIGN OF A NEW STERILE FILLING MACHINE OR A NEW INACTIVATION METHOD.

HOWEVER, A CENTRALIZED PATENT DEPARTMENT – WITH AN OUTSOURCED PATENT AGENT – CO-ORDINATES, FOLLOWS UP AND MAINTAINS THE PATENTS OWNED BY THE CORPORATION.



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R & D DIAGNOSTICS. INSTRUMENTS (I)

> IMMUNOHEMATOLOGY

WADIANA®: NEW SOFTWARE VERSION 3.2. LAUNCH Q3 2009

• ERYTRA®: NEW AUTOMATED IMMUNOHEMATOLOGY

ANALYZER.

EXPECTED LAUNCHING DATE AT ISBT BERLIN

CONGRESS, JUNE 2010

• F50 (AN INTERMEDIATE VERSION BETWEEN WADIANA®

AND ERYTRA®). LAUNCH 2012

NEW STAND- LAUNCH Q4 2010

ALONE COLOUR GEL CARD READER.

• NEW MULTICARD® (AUSTRALIA/SWITZERLAND). INITIAL STEPS.

AUTOMATED PROCESSOR.



R & D DIAGNOSTICS. INSTRUMENTS (II)

COAGULATION

• Q® COAGULOMETER LAUNCHED IN JUNE 2008

• Q® COAGULOMETER VERSION 2.0. LAUNCH IN JUNE 2009.

• NEW HIGH THROUGHPUT LAUNCH IN 2013 COAGULOMETER

> IMMUNOLOGY

• NEW IMPROVED TRITURUS®. LAUNCH IN 2012















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R & D DIAGNOSTICS. REAGENTS

> IMMUNOHEMATOLOGY

- CONTINOUS EVALUATION OF NEW COMMERCIAL ANTIBODIES FOR DG GEL
- INCORPORATION OF AUSTRALIA/SWITZERLAND RED CELL PANELS TO OUR EXISTING PORTFOLIO.

> COAGULATION

- NEW PT FORMULATION. Q4 2009
- SYNTHETIC LIPIDS APTT. Q4 2010
- THROMBIN/FIBRINGGEN, Q4 2011/Q4 2010
- CHROMOGENIC PROTEIN C. Q4 2010
- EVALUATION OF NEW METHODS TO OBTAIN FACTOR DEFICIENT PLASMAS (STARTING Q1 2009)

> IMMUNOLOGY

SERAQUEST DEVELOPMENT OF OWN AUTOIMMUNITY ELISA KITS.



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R & D DIAGNOSTIC REGULATORY & PATENTS

> **REGULATORY**

SAME COMMENTS AS FOR BIOSCIENCE DIVISION.

> PATENTS

SAME COMMENTS AS FOR BIOSCIENCE DIVISION.



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R & D HOSPITAL NEW PRODUCTS

> PARENTERAL NUTRITION NEW FORMULATIONS IN TRICAMERAL BAGS.

TWO NEW FORMULATIONS WHICH CONTAIN AMINOACIDS, LIPIDS AND GLUCOSE IN A THREE- CHAMBER BAG. LICENSING PROCESS ONGOING AND EXPECTED LAUNCH IN Q4 2010

> PARENTERAL NUTRITION NEW FORMULATION OF MEDIUM AND LONG CHAIN LIPIDS EMULSION.

LICENSING PROCESS ONGOING AND EXPECTED LAUNCH IN Q1 2010.

- > ENTERAL NUTRITION DIET FOR TREATMENT OF NON ALCOHOLIC FAT LIVER DISEASES. INITIAL PHASE.
- > LARGE VOLUME PARENTERALS
 - URIC ACID SOLUTION FOR STROKE TREATMENT IN CONJUNCTION WITH tPA
- > PARENTERAL SOLUTION FOR SURGICAL ARRYTHMIA PREVENTION (EMULSION OF MCT AND OMEGA 3).
- > OEM DEVELOPMENT OF DIFFERENT LARGE VOLUME PARENTERALS: PRE-DILUTED PARACETAMOL AND LEVOFLOXACINE



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R & D HOSPITAL CONTAINERS & PLASTICS

- > NON PVC CONTAINERS.
 - > ALL PRODUCTS HAVE BEEN RELICENSED TO NON PVC CONTAINERS
 - > NEW INJECTION PORE TO AVOID THE USE OF NEEDLES. SPECIALLY DESIGNED FOR CYTOSTATIC ADMINISTRATION. LICENSING PROCESS ONGOING.



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R & D HOSPITAL DEVICES & EQUIPMENTS

- > DIFFERENT DEVICES TO USE WITH GRIFOLS EQUIPMENTS
- > NEW BLOOD BAGS VERSIONS FOR THE COLLECTION OF STEM CELLS





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PATENTS SUBMITTED. 2005-2008

	2005	2006	2007	2008	TOTAL
HOSPITAL	1	1			2
DIAGNOSTIC		2	1	2	5
ENGINEERING	1	1	1		3
BIOSCIENCE		1	1	2	4
TOTAL	2	5	3	4	14 (*)

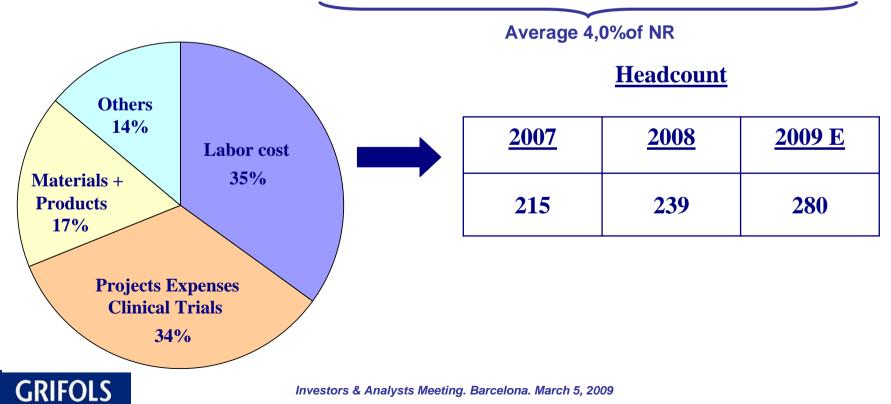
(*) First submission

Patents in force: 445 Patents in process : 129



Full commitment to increase R&D budget

	<u>2007</u>	<u>2008</u>	2009 E
R&D Expenses (Cash) MM €	29,4	30,6	44,5
% of NR	4,2%	3,8%	4,5%



Investors and Analysts Meeting

Barcelona, March 5, 2009

