



***A Specialty
Pharmaceutical Group***

GOLDMAN SACHS GLOBAL HEALTHCARE CONFERENCE, RANCHO PALOS VERDES, 9 JUNE 2015

A strategy of growth and geographical expansion

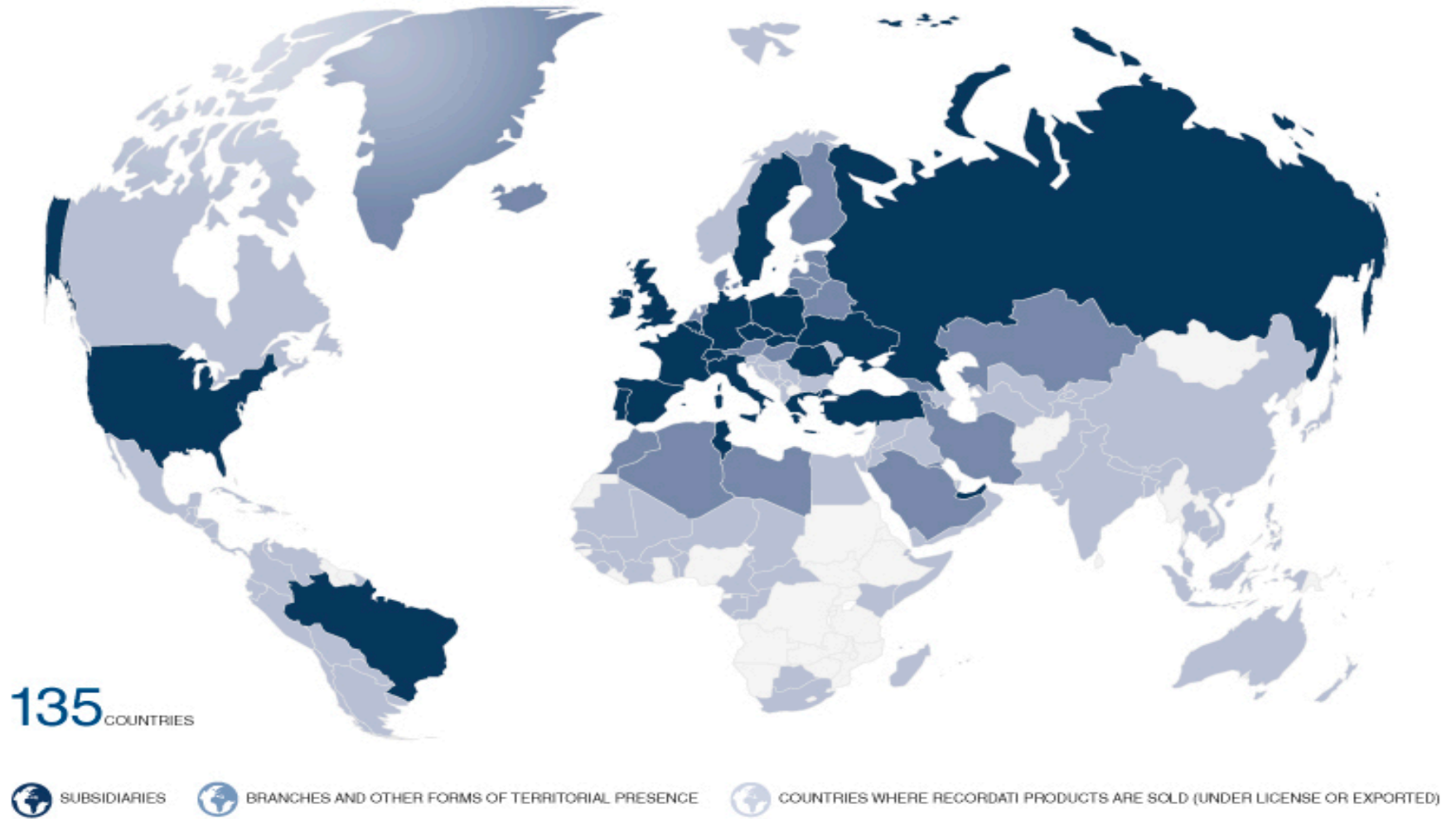
Profile

- An international specialty pharmaceutical group (€ 987.4 m sales in 2014 and 4,000 employees).
- Marketing operations in the main European markets, in Russia, Poland and other Central and Eastern European countries, in Turkey, in the U.S.A. and in North Africa
- Drugs for the treatment of rare diseases marketed worldwide
- Proprietary drugs sold worldwide through licensees
- R&D in specialty care and in treatments for rare diseases

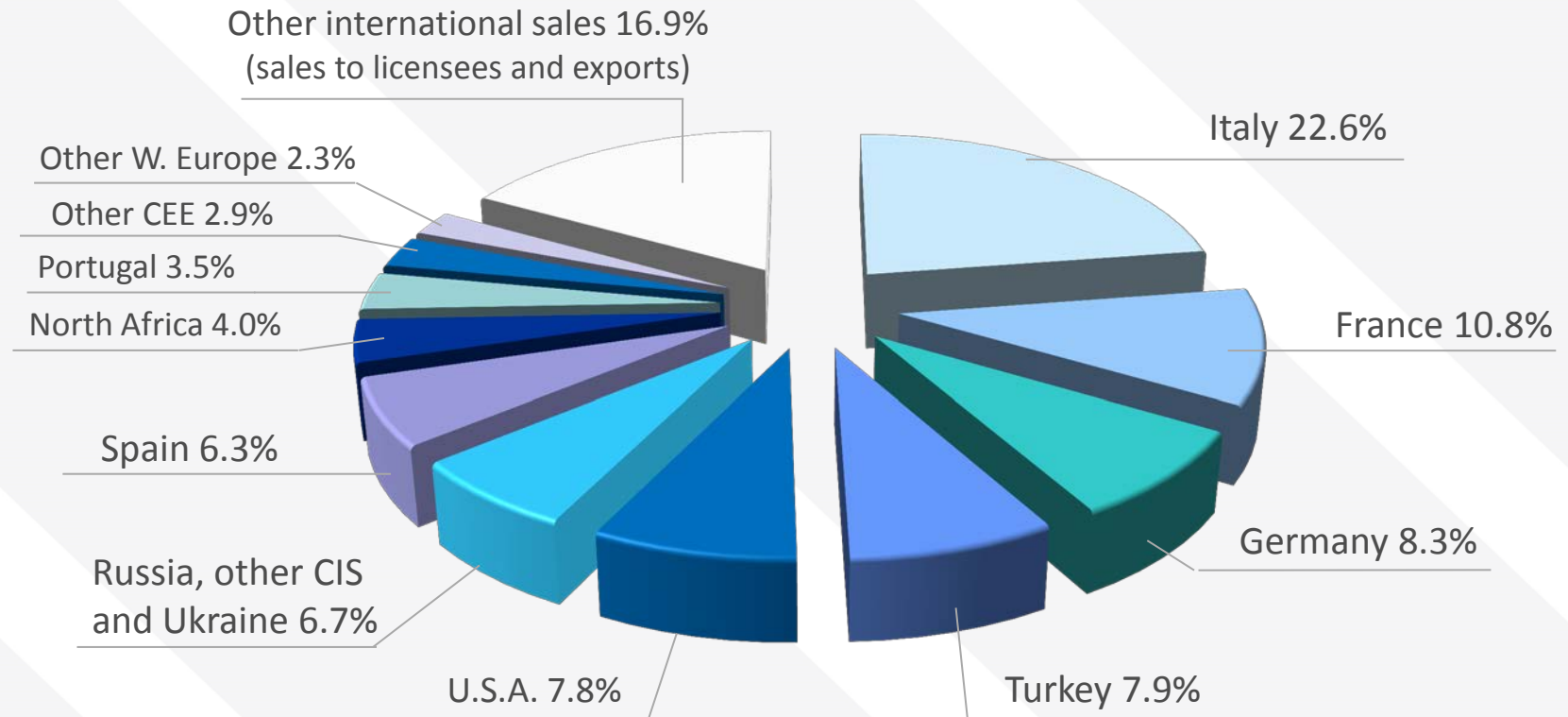
Strategy

- Expand through organic development and through acquisitions
- Develop product portfolio by enhancing product pipeline and new product acquisitions. Prioritize special care.
- Increase presence in new markets with high potential
- Treatments for rare diseases: develop a global presence

Extensive geographical footprint

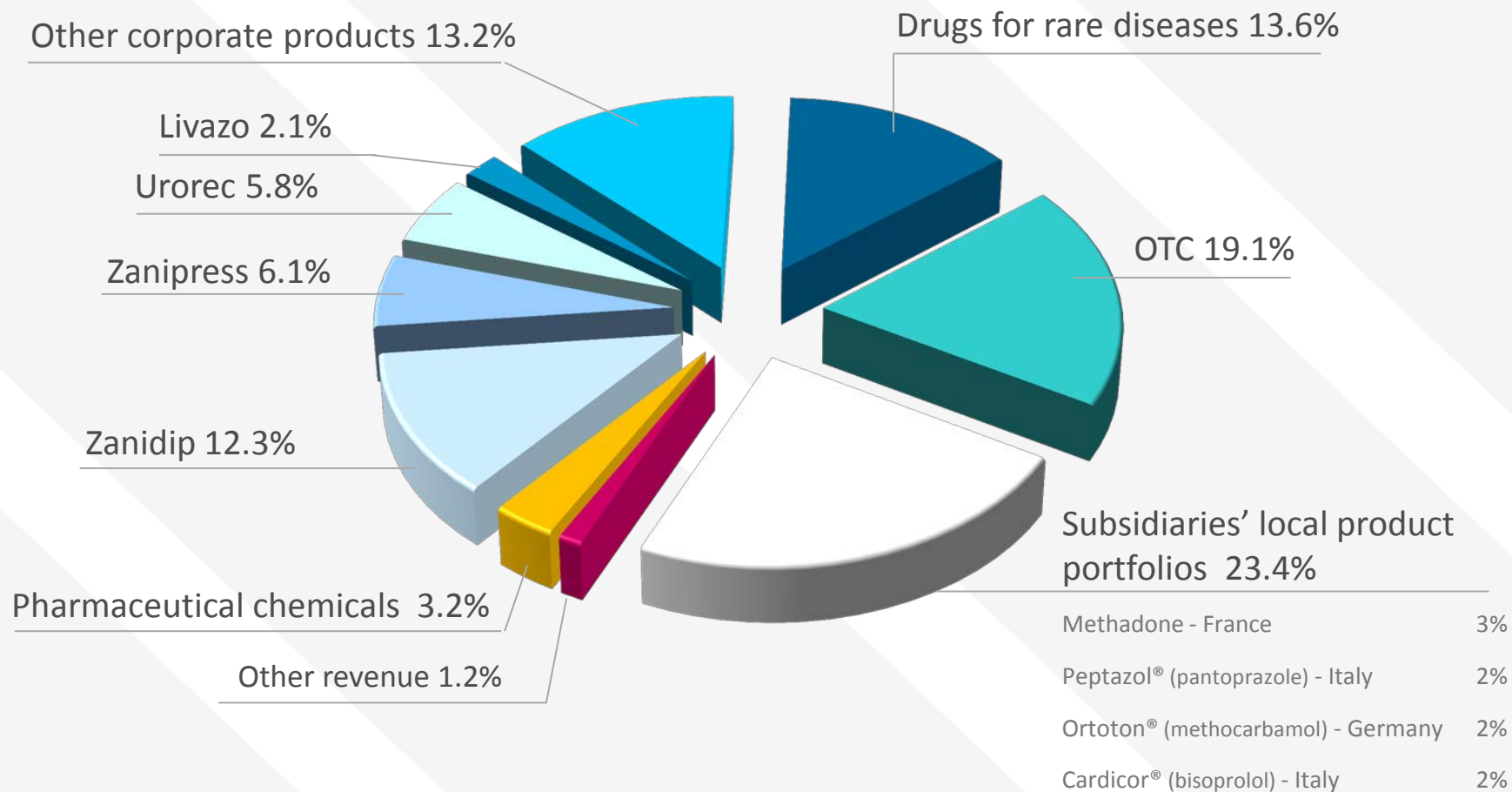


Geographical breakdown of pharmaceutical revenue



Data: First quarter 2015
Pharmaceutical revenue € 267.0 m

A diversified product portfolio



Data: First Quarter 2015
Total revenue € 275.7 m

Core corporate products

Zanipress[®] (lercanidipine+enalapril) **Zanidip[®] (lercanidipine)**

- Lercanidipine is a proprietary latest generation calcium channel blocker indicated for the treatment of hypertension. Enalapril is an ACE inhibitor indicated for the treatment of hypertension.
- Sales of Zanidip[®] (lercanidipine) have eroded (CAGR -12.7%) following its patent expiry at the beginning of 2010.
- Zanipress[®] (lercanidipine+enalapril) now launched in 25 markets and growing steadily
- Zanipress[®] to be rolled-out progressively in new markets
- Zanipress[®] prices will come under pressure due to generic competition
- Sales of the Zanidip[®] / Zanipress[®] franchise expected to remain at between € 160 and € 165 million in 2015 and to decrease to around € 140 million thereafter

Core corporate products

Urorec[®] (silodosin)

- Highly selective α_{1A} receptor antagonist indicated for the treatment of symptoms associated with benign prostatic hyperplasia (BPH).
- Fast onset of action. High efficacy. Very good cardiovascular safety.
- Launched in 28 markets: Armenia, Belarus, Belgium/Luxembourg, Bulgaria, Croatia, Cyprus, Czech Rep., France, Georgia, Germany, Greece, Ireland, Italy, Kuwait, Lebanon, Moldavia, the Netherlands, Poland, Portugal, Romania, Russia, Serbia, Slovakia, Spain, South Africa, Turkey, Ukraine and the United Arab Emirates. Further launches to take place.
- License and co-marketing agreements in place with important players
- BPH market in 16 main countries approx. € 0.8 billion

Core corporate products

Livazo[®] (pitavastatin)

- Highly effective HMG-CoA reductase inhibitor indicated for the treatment of hypercholesterolaemia.
- Thanks to its unique chemical structure Livazo[®] is a potent LDL-lowering drug with a consistent and progressive HDL-raising effect. (*Atherosclerosis Supplements 2010; 11:15-22*)
- Livazo[®], unlike most statins, is only minimally metabolized through a CYP pathway thereby reducing the risk of drug-drug interactions and providing a clear benefit in patients receiving polypharmacy. (*Atherosclerosis Supplements 2010; 11:15-22*)
- Launched in Spain, Portugal, Switzerland, Ukraine and Greece, further launches to take place.
- Statins market in the 11 key countries covered by the agreement is of around € 2.1 billion.

Drugs for rare diseases

A worldwide business

Huge market potential: > 6,000 rare diseases identified for which treatments exist for only around 300.

Progressive country introduction of rare disease plans and access to diagnostic tests will stimulate the market for orphan drugs.

- Acquisition of Orphan Europe end 2007. Establishment of Recordati Rare Diseases in the U.S.A. in 2013 following the acquisition of Lundbeck's U.S. portfolio of rare disease treatments.
- Present throughout Europe, Middle East and the U.S.. Sales coverage of new territories, either directly or through partnerships, ongoing.
- R&D in rare diseases:
 - Carbaglu[®], indication in organic acidemias in the U.S.A., phase III
 - Cystadrops[®], ocular cystinosis, filed in the EU
 - Grasp[®], partnership with Erytech, development of indication in Acute Myeloid Leukemia, phase II b
- Sales of drugs for rare diseases in 2014 total € 123.2 million. CAGR of 18.8% over the past six years

Drugs for rare diseases

NORMOSANG[®] (EU-RoW) /**PANHEMATIN**[®] (US) (human haemin), used to treat acute attacks of hepatic porphyria

CARBAGLU[®] (carglumic acid), indicated in the treatment of hyperammonaemia due to NAGS deficiency and to the main organic acidemias

COSMEGEN[®] (dactinomycin), used mainly in the treatment of three rare cancers, Wilms' tumor, childhood rhabdomyosarcoma and choriocarcinoma.

PEDEA[®] (EU-RoW)/**NEOPROFEN**[®] (US) (ibuprofen I.V.), indicated in the treatment of patent *ductus arteriosus*

CYSTADANE[®] (betaine anhydrous), indicated in the treatment of homocystinuria

CYSTAGON[®] (cysteamine bitartrate), indicated in the treatment of nephropathic cystinosis

ADAGEN[®] (pegademase bovine), indicated in the treatment of SCID-ADA deficiency

VEDROP[®] (water soluble vitamin E), indicated in the treatment of vitamin E deficiency in pediatric patients suffering from congenital chronic cholestasis

WILZIN[®] (zinc acetate), indicated in the treatment of Wilson's disease

A well balanced product pipeline

NAME	ORIGINATOR	INDICATION	DEVELOPMENT STATUS
VITAROS®	Apricus	Erectile dysfunction	Approved by a number of health authorities in Europe
CARBAGLU®	Recordati	Organic acidemias	Approved in EU Phase III in U.S.A.
CYSTADROPS®	Recordati	Ocular cystinosis	Filed in EU
FORTACIN™	Plethora Solutions	Premature ejaculation	Variation of EU approval
methadone		Treatment of cancer-related pain in cases of resistance or intolerance to opioids	Phase III b
CITRAFLEET®	Recordati/Casen	Cleansing of the colon in preparation for colonoscopy.	Split dose administration MA variation filed
GRASPA®	Erytech	Acute lymphoblastic leukemia (ALL) in patients with first recurrence of Philadelphia chromosome negative ALL Acute myeloid leukemia (AML) in patients >65 unfit for chemotherapy	Pre-filing in EU Phase II b
REC 0438	Recordati/UFPeptides	Overactive bladder (OAB) in patients with spinal lesions	Phase I in EU

A well balanced R&D pipeline (cont'd)

- **VITAROS®** (alprostadil) is a topically applied cream formulation of alprostadil for the treatment of erectile dysfunction. A patient-friendly form of alprostadil as an alternative to PDE-5 inhibitors for difficult to treat patients.
- **CARBAGLU®** (carglumic acid), currently approved for the treatment of hyperammonaemia due to NAGS deficiency, approved in Europe and in phase III clinical development in the USA for additional indications in organic acidemias (orphan drug designation granted).
- **CYSTADROPS®** (cysteamine chlorhydrate) are eye drops developed for “ocular manifestations of cystinosis” which cannot be controlled by orally administered cysteamine, specially formulated in a patient-friendly gel form.
- **FORTACIN™** (lidocaine+prilocaine) is an easy-to-use fast acting topical spray formulation of lidocaine and prilocaine for the treatment of premature ejaculation.
- **Methadone**, currently used in France, where it is distributed by Bouchara Recordati, as replacement therapy for major opioid drugs dependence. In 2012 Recordati started, in France, an open, multicenter, randomized, national Phase III b clinical study on methadone for the treatment of cancer-related pain inadequately relieved by opioids.

A well balanced R&D pipeline (cont'd)

- **CITRAFLEET®** is an intestinal evacuant used in preparation for colonoscopy. The objective of the trial is to assess the clinical effectiveness of a split-dose administration schedule compared to that of the current SmPC regimen.
- **GRASPA®** is L-asparaginase encapsulated in homologous human red blood cells. L-asparaginase has been shown to possess a powerful antitumor activity, but this enzyme is highly toxic and a large part of the patient population presents with a hypersensitivity and does not tolerate well the current treatment protocols. This population represents a large currently unmet medical need. Graspas® avoids toxicity and hypersensitivity issues associated with L-asparaginase treatments while maintaining its antitumor activity.
- **REC 0438** represents a structurally different class of compounds and is being studied for the treatment of OAB in patients with spinal lesions

First quarter 2015 highlights

- Revenue € 275.7 million, up 5.9%
- EBITDA € 82.8 million or 30.0% of sales, up 15.9%
- Operating income (EBIT) € 73.5 million or 26.6% of sales, up 18.1%
- Net income € 52.0 million or 18.8% of sales, up 21.5%
- Net debt € 136.8 million, a reduction of € 49.3 million

Main product sales

Corporate products including drugs for rare diseases account for 58.4% of revenue

(million Euro)	1Q 2015	1Q 2014	Change %
Zanidip [®] (lercanidipine)	33.8	31.3	8.1
Zanipress [®] (lercanidipine+enalapril)	16.9	15.7	7.3
Urorec [®] (silodosin)	16.0	14.5	10.8
Livazo [®] (pitavastatin)	5.9	6.4	(8.0)
Other corporate products*	51.0	47.6	7.1
Drugs for rare diseases	37.4	27.8	34.6

* Include the OTC corporate products for an amount of € 14.5 million in 2015 and € 11.5 million in 2014.

Composition of revenue by geography

Solid volume growth

(million Euro)	1Q 2015	1Q 2014	Change %
Italy	60.3	65.1	(7.4)
France	28.7	27.8	3.1
Germany	22.2	19.6	13.1
Turkey	21.1	16.4	28.5
U.S.A.	20.7	11.4	81.0
Russia, other CIS countries and Ukraine	18.0	19.6	(8.3)
Spain	16.9	17.5	(3.3)
North Africa	10.8	10.2	5.6
Portugal	9.3	9.0	3.4
Other CEE countries	7.8	6.5	20.9
Other W. Europe countries	6.3	5.7	10.0
Other international sales	45.1	42.8	5.3
TOTAL PHARMACEUTICALS	267.0	251.6	6.1
PHARMACEUTICAL CHEMICALS	8.7	8.8	(0.5)

(In local currency, millions)	1Q 2015	1Q 2014	Change %
Russia (RUB)	1,104.7	752.4	46.8
Turkish subsidiary (TRY)	55.3	46.8	18.3
U.S.A. (USD)	23.3	15.6	48.8

First quarter 2015 results

Further margin growth

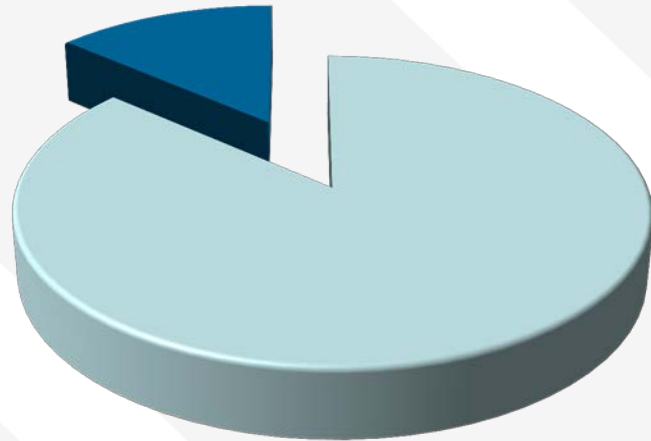
(million Euro)	1Q 2015	1Q 2014	Change %
Revenue	275.7	260.4	5.9
Gross Profit	187.5	173.4	8.1
as % of revenue	68.0	66.6	
SG&A Expenses	93.3	90.0	3.7
as % of revenue	33.9	34.6	
R&D Expenses	19.9	20.8	(4.3)
as % of revenue	7.2	8.0	
Other Income (Expense), net	(0.8)	(0.4)	79.9
as % of revenue	(0.3)	(0.2)	
Operating Income	73.5	62.2	18.1
as % of revenue	26.6	23.9	
Net Income	52.0	42.8	21.5
as % of revenue	18.8	16.4	
EBITDA	82.8	71.4	15.9
as % of revenue	30.0	27.4	

First quarter 2015 results

Operating segments

Revenue

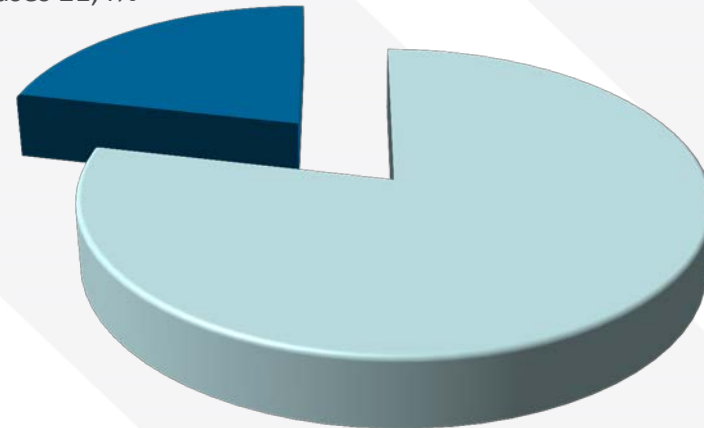
Treatments for rare diseases 13.6%



Primary & specialty care 86.4%

EBIT

Treatments for rare diseases 21,4%



Primary & specialty care 78,6%

EBIT margin on sales:

Treatments for rare diseases, 42.0%

Primary & Specialty care, 24.2%

Financial position and Shareholders' equity

(million Euro)	31 Mar 2015	31 Dec 2014	Change
Cash and short-term financial investments	187.1	137.0	50.1
Bank overdrafts and short-term loans	(8.0)	(8.6)	0.6
Loans – due within one year	(28.3)	(28.3)	0.0
Loans – due after one year	(287.6)	(286.2)	(1.4)
NET FINANCIAL POSITION	(136.8)	(186.0)	49.2
SHAREHOLDERS' EQUITY	865.0	787.4	77.6

2015 targets

Assumptions

- Lercanidipine franchise to remain relatively stable generating sales of between € 160 and € 165 million.
- Sales of Urorec® to increase by around 15%. Sales of Livazo® to increase by more than 10%.
- First half sales in Italy to be impacted by termination of license agreement for Entact® (escitalopram) as from mid-year 2014.
- Sales in Russia to increase by more than 10% in local currency. Sales in Turkey to increase by around 10% in local currency.
- Drugs for the treatment of rare diseases to grow double digit.
- Main FX impact on net revenue due to devaluation of the Russian rouble and revaluation of the US dollar. Net impact is a negative effect of around € 15 million.
- Breakdown of sales by product/business and by geography to remain substantially unchanged.

2017 plan

Assumptions

- Lercanidipine franchise to decline slightly following entry of generic competition for Zanipress[®]. Sales going forward of around € 140 million.
- Sales of Urorec[®] to reach around € 85 million in 2017.
- Sales of Livazo[®] to exceed € 35 million in 2017.
- Other corporate products to exceed € 140 million in 2017 including around € 20 million of new products (Vitaros[®], Fortacin[™]).
- Drugs for the treatment of rare diseases to exceed 15% of total sales (CAGR > 10%).
- OTC products to remain essentially stable as a percent of total sales.
- Objectives for 2017 are based on current currency exchange rates.
- Cash generated to be invested in corporate development activities, primarily in product portfolio enhancement.

2015 targets and 2017 plan

Objectives

- **Revenue:**
 - Target for 2015 to achieve sales of more than € 1 billion, below original expectations due exclusively to negative FX effect due to devaluation of the rouble.
 - Plan for 2017 to achieve sales of around € 1,150 million. Organic growth about one half of 2015-2017 increase in revenues.
- **Further margin improvement:**
 - Exceed original expectations for 2015: EBIT around € 250 million, 25% of sales, Net Income around € 175 million, >17% of sales.
 - Plan for 2017 to achieve EBIT between 25% and 26% of sales, Net Income between 17% and 18% of sales.

Financial projections

2015 targets and 2017 plan

(million Euro)	2014 Actual	2015 Targets		2017 Plan
Revenue	987.4	> 1,000		± 1,150
Operating income (EBIT)	231.0	± 250	EBIT margin	25% - 26%
Net Income	161.2	± 175	Net income margin	17% - 18%

The Recordati share

The Recordati share (ticker REC, **Reuters RECI.MI, Bloomberg REC IM, ISIN IT 0003828271**) has been listed on the Italian Stock Exchange since 1984. It belongs to the FTSE IT Mid Cap and FTSE IT Health Care indexes.

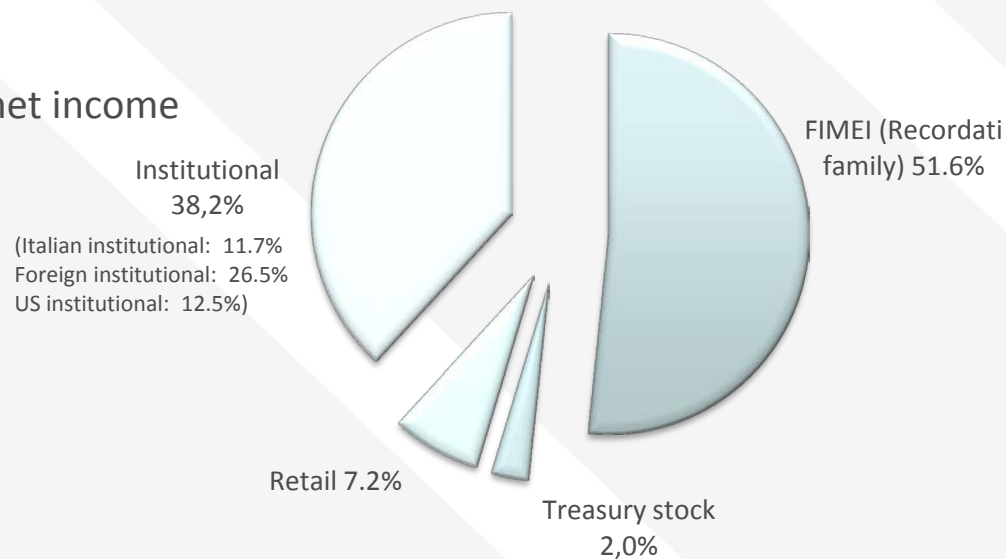
Share capital consists of **209,125,156** ordinary (common) shares with a par value of € 0.125 each.

2014 EPS (diluted): € 0.771

2014 dividend per share: € 0.50

Dividend pay-out ratio: $\pm 60\%$ of group net income

Ownership:



Company declarations, disclaimers and profile

DECLARATION BY THE MANAGER RESPONSIBLE FOR PREPARING THE COMPANY'S FINANCIAL REPORTS

The manager responsible for preparing the company's financial reports Fritz Squindo declares, pursuant to paragraph 2 of Article 154-bis of the Consolidated Law on Finance, that the accounting information contained in this presentation corresponds to the document results, books and accounting records.

Statements contained in this presentation, other than historical facts, are "forward-looking statements" (as such term is defined in the Private Securities Litigation Reform Act of 1995). These statements are based on currently available information, on current best estimates, and on assumptions believed to be reasonable. This information, these estimates and assumptions may prove to be incomplete or erroneous, and involve numerous risks and uncertainties, beyond the Company's control. Hence, actual results may differ materially from those expressed or implied by such forward-looking statements.

All mentions and descriptions of Recordati products are intended solely as information on the general nature of the company's activities and are not intended to indicate the advisability of administering any product in any particular instance.

Recordati, established in 1926, is an international pharmaceutical group, listed on the Italian Stock Exchange (Reuters RECI.MI, Bloomberg REC IM, ISIN IT 0003828271) with a total staff of around 4,000, dedicated to the research, development, manufacturing and marketing of pharmaceuticals. Headquartered in Milan, Italy, Recordati has operations in the main European countries, in Russia, in other Central and Eastern European countries, in Turkey, in the United States of America and in North Africa. An efficient field force of medical representatives promotes a wide range of innovative pharmaceuticals, both proprietary and under license, in a number of therapeutic areas including a specialized business dedicated to treatments for rare diseases. Recordati is a partner of choice for new product licenses for its territories. Recordati is committed to the research of new drug entities within the cardiovascular and urogenital therapeutic areas and of treatments for rare diseases. Consolidated revenue for 2014 is € 987.4 million, operating income is € 231.0 million and net income is € 161.2 million.

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