Condensed Consolidated Statement of Comprehensive Income

For the **nine months** ended 30 September	2011 $m	2010 $m
Revenue	24,935	24,652
Cost of sales	(4,414)	(4,630)
Gross profit	20,521	20,022
Distribution costs	(261)	(248)
Research and development	(3,656)	(3,388)
Selling, general and administrative costs	(8,020)	(7,923)
Other operating income and expense	2,044	620
Operating profit	10,628	9,083
Finance income	426	376
Finance expense	(739)	(765)
Profit before tax	10,315	8,694
Taxation	(1,792)	(2,245)
Profit for the period	8,523	6,449
Other comprehensive income:		
Foreign exchange arising on consolidation	21	13
Foreign exchange differences on borrowings forming net investment hedges	(25)	63
Amortisation of loss on cash flow hedge	2	1
Net available for sale losses taken to equity	(5)	-
Actuarial loss for the period	(53)	(384)
Income tax relating to components of other comprehensive income	4	84
Other comprehensive income for the period, net of tax	(56)	(223)
Total comprehensive income for the period	8,467	6,226
Profit attributable to:		
Owners of the parent	8,497	6,432
Non-controlling interests	26	17
	8,523	6,449
Total comprehensive income attributable to:		
Owners of the parent	8,429	6,193
Non-controlling interests	38	33
	8,467	6,226
Basic earnings per $0.25 Ordinary Share	$6.17	$4.45
Diluted earnings per $0.25 Ordinary Share	$6.14	$4.43
Weighted average number of Ordinary Shares in issue (millions)	1,377	1,445
Diluted weighted average number of Ordinary Shares in issue (millions)	1,383	1,452

Condensed Consolidated Statement of Comprehensive Income

For the **quarter** ended 30 September	2011 $m	2010 $m
Revenue	8,213	7,898
Cost of sales	(1,593)	(1,524)
Gross profit	6,620	6,374
Distribution costs	(93)	(82)
Research and development	(1,296)	(1,077)
Selling, general and administrative costs	(2,644)	(3,011)
Other operating income and expense	1,675	202
Operating profit	4,262	2,406
Finance income	153	123
Finance expense	(246)	(271)
Profit before tax	4,169	2,258
Taxation	(684)	(704)
Profit for the period	3,485	1,554
Other comprehensive income:		
Foreign exchange arising on consolidation	(225)	391
Foreign exchange differences on borrowings forming net investment hedges	88	(133)
Amortisation of loss on cash flow hedge	1	-
Net available for sale (losses)/gains taken to equity	(23)	5
Actuarial loss for the period	(209)	(56)
Income tax relating to components of other comprehensive income	10	67
Other comprehensive income for the period, net of tax	(358)	274
Total comprehensive income for the period	3,127	1,828
Profit attributable to:		
Owners of the parent	3,477	1,548
Non-controlling interests	8	6
	3,485	1,554
Total comprehensive income attributable to:		
Owners of the parent	3,111	1,812
Non-controlling interests	16	16
	3,127	1,828
Basic earnings per $0.25 Ordinary Share	$2.56	$1.08
Diluted earnings per $0.25 Ordinary Share	$2.54	$1.07
Weighted average number of Ordinary Shares in issue (millions)	1,354	1,437
Diluted weighted average number of Ordinary Shares in issue (millions)	1,359	1,446

Condensed Consolidated Statement of Financial Position

	At 30 Sep 2011 $m	At 31 Dec 2010 $m	At 30 Sep 2010 $m
ASSETS			
Non-current assets			
Property, plant and equipment	6,526	6,957	7,096
Goodwill	9,874	9,871	9,878
Intangible assets	11,661	12,158	12,945
Derivative financial instruments	355	324	420
Other investments	207	211	205
Deferred tax assets	1,486	1,475	1,277
	30,109	30,996	31,821
Current assets			
Inventories	1,955	1,682	1,810
Trade and other receivables	8,308	7,847	7,735
Other investments	924	1,482	1,517
Derivative financial instruments	29	9	49
Income tax receivable	1,391	3,043	3,448
Cash and cash equivalents	9,860	11,068	10,010
	22,467	25,131	24,569
Total assets	52,576	56,127	56,390
LIABILITIES			
Current liabilities			
Interest-bearing loans and borrowings	(2,055)	(125)	(1,376)
Trade and other payables	(8,028)	(8,661)	(7,796)
Derivative financial instruments	-	(8)	(82)
Provisions	(1,083)	(1,095)	(884)
Income tax payable	(3,491)	(6,898)	(6,714)
	(14,657)	(16,787)	(16,852)
Non-current liabilities			
Interest-bearing loans and borrowings	(7,394)	(9,097)	(9,231)
Deferred tax liabilities	(2,923)	(3,145)	(3,158)
Retirement benefit obligations	(2,388)	(2,472)	(3,739)
Provisions	(555)	(843)	(799)
Other payables	(505)	(373)	(299)
	(13,765)	(15,930)	(17,226)
Total liabilities	(28,422)	(32,717)	(34,078)
Net assets	24,154	23,410	22,312
EQUITY			
Capital and reserves attributable to equity holders of the Company			
Share capital	332	352	356
Share premium account	3,048	2,672	2,623
Other reserves	1,937	1,917	1,913
Retained earnings	18,614	18,272	17,233
	23,931	23,213	22,125
Non-controlling interests	223	197	187
Total equity	24,154	23,410	22,312

Condensed Consolidated Statement of Cash Flows

For the **nine months** ended 30 September	2011 $m	Restated 2010 $m
Cash flows from operating activities		
Profit before taxation	10,315	8,694
Finance income and expense	313	389
Depreciation, amortisation and impairment	1,580	1,434
Increase in working capital and short-term provisions	(1,528)	(1,016)
Other non-cash movements[1]	(1,806)	249
Cash generated from operations	8,874	9,750
Interest paid	(467)	(515)
Tax paid	(3,655)	(2,115)
Net cash inflow from operating activities	4,752	7,120
Cash flows from investing activities		
Movement in short-term investments and fixed deposits[2]	542	(80)
Purchase of property, plant and equipment	(593)	(473)
Disposal of property, plant and equipment	56	67
Purchase of intangible assets	(326)	(1,241)
Disposal of intangible assets	-	210
Purchase of non-current asset investments	(8)	(27)
Disposal of non-current asset investments	-	2
Acquisitions of business operations	-	(348)
Net cash received on disposal of subsidiaries	1,772	-
Interest received	131	126
Payments made by subsidiaries to non-controlling interests	(16)	(10)
Net cash inflow/(outflow) from investing activities	1,558	(1,774)
Net cash inflow before financing activities	6,310	5,346
Cash flows from financing activities		
Proceeds from issue of share capital	378	445
Repurchase of shares for cancellation	(4,256)	(1,742)
Repayment of loans	-	(717)
Dividends paid	(3,764)	(3,361)
Movement in derivative financial instruments[2]	3	(114)
Movement in short-term borrowings	(1)	(25)
Net cash outflow from financing activities	(7,640)	(5,514)
Net decrease in cash and cash equivalents in the period	(1,330)	(168)
Cash and cash equivalents at the beginning of the period	10,981	9,828
Exchange rate effects	(30)	16
Cash and cash equivalents at the end of the period	9,621	9,676
Cash and cash equivalents consists of:		
Cash and cash equivalents	9,860	10,010
Overdrafts	(239)	(334)
	9,621	9,676

[1] Included in other non-cash movements is the profit on disposal of Astra Tech (see Note 4).

[2] 2010 restated to reclassify $114m movement in derivative financial instruments associated with financing activities.

Condensed Consolidated Statement of Changes in Equity

	Share capital $m	Share premium account $m	Other* reserves $m	Retained earnings $m	Total $m	Non-controlling interests $m	Total equity $m
At 1 January 2010	363	2,180	1,919	16,198	20,660	161	20,821
Profit for the period	-	-	-	6,432	6,432	17	6,449
Other comprehensive income	-	-	-	(239)	(239)	16	(223)
Transfer to other reserve	-	-	(15)	15	-	-	-
Transactions with owners:							
Dividends	-	-	-	(3,494)	(3,494)	-	(3,494)
Issue of AstraZeneca PLC Ordinary shares	2	443	-	-	445	-	445
Repurchase of AstraZeneca PLC Ordinary shares	(9)	-	9	(1,742)	(1,742)	-	(1,742)
Share-based payments	-	-	-	63	63	-	63
Transfer from non-controlling interests to payables	-	-	-	-	-	(6)	(6)
Dividend paid to non-controlling interest	-	-	-	-	-	(1)	(1)
Net movement	(7)	443	(6)	1,035	1,465	26	1,491
At 30 September 2010	**356**	**2,623**	**1,913**	**17,233**	**22,125**	**187**	**22,312**

	Share capital $m	Share premium account $m	Other* reserves $m	Retained earnings $m	Total $m	Non-controlling interests $m	Total equity $m
At 1 January 2011	352	2,672	1,917	18,272	23,213	197	23,410
Profit for the period	-	-	-	8,497	8,497	26	8,523
Other comprehensive income	-	-	-	(68)	(68)	12	(56)
Transfer to other reserve	-	-	(2)	2	-	-	-
Transactions with owners:							
Dividends	-	-	-	(3,752)	(3,752)	-	(3,752)
Issue of AstraZeneca PLC Ordinary shares	2	376	-	-	378	-	378
Repurchase of AstraZeneca PLC Ordinary shares	(22)	-	22	(4,256)	(4,256)	-	(4,256)
Share-based payments	-	-	-	(81)	(81)	-	(81)
Transfer from non-controlling interests to payables	-	-	-	-	-	(8)	(8)
Dividend paid to non-controlling interests	-	-	-	-	-	(4)	(4)
Net movement	(20)	376	20	342	718	26	744
At 30 September 2011	**332**	**3,048**	**1,937**	**18,614**	**23,931**	**223**	**24,154**

* Other reserves includes the capital redemption reserve and the merger reserve.

1 BASIS OF PREPARATION AND ACCOUNTING POLICIES

These condensed consolidated interim financial statements ("interim financial statements") for the nine months ended 30 September 2011 have been prepared in accordance with IAS 34 *Interim Financial Reporting* as adopted by the European Union. The annual financial statements of the Group are prepared in accordance with International Financial Reporting Standards (IFRSs) as adopted by the European Union and as issued by the International Accounting Standards Board. As required by the Disclosure and Transparency Rules of the Financial Services Authority, the interim financial statements have been prepared applying the accounting policies and presentation that were applied in the preparation of the Company's published consolidated financial statements for the year ended 31 December 2010, except where new or revised accounting standards have been applied. There has been no significant impact on the Group profit or net assets on adoption of new or revised accounting standards in the period.

The Group has considerable financial resources available. The Group's revenues are largely derived from sales of products which are covered by patents and for which, historically at least, demand has been relatively unaffected by changes in the general economy. As a consequence, the Directors believe that the Group is well placed to manage its business risks successfully and as such, the interim financial statements have been prepared on a Going Concern basis.

The information contained in Note 5 updates the disclosures concerning legal proceedings and contingent liabilities in the Group's Annual Report and Form 20-F Information 2010.

The comparative figures for the financial year ended 31 December 2010 are not the Company's statutory accounts for that financial year. Those accounts have been reported on by the Company's auditors and delivered to the registrar of companies. The report of the auditors was (i) unqualified, (ii) did not include a reference to any matters to which the auditors drew attention by way of emphasis without qualifying their report, and (iii) did not contain a statement under section 498(2) or (3) of the Companies Act 2006.

2 NET FUNDS

The table below provides an analysis of net funds and a reconciliation of net cash flow to the movement in net funds.

	At 1 Jan 2011 $m	Cash flow $m	Non-cash mvmts $m	Exchange mvmts $m	At 30 Sep 2011 $m
Loans due after one year	(9,097)	-	1,729	(26)	(7,394)
Current instalments of loan	-	-	(1,777)	-	(1,777)
Total loans	(9,097)	-	(48)	(26)	(9,171)
Other investments - current	1,482	(542)	(16)	-	924
Net derivative financial instruments	325	(3)	62	-	384
Cash and cash equivalents	11,068	(1,179)	-	(29)	9,860
Overdrafts	(87)	(151)	-	(1)	(239)
Short-term borrowings	(38)	1	-	(2)	(39)
	12,750	(1,874)	46	(32)	10,890
Net funds	3,653	(1,874)	(2)	(58)	1,719

Non-cash movements in the period include fair value adjustments under IAS 39.

3 RESTRUCTURING COSTS

Profit before tax for the nine months ended 30 September 2011 is stated after charging restructuring costs of $502 million ($777 million in the first nine months of 2010). These have been charged to profit as follows:

	3rd Quarter 2011 $m	3rd Quarter 2010 $m	9 Months 2011 $m	9 Months 2010 $m
Cost of sales	(14)	19	18	110
Research and development	124	91	293	463
Selling, general and administrative costs	111	102	191	204
Total	**221**	**212**	**502**	**777**

4 DISPOSAL OF ASTRA TECH

In August 2011, the Group announced the sale of the Astra Tech business to Dentsply International for approximately $1.8 billion in cash. At 30 September 2011, the Group has reported a profit on disposal of $1,483 million and a total cash inflow of $1,772 million as a result of this transaction.

	$m
Consideration	1,795
Net assets	(279)
Fees and other disposal costs	(59)
Exchange recycled on disposal	26
Profit on disposal	**1,483**

	$m
Consideration	1,795
Cash held in Astra Tech on disposal	(23)
Cash inflow on disposal	**1,772**

5 LEGAL PROCEEDINGS AND CONTINGENT LIABILITIES

AstraZeneca is involved in various legal proceedings considered typical to its business, including litigation and investigations relating to product liability, commercial disputes, infringement of intellectual property rights, the validity of certain patents, anti-trust law and sales and marketing practices. The matters discussed below constitute the more significant developments since publication of the disclosures concerning legal proceedings in the Company's Annual Report and Form 20-F Information 2010 and Interim Management Statement 2011 as part of the Company's Half-Yearly Financial Report for the six-month period to 30 June 2011. Unless noted otherwise below or in the Annual Report and Form 20-F Information 2010 and Interim Management Statement 2011 as part of the Company's Half-Yearly Financial Report for the six-month period to 30 June 2011, no provisions have been established in respect of the claims discussed below.

As discussed in the Company's Annual Report and Form 20-F Information 2010, for the majority of claims in which AstraZeneca is involved it is not possible to make a reasonable estimate of the expected financial effect, if any, that will result from ultimate resolution of the proceedings. In these cases, AstraZeneca discloses information with respect only to the nature and facts of the cases but no provision is made.

In cases that have been settled or adjudicated, or where quantifiable fines and penalties have been assessed and which are not subject to appeal, or where a loss is probable and we are able to make a reasonable estimate of the loss, we record the loss absorbed or make a provision for our best estimate of the expected loss.

The position could change over time and the estimates that we have made and upon which we have relied in calculating these provisions are inherently imprecise. There can, therefore, be no assurance that any losses that result from the outcome of any legal proceedings will not exceed the amount of the provisions that have been booked in the accounts. The major factors causing this uncertainty are described more fully in the Annual Report and Form 20-F Information 2010 and herein.

AstraZeneca has full confidence in, and will vigorously defend and enforce its intellectual property.

Matters disclosed in respect of the third quarter of 2011 and October 2011

***Arimidex* (anastrozole)**
Patent Proceedings pursuant to Patented Medicines (Notice of Compliance) Regulations—Canada (NOC Proceedings)
As previously disclosed, the Canadian Federal Court conducted a hearing in the NOC Proceeding filed by AstraZeneca against Mylan Pharmaceuticals ULC (Mylan) in respect of AstraZeneca's Canadian substance Patent No. 1,337,420 (the '420 patent). In August 2011, the Court granted a Prohibition Order preventing the Minister of Health from granting a marketing authorisation to Mylan until the expiry of the '420 patent on 24 October 2012. Mylan has appealed.

As previously disclosed, in May 2011, AstraZeneca commenced a NOC Proceeding against Pharmascience Inc. in respect of the '420 patent. In August 2011, the proceeding was stayed until a final decision, including on appeal, is reached in the above-noted Mylan NOC Proceeding.

As previously disclosed, in May 2011, AstraZeneca commenced a NOC Proceeding against Teva Canada Limited in respect of the '420 patent. In September 2011, the proceeding was stayed until the expiry of the '420 patent.

As previously disclosed, in July 2011, AstraZeneca commenced a NOC Proceeding against Apotex Inc. (Apotex) in respect of the '420 patent. In September 2011, the proceeding was discontinued after Apotex withdrew its Notice of Allegation.

***Atacand* (candesartan cilexetil)**

Patent litigation – EU

As previously disclosed, in Portugal, in addition to the previously disclosed cases, approvals for generic candesartan cilexetil or candesartan cilexetil and hydrochlorothiazide have been granted to Pharmakern Portugal - Produtos Farmacêuticos, Sociedade Unipessoal, Lda. (Pharmakern); Sandoz Farmacêutica Lda. (Sandoz); Bluepharma Genéricos Comércio de Medicamentos S.A. and Bluepharma Indústria Pharmacêutica S.A (together Bluepharma); and Sanofi-Aventis - Produtos Farmacêuticos, Lda (Sanofi). AstraZeneca filed preliminary injunctions to suspend those marketing approvals as well as corresponding administrative main actions during the third quarter of 2011.

In the cases against Laboratórios Azevados – Industria Farmacêutica; Ceamed Servico e Consultadoria Farmacêutica Lda; Teva Pharma – Produtos Farmacêuticos Lda; Mylan Lda; Laboratórios Anova - Produtos Farmacêuticos, Lda; Ranbaxy Portugal - Comércio e Desenvolvimento de Produtos Farmacêuticos, Unipessoal Lda; and Ratiopharm - Comércio e Indústria de Produtos Farmacêuticos the Court of First Instance issued negative decisions, denying the preliminary injunction requests. Appeals have been filed and we await decisions from the Court of Appeal.

***Atacand Plus* (candesartan cilexetil / hydrochlorothiazide)**

Patent Proceedings pursuant to Patented Medicines (Notice of Compliance) Regulations — Canada (NOC Proceedings)

In August 2011, AstraZeneca settled the previously disclosed NOC Proceeding pending with Pharmascience Inc. (PMS) with respect to Canadian Patent Nos. 2,040,955 (the '955 patent), 2,083,305 (the '305 patent) and 2,125,251 (the '251 patent). The settlement resolves the litigation and allows PMS to enter the Canadian market on 23 September 2012, or earlier, in certain circumstances.

In August 2011, AstraZeneca settled the previously disclosed NOC Proceeding pending with Teva Canada Limited (Teva) with respect to the '955 patent, the '305 patent and the '251 patent. The settlement resolves the litigation and allows Teva to enter the Canadian market on 23 September 2012, or earlier, in certain circumstances.

***Crestor* (rosuvastatin calcium)**

Patent litigation – US

Abbreviated New Drug Applications (ANDAs)—US Patent No. RE37.314 (the '314 patent)

On 5 October 2011, the US Court of Appeals for the Federal Circuit held oral argument on the appeal of the June 2010 decision by the US District Court for the District of Delaware, which found the '314 patent valid, enforceable and infringed by the eight generic defendants.

Abbreviated New Drug Applications (ANDAs)—US Patent Nos. 6,858,618 (the '618 patent) and 7,030,152 (the '152 patent)

The US Court of Appeals for the Federal Circuit scheduled oral argument for 7 November 2011, on the appeal by AstraZeneca and The Brighams & Women's Hospital (BWH) (AstraZeneca's licensor of the '152 patent) (together the Plaintiffs) of the dismissal by the US District Court for the District of Delaware of the *Crestor* ANDA patent infringement actions based on the '152 and the '618 patents for lack of subject matter jurisdiction.

Teva Pharmaceutical Industries LTD. (Teva LTD) Infringement suit in the Eastern District of Pennsylvania

On 6 October 2011, the US Court of Appeals for the Federal Circuit held oral argument on the appeal of the decision by the US District Court for the District of Pennsylvania granting AstraZeneca's motion for summary judgment and invalidating Teva LTD's formulation patent.

Patent Proceedings pursuant to Patented Medicines (Notice of Compliance) Regulations — Canada (NOC Proceedings)

As previously disclosed, in July 2010, AstraZeneca received a Notice of Allegation from Ranbaxy Pharmaceuticals Canada Inc. (Ranbaxy) regarding Canadian Patent Nos. 2,072,945 (the '945 patent), 2,313,783 (the '783 patent) and 2,315,141 (the '141 patent). In July 2011, AstraZeneca reached a comprehensive settlement agreement resolving the litigation and as part of the agreement, Ranbaxy may enter the Canadian market in April 2012, or earlier, in certain circumstances.

As previously disclosed, in July 2011, AstraZeneca received a Notice of Allegation (NOA) from Laboratoire Riva Inc. (Riva) under the Canadian Patented Medicines (Notice of Compliance) Regulations in respect of the '945 patent, the '783 patent and the '141 patent. AstraZeneca commenced a proceeding in response in August 2011.

Patent litigation/Data exclusivity – Brazil

As previously disclosed, the court denied AstraZeneca's request for data exclusivity for *Crestor*. AstraZeneca requested an interlocutory appeal of the decision, which was denied. AstraZeneca filed a motion for reconsideration in September 2011.

***Nexium* (esomeprazole magnesium)**

Patent litigation – US

Abbreviated New Drug Applications (ANDAs)

As previously disclosed, in June 2011, AstraZeneca received a Paragraph IV Certification notice-letter from Hetero Drug Limited Unit III (Hetero) stating that Hetero had submitted an ANDA for approval to market 20 and 40mg esomeprazole magnesium delayed-release capsules. In August 2011, AstraZeneca filed an ANDA patent infringement action against Hetero in the US District Court for the District of New Jersey. In September 2011, the proceeding was stayed.

Patent Litigation – EU: 10-year countries
As previously disclosed, in the UK, in October 2010, AstraZeneca was served an invalidity case in which Ranbaxy (UK) Ltd (Ranbaxy) claimed that the *Nexium* esomeprazole magnesium patent (the EP 1020461 patent) and the esomeprazole magnesium trihydrate patent (the EP 0984957 patent) were invalid in the UK. Ranbaxy further requested the court to find that its generic esomeprazole product would not infringe either patent if launched in the UK. In March 2011, AstraZeneca filed a suit against Ranbaxy claiming that its generic esomeprazole product infringes the EP 1020461 patent. The trial of the non-infringement/infringement part took place in June 2011. In July 2011, the court held that Ranbaxy's generic esomeprazole product does not infringe the EP 1020461 patent. AstraZeneca has not appealed. The invalidity part of the proceedings had been stayed pending the non-infringement trial. There has been no update as to when the invalidity part will be heard.

In September 2011, in the Netherlands, AstraZeneca was served a declaratory action in which Ranbaxy (UK) Ltd requests the court to find that its generic esomeprazole product would not infringe the EP 1020461 patent or the EP 0984957 patent if commercialised in the Netherlands. In September 2011, AstraZeneca was served a nullity action in which Actavis Group PTC ehf claims that two esomeprazole formulation patents (EP 984773 and EP 1124539) are invalid. AstraZeneca is preparing its responses.

Patent Litigation – EU: 6-year countries
As previously disclosed, in Finland, AstraZeneca initiated a declaratory action requesting the District Court of Helsinki to find that Ranbaxy (UK) Ltd would infringe a patent relating to esomeprazole if commercialising its generic esomeprazole magnesium products. The trial took place in May 2011. In June 2011, the Court denied AstraZeneca's claim. AstraZeneca has appealed. In July 2009, AstraZeneca initiated similar declaratory actions against Sandoz Oy AB and Sandoz A/S. In September 2009, Hexal AG, Sandoz Oy AB and Sandoz A/S (all in the Sandoz group) initiated an invalidity case requesting the court to invalidate the same patent. These cases were heard together in September 2011. On 18 July 2011, AstraZeneca applied for an interlocutory injunction to be granted against Sandoz A/S and Novartis Finland Oy and the interlocutory injunction was granted on 26 July 2011.

On 13 July 2011, in Poland, AstraZeneca filed an application for an interlocutory injunction to be granted against Krka d.d.Novo Mesto (Krka). On 4 August 2011, the Regional Court of Lodz granted the interlocutory injunction. Krka can appeal. On 8 August 2011, the Regional Court of Warsaw dismissed the declaratory action of Lek Farmacevtska Druzba d.d. (Lek) and Sandoz GmbH (Sandoz) (both in the Sandoz group). Lek and Sandoz can appeal.

As previously disclosed, in Ireland, AstraZeneca reached a settlement with Krka, d.d. Novo Mesto and Pinewood Laboratories Ltd (together Krka), in October 2011, discontinuing the legal actions under which AstraZeneca claimed that Krka's generic esomeprazole magnesium product infringes EP 1020461 and under which Krka claimed that EP 1020461 is invalid in Ireland.

Patent litigation – Norway
As previously disclosed, in Norway, the Appeal Court held the esomeprazole magnesium patent as valid. Hexal AG, Sandoz AS and Sandoz A/S (all in the Sandoz group) applied for leave to appeal to the Supreme Court. On 29 August 2011, the Supreme Court denied the requested leave to appeal.

Patent litigation – Singapore
As previously disclosed, in July 2011, AstraZeneca initiated patent infringement proceedings against Ranbaxy (Malaysia) SDN BHD (Ranbaxy) based on an esomeprazole related patent. In August 2011, Ranbaxy initiated an invalidity case regarding the esomeprazole magnesium patent.

Patent litigation – Turkey
In July 2011, AstraZeneca initiated patent infringement proceedings against Logus Ilac, Integri Ilac, Vem Ilac, Biofarma Ilac and Sandoz Ilac San.ve Tic.AS based on esomeprazole related patents. In September 2011, the court rejected AstraZeneca's claim in the case against Integri Ilac. AstraZeneca cannot appeal.

Patent litigation – Australia
In September 2011, AstraZeneca was served an invalidity case in which Ranbaxy (UK) Ltd claimed that the esomeprazole magnesium patent and a formulation patent relating to esomeprazole are invalid in Australia.

Patent Proceedings – EU
As previously disclosed, the Opposition Division of the European Patent Office (EPO) decided to revoke two patents that relate to *Nexium* (the EP 1020461 patent) and *Nexium i.v.* (the EP 1020460 patent) in June and July 2011 following Notices of Opposition filed by parties opposed to the grant of these patents. AstraZeneca has now appealed these decisions. Formal Notices of Appeal were filed for the EP 1020460 patent on 15 July 2011 and for the EP 1020461 patent on 1 August 2011.

Patent Proceedings - China
In September 2011, AstraZeneca received notice that an individual had filed a request with the Chinese Patent Office for invalidation of a Chinese substance patent relating to *Nexium* (Chinese Patent for Invention No. 94190335.4).

***Nexium i.v.* (esomeprazole sodium)**
Patent litigation – US
Abbreviated New Drug Applications (ANDAs)
In October 2011, AstraZeneca entered into an agreement with Sun Pharma Global FZE and affiliates (together Sun) to settle the previously disclosed patent infringement suit that AstraZeneca filed against Sun in the US District Court for the District of New Jersey with respect to Sun's ANDA for esomeprazole sodium intravenous formulation. As part of the settlement agreement, AstraZeneca has granted Sun a licence to enter the US market with its generic esomeprazole sodium formulation on 1 January 2014, subject to regulatory approval, or earlier in certain circumstances.

***Seroquel* (quetiapine fumarate)**

Product liability

As of October 2011, approximately 28,618 claims have been settled in principle, 28,450 of which are subject to written agreements and AstraZeneca was aware of approximately 75 *Seroquel* US product liability claims that have not been settled in principle.

As of 30 September 2011, legal defence costs of approximately $759m have been incurred in connection with *Seroquel*-related product liability claims. As previously disclosed, AstraZeneca settled its claims against several of its insurers for a substantial part of those legal defence costs.

As previously disclosed, disputes continue with other insurers about the availability of coverage under other insurance policies for legal defence costs and settlements. These policies have aggregate coverage limits of $300m. On 5 September 2011, AstraZeneca Insurance Company Limited commenced formal legal proceedings in the High Court in London against certain of these insurers for recovery of money which AstraZeneca believes is due under certain of these policies which have aggregate coverage limits of $200m.

As of 30 September 2011, out of the legal defence costs of $759m mentioned above, AstraZeneca believes that approximately $144m is covered by these insurance policies.

While no insurance receivable can be recognised under applicable accounting standards at this time, AstraZeneca believes that it is more likely than not that further insurance recoveries will be secured under the additional policies, but there can be no assurance of this or the amount of any potential future recovery.

Regulatory Related Matters – US

On 9 September 2011, AstraZeneca filed a Citizen Petition with the US Food and Drug Administration (FDA) for *Seroquel* asking the FDA not to approve any generic quetiapine drug product that omits certain hyperglycemia and suicidality warning language from its label that the FDA required AstraZeneca to include in the *Seroquel* labelling. The FDA is required to issue a decision on this petition by 7 March 2012.

Patent litigation – EU

In Portugal, in addition to the previously disclosed cases, approvals were granted to Germed Farmacêutica Lda (Germed) and ToLife Produtos Farmacêuticos S.A. in September 2011. Preliminary injunctions to suspend those marketing approvals as well as corresponding administrative main actions were filed in October 2011. In the cases against Alter S.A. and Mepha Investigacão, Desenvolvimento e Fabricacão Farmacêutica, Lda, the Court of First Instance issued negative decisions, denying the preliminary injunction requests. Appeals were filed early in October 2011 and AstraZeneca is waiting for the decision of the Court of Appeal.

In Italy, as previously disclosed, after a mistake by the Italian Patent Office, AstraZeneca filed motions for preliminary injunctions against five generic companies. In September 2011, the parties agreed to settle the cases.

Seroquel XR

Patent litigation – US

Abbreviated New Drug Applications (ANDAs)

As previously disclosed, in May 2011, AstraZeneca filed an ANDA patent infringement action in the US District Court for the District of New Jersey against Intellipharmaceutics Corp. (IPC) alleging infringement of US Patent No. 5,948,437 (the '437 patent). After IPC filed a motion seeking to have the case dismissed for lack of personal jurisdiction or alternatively, for the action to be transferred to New York, AstraZeneca filed a second, essentially identical lawsuit in the US District Court for the Southern District of New York. In August 2011, the US District Court for the Southern District of New York dismissed the second suit without prejudice. AstraZeneca filed a motion for reconsideration and reinstatement of the original filing date for the action, which was granted in September 2011.

As previously reported, AstraZeneca filed patent infringement actions in the US District Court for the District of New Jersey against various entities of Handa Pharmaceuticals, LLC (Handa), Accord Healthcare Inc. (Accord), Anchen Pharmaceuticals, Inc. (Anchen), Torrent Pharmaceuticals Ltd. (Torrent), Osmotica Pharmaceutical Corporation (Osmotica), and Mylan Pharmaceuticals Inc. (Mylan).

On 29 September 2011, AstraZeneca settled its patent infringement action against Handa by granting Handa a licence to the '437 patent effective 1 November 2016, or earlier under certain circumstances. On 4 October 2011, the Court dismissed the action against Handa.

Beginning 3 October 2011, the US District Court for the District of New Jersey conducted a trial of the pending patent infringement actions against Accord, Anchen, Torrent, Osmotica and Mylan. On 5 October 2011, AstraZeneca settled its patent infringement action against Accord by granting Accord a licence to the '437 patent effective 1 November 2016, or earlier under certain circumstances. On 7 October 2011, the Court dismissed the action against Accord.

Patent Proceedings pursuant to Patented Medicines (Notice of Compliance) regulations—Canada (NOC Proceedings)

In August 2011, AstraZeneca commenced a NOC Proceeding in response to the June 2011 Notice of Allegation from Teva Canada Limited under the Patented Medicines (Notice of Compliance) Regulations respecting Canadian Patent No. 2,251,944.

Patent litigation – EU

In Spain, Accord Healthcare S.L.U. and Sandoz Farmaceutica S.A. issued revocation proceedings against AstraZeneca in July 2011, which AstraZeneca received notification of in August 2011. A trial date has not yet been scheduled.

In Romania, Teva Pharmaceuticals S.R.L. issued revocation proceedings against AstraZeneca in September 2011, which AstraZeneca received notification of in October 2011.

Patent litigation – Turkey
In Turkey, Sanovel İlaç San. ve Tic.A.Ş, has filed a negative declaratory action for non-infringement. A trial date has not yet been scheduled.

Regulatory Related Matters – US
On 9 September 2011, AstraZeneca filed a Citizen Petition with the US Food and Drug Administration (FDA) for *Seroquel XR* that asks the FDA not to approve any generic quetiapine drug product that omits certain hyperglycemia and suicidality warning language from its label that the FDA required AstraZeneca to include in the *Seroquel XR* labelling. The FDA is required to issue a decision on this petition by 7 March 2012.

***Vimovo* (fixed-dose combination of naproxen and esomeprazole)**
Abbreviated New Drug Applications (ANDAs)
As previously disclosed, in June 2011, Dr. Reddy's Laboratories and Dr. Reddy's Laboratories, Ltd. (together DRL) answered the ANDA patent infringement suit filed by AstraZeneca and Pozen, Inc. (Pozen) (AstraZeneca's licensor) in the US District Court for the District of New Jersey against DRL alleging infringement of US Patent No 6,926,907 (the '907 patent). AstraZeneca received a Paragraph IV Certification notice-letter from DRL dated 19 September 2011, indicating it also seeks approval to market generic versions of 375/20mg and 500/20mg *Vimovo* tablets before expiration of US Patent Nos. 5,714,504 (the '504 patent); 6,875,872 (the '872 patent); 5,900,424 (the '424 patent); 6,369,085 (the '085 patent); 7,411,070 (the '070 patent); and 7,745,466 (the '466 patent). AstraZeneca is evaluating DRL's certifications.

On 26 September 2011, Lupin Ltd. and Lupin Pharmaceuticals, Inc. (together Lupin) answered the ANDA patent infringement complaint filed by AstraZeneca and Pozen in the US District Court for the District of New Jersey against Lupin alleging patent infringement of the '504, '085, '872, '907, '070, and '466 patents.

AstraZeneca received a Paragraph IV Certification notice-letter from Anchen Pharmaceuticals, Inc. (Anchen) dated 16 September 2011, indicating it seeks approval to market generic versions of 375/20mg and 500/20mg *Vimovo* tablets before expiration of the '085, '424, '907, '070, and '466 patents. AstraZeneca is evaluating Anchen's certifications.

Other Commercial Litigation
***Synagis* (palivizumab)**
In August 2011, AstraZeneca's biologics unit MedImmune filed a declaratory action against Abbott International LLC (Abbott) in the Federal District Court in Maryland. The action sought a declaratory judgment related to a contract dispute between the parties. The parties are disputing when the transfer price of *Synagis* would revert to a lower amount based on the occurrence of one or more events related to the development of motavizumab (the Reversion Event). On 15 September 2011, Abbott filed a motion to dismiss the declaratory action filed in the Federal District Court in Maryland. On the same date, Abbott also filed a parallel action in Illinois state court for breach of contract and for a declaratory judgment that the Reversion Event shall be deemed to have occurred on 1 July 2011 or alternatively, shall occur as of 31 December 2011.

On 26 September 2011, MedImmune voluntarily dismissed the complaint in the Federal District Court and re-filed the same declaratory action seeking the same relief in the state court in Montgomery County, Maryland. MedImmune expects to file a motion to dismiss the Illinois action as it believes the contract requires any litigation to be litigated in the Federal or state courts of Maryland. Abbott's response to the state court action in Maryland is not due until late November 2011.

***Toprol XL* (metoprolol succinate)**
As previously disclosed, AstraZeneca is defending anti-trust claims regarding *Toprol XL*, brought by both direct purchasers and end-payers. AstraZeneca has taken a provision of $21 million in connection with an agreement in principle to settle the claims of the putative class of direct purchasers. AstraZeneca continues to defend against the claims of those who have opted-out of the class and against the claims alleged by end-payers.

Other Pricing Litigation
Average Wholesale Price Litigation
As previously disclosed, AstraZeneca is a defendant, along with many other pharmaceutical manufacturers, in several sets of cases involving allegations that, by causing the publication of allegedly inflated wholesale list prices, the defendants caused entities to overpay for prescription drugs. In October 2011, AstraZeneca agreed in principle to settle the lawsuits brought by the Attorneys General of the States of Kansas and Mississippi, subject to documentation. Provision has been made in the fourth quarter in respect of these settlements.

Average Manufacturer's Price Qui Tam Litigation (Streck)
AstraZeneca is one of several manufacturers named as a defendant in a lawsuit filed in US federal court in Philadelphia by Ronald J. Streck, under the *qui tam* (whistleblower) provisions of the federal and certain state False Claims Acts. The action was initially filed in October 2008 but remained under seal until May 2011, following the government's decision not to intervene in the case with regard to certain manufacturers, including AstraZeneca. AstraZeneca was served with a copy of the amended complaint on 7 September 2011. The lawsuit seeks to recover, among other things, damages, civil penalties, and attorneys' fees for alleged inaccurate reporting of Average Manufacturer's Prices to the Centers for Medicaid and Medicare Services. AstraZeneca intends to vigorously defend against these claims.

Other Actual and Threatened Government Investigations and Related Litigation
On 5 October 2011, AstraZeneca LP and AstraZeneca Pharmaceuticals LP received a subpoena from the Department of Justice in connection with an investigation of the possible submission of false or otherwise improper pricing information to the Centers for Medicare and Medicaid Services. The precise parameters of this inquiry are unknown, and AstraZeneca is not in a position at this time to predict the scope, duration or outcome of this matter, including whether it will result in any liability to AstraZeneca.

AstraZeneca LP and AstraZeneca Pharmaceuticals LP are also in receipt of a Civil Investigative Demand issued by the Attorney General of Texas in connection with an investigation of the possible submission of false or otherwise improper reporting used to calculate the Medicaid rebates paid to the State of Texas. The precise parameters of this inquiry are unknown, and AstraZeneca is not in a position at this time to predict the scope, duration or outcome of this matter, including whether it will result in any liability to AstraZeneca.

Serbia
In August 2011, AstraZeneca UK Limited's Representative Office in Belgrade, Serbia was served with a criminal indictment relating to allegations that local employees of AstraZeneca made allegedly improper payments to physicians at the Institute of Oncology and Radiology of Serbia. AstraZeneca has filed a number of preliminary procedural objections, which ask the Serbian criminal court to dismiss the indictment against the Representative Office.

6 NINE MONTHS TERRITORIAL REVENUE ANALYSIS

	9 Months 2011 $m	9 Months 2010 $m	% Growth Actual	% Growth Constant Currency
US	9,783	10,273	(5)	(5)
Western Europe[1]	6,496	6,821	(5)	(10)
Canada	1,241	1,102	13	6
Japan	2,138	1,854	15	4
Other Established ROW	922	745	24	6
Established ROW[2]	4,301	3,701	16	5
Emerging Europe	927	859	8	5
China	947	780	21	16
Emerging Asia Pacific	732	651	12	6
Other Emerging ROW	1,749	1,567	12	11
Emerging ROW[3]	4,355	3,857	13	10
Total Revenue	24,935	24,652	1	(3)

[1] Western Europe comprises France, Germany, Italy, Sweden, UK and others.
[2] Established ROW comprises Australia, Canada, Japan and New Zealand.
[3] Emerging ROW comprises Brazil, China, India, Mexico, Russia, Turkey and all other ROW countries.

7 THIRD QUARTER TERRITORIAL REVENUE ANALYSIS

	3rd Quarter 2011 $m	3nd Quarter 2010 $m	% Growth Actual	% Growth Constant Currency
US	3,187	3,179	-	-
Western Europe[1]	2,067	2,150	(4)	(15)
Canada	401	379	6	(1)
Japan	771	632	22	10
Other Established ROW	332	251	32	11
Established ROW[2]	1,504	1,262	19	7
Emerging Europe	291	263	11	4
China	322	269	20	13
Emerging Asia Pacific	248	222	12	5
Other Emerging ROW	594	553	7	7
Emerging ROW[3]	1,455	1,307	11	7
Total Revenue	8,213	7,898	4	(2)

[1] Western Europe comprises France, Germany, Italy, Sweden, UK and others.
[2] Established ROW comprises Australia, Canada, Japan and New Zealand.
[3] Emerging ROW comprises Brazil, China, India, Mexico, Russia, Turkey and all other ROW countries.

8 NINE MONTHS PRODUCT REVENUE ANALYSIS

	World			US		Western Europe			Established ROW			Emerging ROW		
	9 Months 2011 $m	Actual Growth %	Constant Currency Growth %	9 Months 2011 $m	Actual Growth %	9 Months 2011 $m	Actual Growth %	Constant Currency Growth %	9 Months 2011 $m	Actual Growth %	Constant Currency Growth %	9 Months 2011 $m	Actual Growth %	Constant Currency Growth %
Gastrointestinal:														
Nexium	3,362	(10)	(12)	1,783	(12)	617	(32)	(36)	408	24	12	554	19	18
Losec/Prilosec	698	(6)	(13)	30	(24)	185	(7)	(14)	315	1	(9)	168	(14)	(17)
Others	112	5	2	67	3	35	6	-	6	20	20	4	-	(25)
Total Gastrointestinal	**4,172**	**(9)**	**(12)**	**1,880**	**(12)**	**837**	**(27)**	**(31)**	**729**	**13**	**2**	**726**	**9**	**8**
Cardiovascular:														
Crestor	4,851	18	14	2,231	18	920	12	5	1,197	27	16	503	11	8
Atacand	1,104	-	(5)	139	(16)	548	-	(6)	173	5	(4)	244	5	4
Seloken/Toprol-XL	750	(22)	(24)	315	(45)	63	(6)	(13)	28	(3)	(14)	344	19	14
Plendil	196	2	(2)	7	(42)	18	(14)	(19)	10	-	(10)	161	8	4
Tenormin	202	(2)	(9)	9	(10)	45	(2)	(9)	91	(1)	(11)	57	(2)	(5)
Zestril	109	(7)	(11)	8	-	54	(11)	(16)	12	(8)	(15)	35	-	(3)
Onglyza™	140	278	278	103	243	24	380	380	4	300	300	9	800	800
Brilinta/Brilique	16	n/m	n/m	11	n/m	4	n/m	n/m	-	-	-	1	n/m	n/m
Others	190	(3)	(8)	-	(100)	91	6	(1)	18	-	(11)	81	7	4
Total Cardiovascular	**7,558**	**9**	**5**	**2,823**	**5**	**1,767**	**7**	**-**	**1,533**	**21**	**10**	**1,435**	**11**	**7**
Respiratory:														
Symbicort	2,309	15	10	604	14	1,075	6	-	295	54	39	335	24	20
Pulmicort	669	5	1	218	(8)	143	(9)	(15)	86	10	-	222	34	30
Rhinocort	162	(7)	(11)	58	(22)	29	(3)	(10)	15	36	18	60	-	(5)
Others	162	(16)	(22)	6	(84)	83	(6)	(13)	19	6	-	54	6	-
Total Respiratory	**3,302**	**10**	**5**	**886**	**1**	**1,330**	**3**	**(3)**	**415**	**39**	**26**	**671**	**22**	**18**
Oncology:														
Arimidex	590	(52)	(55)	37	(92)	215	(51)	(53)	224	8	(2)	114	(1)	(4)
Zoladex	881	8	4	31	(9)	199	(5)	(10)	357	10	(1)	294	20	24
Casodex	408	(5)	(13)	(1)	(107)	63	(28)	(32)	262	4	(6)	84	8	5
Iressa	405	46	35	2	(33)	93	221	200	144	13	2	166	41	33
Others	482	57	52	200	94	151	57	48	47	12	-	84	27	24
Total Oncology	**2,766**	**(10)**	**(14)**	**269**	**(57)**	**721**	**(16)**	**(21)**	**1,034**	**8**	**(2)**	**742**	**19**	**18**
Neuroscience:														
Seroquel IR	3,190	2	-	2,434	4	418	-	(7)	165	(6)	(16)	173	(10)	(14)
Seroquel XR	1,092	30	26	565	18	363	44	35	66	57	43	98	46	43
Local Anaesthetics	454	2	(5)	10	(58)	185	(5)	(11)	151	14	2	108	16	11
Zomig	312	(2)	(6)	117	(10)	131	2	(5)	52	4	(4)	12	33	22
Diprivan	227	(6)	(12)	12	(68)	33	(15)	(21)	63	19	9	119	7	1
Vimovo	20	300	280	14	180	2	n/m	n/m	3	n/m	n/m	1	n/m	n/m
Others	26	(10)	(14)	1	-	14	(30)	(35)	3	-	-	8	50	50
Total Neuroscience	**5,321**	**6**	**4**	**3,153**	**5**	**1,146**	**9**	**2**	**503**	**11**	**-**	**519**	**9**	**4**
Infection & Other:														
Synagis	564	(12)	(12)	309	(16)	254	(6)	(6)	-	-	-	1	-	-
Merrem	469	(26)	(29)	33	(69)	151	(42)	(45)	46	12	2	239	6	2
FluMist	127	3	3	126	3	-	-	-	-	-	-	1	-	-
Others	101	(17)	(19)	58	(33)	9	100	50	14	40	(10)	20	(4)	17
Total Infection & Other	**1,261**	**(17)**	**(18)**	**526**	**(23)**	**414**	**(23)**	**(24)**	**60**	**18**	**-**	**261**	**5**	**4**
Aptium Oncology	169	2	2	169	2	-	-	-	-	-	-	-	-	-
Astra Tech	386	(1)	(6)	77	3	281	(1)	(7)	27	(7)	(21)	1	-	-
Total	**24,935**	**1**	**(3)**	**9,783**	**(5)**	**6,496**	**(5)**	**(10)**	**4,301**	**16**	**5**	**4,355**	**13**	**10**

9 THIRD QUARTER PRODUCT REVENUE ANALYSIS

	World			US		Western Europe			Established ROW			Emerging ROW		
	3rd Quarter 2011 $m	Actual Growth %	Constant Currency Growth %	3rd Quarter 2011 $m	Actual Growth %	3rd Quarter 2011 $m	Actual Growth %	Constant Currency Growth %	3rd Quarter 2011 $m	Actual Growth %	Constant Currency Growth %	3rd Quarter 2011 $m	Actual Growth %	Constant Currency Growth %
Gastrointestinal:														
Nexium	1,089	(12)	(16)	570	(16)	163	(42)	(50)	169	52	35	187	12	12
Losec/Prilosec	224	(4)	(13)	9	13	58	(3)	(15)	102	-	(11)	55	(13)	(17)
Others	37	-	(5)	20	(13)	13	18	9	2	-	-	2	100	-
Total Gastrointestinal	**1,350**	**(11)**	**(15)**	**599**	**(16)**	**234**	**(34)**	**(42)**	**273**	**27**	**13**	**244**	**6**	**4**
Cardiovascular:														
Crestor	1,659	21	14	753	20	307	16	2	431	31	17	168	10	7
Atacand	364	1	(8)	44	(15)	188	11	(4)	51	(9)	(20)	81	-	(2)
Seloken/Toprol-XL	273	-	(4)	123	(17)	22	5	(10)	9	(10)	(20)	119	28	20
Plendil	66	5	-	3	(25)	6	-	(17)	4	-	-	53	8	4
Tenormin	68	1	(7)	3	-	15	7	(7)	31	-	(10)	19	-	(5)
Zestril	37	6	(3)	3	50	18	(5)	(16)	3	(25)	(50)	13	30	30
Onglyza™	59	211	211	44	175	9	350	350	2	100	100	4	n/m	n/m
Brilinta/Brilique	13	n/m	n/m	11	n/m	2	n/m	n/m	-	-	-	-	-	-
Others	61	3	(5)	-	-	28	8	(8)	6	20	-	27	(4)	(4)
Total Cardiovascular	**2,600**	**16**	**9**	**984**	**15**	**595**	**14**	**-**	**537**	**22**	**9**	**484**	**12**	**8**
Respiratory:														
Symbicort	755	18	9	201	15	351	16	3	97	39	23	106	15	9
Pulmicort	185	3	(3)	52	(15)	40	(7)	(19)	27	4	(8)	66	32	28
Rhinocort	52	(5)	(13)	15	(29)	8	-	(13)	6	20	-	23	10	-
Others	52	(15)	(23)	2	(85)	27	-	(11)	7	17	-	16	7	-
Total Respiratory	**1,044**	**12**	**4**	**270**	**-**	**426**	**12**	**(1)**	**137**	**28**	**13**	**211**	**19**	**12**
Oncology:														
Arimidex	176	(38)	(44)	8	(81)	54	(59)	(65)	77	10	(1)	37	(3)	(5)
Zoladex	304	13	8	9	(31)	67	5	(5)	123	14	4	105	27	29
Casodex	137	-	(9)	-	(100)	18	(31)	(42)	90	7	(4)	29	21	17
Iressa	145	42	29	1	-	34	143	114	50	14	2	60	40	30
Others	168	56	47	68	94	53	56	38	16	7	(7)	31	29	25
Total Oncology	**930**	**3**	**(4)**	**86**	**(9)**	**226**	**(17)**	**(26)**	**356**	**11**	**-**	**262**	**24**	**21**
Neuroscience:														
Seroquel IR	1,034	1	(2)	791	1	138	6	(7)	56	4	(9)	49	(18)	(22)
Seroquel XR	366	31	24	184	18	124	43	25	23	53	40	35	67	57
Local Anaesthetics	149	7	(4)	1	(83)	59	4	(9)	55	25	11	34	6	(3)
Zomig	108	5	(3)	40	(5)	46	12	(2)	17	(6)	(11)	5	150	100
Diprivan	71	(16)	(24)	-	(100)	10	(9)	(18)	21	-	(5)	40	-	(10)
Vimovo	10	100	80	6	20	2	n/m	n/m	2	n/m	n/m	-	-	-
Others	7	(22)	(22)	-	-	3	(33)	(33)	-	-	-	4	-	-
Total Neuroscience	**1,745**	**6**	**2**	**1,022**	**2**	**382**	**15**	**1**	**174**	**14**	**3**	**167**	**6**	**(3)**
Infection & Other:														
Synagis	108	(22)	(22)	8	(27)	100	(22)	(22)	-	-	-	-	-	-
Merrem	139	(32)	(37)	5	(86)	39	(50)	(56)	13	8	-	82	4	(1)
FluMist	124	3	3	124	4	-	-	-	-	-	-	-	**-**	**-**
Others	29	(3)	-	14	(7)	3	n/m	n/m	7	75	75	5	(60)	(40)
Total Infection & Other	**400**	**(19)**	**(21)**	**151**	**(16)**	**142**	**(30)**	**(33)**	**20**	**25**	**19**	**87**	**(7)**	**(9)**
Aptium Oncology	56	33	33	56	33	-	-	-	-	-	-	-	-	-
Astra Tech	88	(28)	(36)	19	(24)	62	(29)	(37)	7	(30)	(50)	-	(100)	(100)
Total	**8,213**	**4**	**(2)**	**3,187**	**-**	**2,067**	**(4)**	**(15)**	**1,504**	**19**	**7**	**1,455**	**11**	**7**

Shareholder Information

ANNOUNCEMENTS AND MEETINGS

Announcement of fourth quarter and full year 2011 results	2 February 2012
Announcement of first quarter 2012 results	26 April 2012
Annual General Meeting	26 April 2012
Announcement of second quarter and half year 2012 results	26 July 2012
Announcement of third quarter and nine months 2012 results	25 October 2012

DIVIDENDS

The record date for the first interim dividend payable on 12 September 2011 was 5 August 2011. Shares traded ex-dividend from 3 August 2011.

The record date for the second interim dividend for 2011, payable on 19 March 2012 will be 17 February 2012. Shares will trade ex-dividend from 15 February 2012.

Future dividends will normally be paid as follows:

First interim	Announced in July and paid in September
Second interim	Announced in January and paid in March

TRADEMARKS

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CAUTIONARY STATEMENT REGARDING FORWARD-LOOKING STATEMENTS

In order, among other things, to utilise the 'safe harbour' provisions of the US Private Securities Litigation Reform Act 1995, we are providing the following cautionary statement: The interim financial statements contain certain forward-looking statements with respect to the operations, performance and financial condition of the Group. Although we believe our expectations are based on reasonable assumptions, any forward-looking statements, by their very nature, involve risks and uncertainties and may be influenced by factors that could cause actual outcomes and results to be materially different from those predicted. The forward-looking statements reflect knowledge and information available at the date of preparation of the interim financial statements and AstraZeneca undertakes no obligation to update these forward-looking statements. We identify the forward-looking statements by using the words 'anticipates', 'believes', 'expects', 'intends' and similar expressions in such statements. Important factors that could cause actual results to differ materially from those contained in forward-looking statements, certain of which are beyond our control, include, among other things: the loss or expiration of patents, marketing exclusivity or trademarks; the risk of substantial adverse litigation/government investigation claims and insufficient insurance coverage; exchange rate fluctuations; the risk that R&D will not yield new products that achieve commercial success; the risk that strategic alliances and acquisitions will be unsuccessful; the impact of competition, price controls and price reductions; taxation risks; the risk of substantial product liability claims; the impact of any failure by third parties to supply materials or services; the risk of failure to manage a crisis; the risk of delay to new product launches; the difficulties of obtaining and maintaining regulatory approvals for products; the risk of failure to observe ongoing regulatory oversight; the risk that new products do not perform as we expect; the risk of environmental liabilities; the risks associated with conducting business in emerging markets; the risk of reputational damage; the risk of product counterfeiting; the risk that regulatory approval processes for biosimilars could have an adverse effect on future commercial prospects; and the impact of increasing implementation and enforcement of more stringent anti-bribery and anti-corruption legislation.