

NOTICES CONCERNING THE EUROPEAN ECONOMIC AREA

EFTA SURVEILLANCE AUTHORITY

Medicinal products — List of marketing authorisations granted by the EEA EFTA States for the first half of 2008

(2009/C 52/05)

Subcommittee I — On the free movement of goods**To be noted by the EEA Joint Committee**

With reference to EEA Joint Committee Decision No 74/1999 of 28 May 1999, the EEA Joint Committee is invited to note the following lists concerning marketing authorisations for medicinal products for the period 1 January-30 June 2008, at their meeting on 7 November 2008:

<i>Annex I</i>	List of new marketing authorisations
<i>Annex II</i>	List of renewed marketing authorisations
<i>Annex III</i>	List of extended marketing authorisations
<i>Annex IV</i>	List of withdrawn marketing authorisations

ANNEX I

List of new marketing authorisations

The following marketing authorisations have been granted in the EEA EFTA States during the period **1 January-30 June 2008**:

EU-Number	Product	Country	Date of authorisation
EU/1/05/312/001	Xyrem	Liechtenstein	30.6.2008
EU/1/06/334/001-004	Evoltra	Liechtenstein	30.6.2008
EU/1/06/370/001-024	Exforge	Liechtenstein	30.6.2008
EU/1/06/371/001-024	Dafiro	Liechtenstein	30.6.2008
EU/1/06/372/001-024	Copalia	Liechtenstein	30.6.2008
EU/1/06/373/001-024	Imprida	Liechtenstein	30.6.2008
EU/1/07/001-019/IS	Retacrit	Iceland	16.1.2008
EU/1/07/393/001	Soliris	Liechtenstein	30.6.2008
EU/1/07/400/001-016	Mircera	Liechtenstein	30.6.2008
EU/1/07/402/001/NO	Increlex	Norway	31.1.2008
EU/1/07/421/001-009/IS	Pioglitazone	Iceland	9.1.2008
EU/1/07/422/001/NO-004/NO	Tasigna	Norway	17.1.2008
EU/1/07/422/001-004	Tasigna	Liechtenstein	30.6.2008
EU/1/07/423/001/NO-003/NO	Vectibix	Norway	21.1.2008
EU/1/07/423/001-003	Vectibix	Liechtenstein	30.6.2008
EU/1/07/423/001-003/IS	Vectibix	Iceland	28.1.2008
EU/1/07/427/001/NO-037/NO	Olanzapine Teva	Norway	7.1.2008
EU/1/07/427/001-047/IS	Olanzapine Teva	Iceland	7.1.2008
EU/1/07/428/001	Abraxane	Liechtenstein	29.2.2008
EU/1/07/428/001/IS	Abraxane	Iceland	24.1.2008
EU/1/07/428/001/NO	Abraxane	Norway	21.1.2008
EU/1/07/430/001	Atripla	Liechtenstein	30.6.2008
EU/1/07/430/001/IS	Atripla	Iceland	28.1.2008
EU/1/07/430/001/NO	Atripla	Norway	29.1.2008
EU/1/07/431/001/NO-019/NO	Retacrit	Norway	17.1.2008
EU/1/07/431/001-019	Retacrit	Liechtenstein	30.6.2008
EU/1/07/432/001/NO-019/NO	Silapo	Norway	17.1.2008
EU/1/07/432/001-019	Silapo	Liechtenstein	29.2.2008
EU/1/07/432/001-019/IS	Silapo	Iceland	16.1.2008
EU/1/07/433/001/IS	Nevanac	Iceland	8.1.2008
EU/1/07/434/001/NO-003/NO	Avamys	Norway	23.1.2008

EU-Number	Product	Country	Date of authorisation
EU/1/07/434/001-003	Avamys	Liechtenstein	29.2.2008
EU/1/07/434/001-003/IS	Avamys	Iceland	25.1.2008
EU/1/07/435/001/NO-018/NO	Tesavel	Norway	24.1.2008
EU/1/07/435/001-018	Tesavel	Liechtenstein	30.4.2008
EU/1/07/435/001-018/IS	Tesavel	Iceland	24.1.2008
EU/1/07/436/001/NO-002/NO	Isentress	Norway	16.1.2008
EU/1/07/436/001-002	Isentress	Liechtenstein	29.2.2008
EU/1/07/436/001-002/IS	Isentress	Iceland	18.1.2008
EU/1/07/437/001/NO-002/NO	Ivemend	Norway	31.1.2008
EU/1/07/437/001-002	Ivemend	Liechtenstein	29.2.2008
EU/1/07/437/001-002/IS	Ivemend	Iceland	22.1.2008
EU/1/07/438/001/NO-004/NO	Myfenac	Norway	14.3.2008
EU/1/07/438/001-004	Myfenax	Liechtenstein	30.4.2008
EU/1/07/438/001-004/IS	Myfenax	Iceland	17.3.2008
EU/1/07/439/001/NO-004/NO	Mycophenolate mofetil Teva	Norway	14.3.2008
EU/1/07/439/001-004	Mycophenolate mofetil Teva	Liechtenstein	30.4.2008
EU/1/07/439/001-004/IS	Mycophenolate mofetil Teva	Iceland	17.3.2008
EU/1/07/440/001/NO-002/NO	Tyverb	Norway	17.6.2008
EU/1/07/440/001-002/IS	Tyverb	Iceland	26.6.2008
EU/1/08/441/001/NO-010/NO	Effentora	Norway	15.4.2008
EU/1/08/441/001-010	Effentora	Liechtenstein	30.4.2008
EU/1/08/441/001-010/IS	Effentora	Iceland	17.4.2008
EU/1/08/442/001/NO-008/NO	Pradaxa	Norway	14.4.2008
EU/1/08/442/001-008	Pradaxa	Liechtenstein	30.4.2008
EU/1/08/442/001-008/IS	Pradaxa	Iceland	26.3.2008
EU/1/08/443/001	Thalidomide Pharmion	Liechtenstein	30.4.2008
EU/1/08/443/001/IS	Thalidomide Pharmion	Iceland	21.4.2008
EU/1/08/443/001/NO	Thalidomide Pharmion	Norway	16.5.2008
EU/1/08/446/001/NO-003/NO	Privigen	Norway	19.5.2008
EU/1/08/446/001-003	Privigen	Liechtenstein	30.6.2008
EU/1/08/446/001-003/IS	Privigen	Iceland	23.5.2008
EU/1/08/447/001/NO-004/NO	Adenuric	Norway	24.6.2008
EU/1/08/447/001-004	Adenuric	Liechtenstein	30.4.2008
EU/1/08/447/001-004/IS	Adenuric	Iceland	16.5.2008
EU/1/08/448/001/NO-002/NO	Mycamine	Norway	21.5.2008

EU-Number	Product	Country	Date of authorisation
EU/1/08/448/001-002	Mycamine	Liechtenstein	30.4.2008
EU/1/08/448/001-002/IS	Mycamine	Iceland	28.5.2008
EU/1/08/451/001/NO-004/NO	Volibris	Norway	29.4.2008
EU/1/08/451/001-002/IS	Volibris	Iceland	21.5.2008
EU/1/08/451/001-004	Volibris	Liechtenstein	30.4.2008
EU/1/08/452/001	Pandemrix	Liechtenstein	30.6.2008
EU/1/08/452/001/IS	Pandemrix	Iceland	23.5.2008
EU/1/08/452/001/NO	Pandemrix	Norway	20.5.2008
EU/1/08/453/001	Prepandrix	Liechtenstein	30.6.2008
EU/1/08/453/001/IS	Prepandrix	Iceland	23.5.2008
EU/1/08/454/001/NO-004/NO	Extavia	Norway	3.6.2008
EU/1/08/454/001-004	Extavia	Liechtenstein	30.6.2008
EU/1/08/454/001-004/IS	Extavia	Iceland	30.5.2008
EU/2/08/081/001-003	Posatex	Liechtenstein	30.6.2008
EU/2/04/044/007/IS	Aivlosin	Iceland	19.6.2008
EU/2/07/073/001/NO-004/NO	Nobilis Influenza H7N1	Norway	4.1.2008
EU/2/07/076/001/NO-004/NO	Nobilis Influenza H5N6	Norway	13.3.2008
EU/2/07/076/001-004	Nobilis Influenza H5N6	Liechtenstein	29.2.2008
EU/2/07/077/001/NO-005/NO	Meloxivet	Norway	7.2.2008
EU/2/07/077/001-004/IS	Meloxivet	Iceland	27.6.2008
EU/2/07/078/001/NO-003/NO	Rheumocam	Norway	25.2.2008
EU/2/07/078/001-002/IS	Rheumocam	Iceland	25.1.2008
EU/2/07/078/001-003	Rheumocam	Liechtenstein	29.2.2008
EU/2/07/079/001/NO-004/NO	Ingelvac CircoFLEX	Norway	26.3.2008
EU/2/07/079/001-004	Ingelvac CircoFLEX	Liechtenstein	29.2.2008
EU/2/07/079/001-004/IS	Ingelvac CircoFLEX	Iceland	19.3.2008

ANNEX II

List of renewed marketing authorisations

The following marketing authorisations have been renewed in the EEA EFTA States during the period **1 January-30 June 2008**:

EU-Number	Product	Country	Date of authorisation
EU/1/02/246/001/NO-003/NO	Carbaglu	Norway	30.5.2008
EU/1/02/246/001-003	Carbaglu	Liechtenstein	30.6.2008
EU/1/02/246/001-003/IS	Carbaglu	Iceland	29.5.2008
EU/1/03/247/001/NO-002/NO	Forsteo	Norway	1.7.2008
EU/1/03/247/001-002	Forsteo	Liechtenstein	30.6.2008
EU/1/03/247/001-002/IS	Forsteo	Iceland	21.5.2008
EU/1/03/248/001/NO-012/NO	Levitra	Norway	25.2.2008
EU/1/03/248/001-012/IS	Levitra	Iceland	19.2.2008
EU/1/03/249/001/NO-012/NO	Vivanza	Norway	25.2.2008
EU/1/03/249/001-012	Vivanza	Liechtenstein	29.2.2008
EU/1/03/249/001-012/IS	Vivanza	Iceland	19.2.2008
EU/1/03/250/001	Ytracis	Liechtenstein	29.2.2008
EU/1/03/250/001/IS	Ytracis	Iceland	28.2.2008
EU/1/03/250/001/NO	Ytracis	Norway	25.2.2008
EU/1/03/251/001/NO-002/NO	Hepsera	Norway	4.4.2008
EU/1/03/251/001-002	Hepsera	Liechtenstein	30.4.2008
EU/1/03/251/001-002/IS	Hepsera	Iceland	10.4.2008
EU/1/03/253/001/NO-003/NO	Aldurazyme	Norway	30.5.2008
EU/1/03/253/001-003	Aldurazyme	Liechtenstein	30.6.2008
EU/1/03/253/001-003/IS	Aldurazyme	Iceland	22.5.2008
EU/1/06/367/001/NO-012/NO	Diacomit	Norway	7.1.2008
EU/1/06/367/001-012	Diacomit	Liechtenstein	29.2.2008
EU/1/06/367/001-012/IS	Diacomit	Iceland	7.1.2008
EU/1/06/380/001	Prezista	Liechtenstein	29.2.2008
EU/1/06/380/001/IS	Prezista	Iceland	22.1.2008
EU/1/06/380/001/NO	Prezista	Norway	21.1.2008
EU/1/97/045/001-004	Helicobacter Test INFAI	Liechtenstein	30.4.2008
EU/1/97/045/001-004/IS	Helicobacter Test INFAI	Iceland	9.4.2008

EU-Number	Product	Country	Date of authorisation
EU/1/97/054/001&004-005/IS	Viracept	Iceland	21.1.2008
EU/1/97/054/001, 004-005	Viracept	Liechtenstein	29.2.2008
EU/1/97/054/001/NO, 004/NO-005/NO	Viracept	Norway	17.1.2008
EU/1/97/055/001/NO-003/NO	Viramune	Norway	21.1.2008
EU/1/97/055/001-003	Viramune	Liechtenstein	30.4.2008
EU/1/97/055/001-003/IS	Viramune	Iceland	22.1.2008
EU/1/97/057/001/IS	Quadramet	Iceland	7.1.2008
EU/1/97/057/001/NO	Quadramet	Norway	17.1.2008
EU/1/98/058/001/NO-002/NO	Combivir	Norway	26.2.2008
EU/1/98/058/001-002	Combivir	Liechtenstein	29.2.2008
EU/1/98/058/001-002/IS	Combivir	Iceland	12.3.2008
EU/1/98/063/001/NO-006/NO	Rebif	Norway	3.6.2008
EU/1/98/063/001-007	Rebif	Liechtenstein	30.6.2008
EU/1/98/063/001-007/IS	Rebif	Iceland	2.6.2008
EU/1/98/065/001-002	Otpison	Liechtenstein	30.6.2008
EU/1/98/066/001/NO-026/NO	Exelon	Norway	3.6.2008
EU/1/98/066/001-026	Exelon	Liechtenstein	30.6.2008
EU/1/98/066/001-026/IS	Exelon	Iceland	29.5.2008
EU/1/98/067/001/NO-002/NO	MabThera	Norway	24.6.2008
EU/1/98/067/001-002	Mabthera	Liechtenstein	30.6.2008
EU/1/98/067/001-002/IS	MabThera	Iceland	29.5.2008
EU/1/98/069/001a/NO-007a/NO; EU/1/98/069/001b/NO-007b/NO; EU/1/98/069/008/NO-010/NO	Plavix	Norway	2.7.2008
EU/1/98/069/001a-001b, 002a-002b, 003a-003b, -004a-004b, 005a-005b, 006a-006b, 007a-007b, 008-010	Plavix	Liechtenstein	30.6.2008
EU/1/98/069/001a-007a& 001b-007b&008-010/IS	Plavix	Iceland	27.6.2008
EU/1/98/070/001a/NO-007a/NO; EU/1/98/070/001b/NO-007b/NO; EU/1/98/070/008/NO-010/NO	Iscover	Norway	2.7.2008

EU-Number	Product	Country	Date of authorisation
EU/1/98/070/001a-001b, 002a-002b, 003a-003b, -004a-004b, 005a-005b, 006a-006b, 007a-007b, 008-010	Iscover	Liechtenstein	30.6.2008
EU/1/98/070/001a-007a& 001b-007b&008-010/IS	Iscover	Iceland	27.6.2008
EU/1/98/071/001-006	Xenical	Liechtenstein	30.6.2008
EU/1/98/092/001/NO-023/NO	Prometax	Norway	3.6.2008
EU/1/98/092/001-026	Prometax	Liechtenstein	30.6.2008
EU/1/98/092/001-026/IS	Prometax	Iceland	16.6.2008
EU/2/02/032/001/NO	Vaxxitek HVT + IBD	Norway	24.1.2008
EU/2/02/033/001/NO	Dexdomitor	Norway	21.1.2008
EU/2/02/034/001/NO	Nobivac Bb for cats	Norway	24.1.2008
EU/2/02/035/001/NO-007/NO	SevoFlo	Norway	17.1.2008
EU/2/02/036/001/NO-002/NO	Nobilis OR inac	Norway	26.2.2008
EU/2/02/036/001-002	Nobilis OR inac	Liechtenstein	30.4.2008
EU/2/02/036/001-002/IS	Nobilis OR inac	Iceland	21.1.2008
EU/2/03/037/005	ProteqFlu	Liechtenstein	29.2.2008
EU/2/03/037/005/IS	ProteqFlu	Iceland	21.1.2008
EU/2/03/037/005/NO	ProteqFlu	Norway	27.3.2008
EU/2/03/038/005	ProteqFlu-Te	Liechtenstein	29.2.2008
EU/2/03/038/005/IS	ProteqFlu-Te	Iceland	21.1.2008
EU/2/03/038/005/NO	ProteqFlu Te	Norway	27.3.2008
EU/2/03/039/001/NO-030/NO	Advocate	Norway	22.4.2008
EU/2/03/039/001-004&013-14&019-022/IS	Advocate for cats	Iceland	13.3.2008
EU/2/03/039/001-030	Advocate	Liechtenstein	29.2.2008
EU/2/03/039/005-012&015-018&023-030/IS	Advocate for dogs	Iceland	13.3.2008
EU/2/03/040/001-002	Gonazon	Liechtenstein	30.6.2008
EU/2/96/002/001/NO-003/NO	Fevaxyn Pentofel	Norway	12.2.2008
EU/2/97/001-018/IS	Dicural	Iceland	21.1.2008
EU/2/97/003/001/NO-018/NO	Dicural	Norway	25.2.2008
EU/2/97/003/001-018	Dicural	Liechtenstein	29.2.2008

EU-Number	Product	Country	Date of authorisation
EU/2/97/004/001&010/IS	Metacam for cattle and pigs	Iceland	4.1.2008
EU/2/97/004/003-005&012-013&016-025&029/IS	Metacam for dogs	Iceland	4.1.2008
EU/2/97/004/006&011/IS	Metacam for dogs and cats	Iceland	4.1.2008
EU/2/97/004/007-008&014-015&027-028&031-032/IS	Metacam for cattle, pigs and horses	Iceland	4.1.2008
EU/2/97/004/009&030/IS	Metacam for horses	Iceland	4.1.2008
EU/2/97/004/026/IS	Metcam for cats	Iceland	4.1.2008
EU/2/98/006/001-010	Nobilis IB 4-91	Liechtenstein	30.6.2008
EU/2/98/006/001-010/IS	Nobilis IB 4-91	Iceland	23.5.2008
EU/2/98/007/001/NO-003/NO	Clomicalm	Norway	26.5.2008
EU/2/98/007/001-003	Clomicalm	Liechtenstein	30.4.2008
EU/2/98/007/001-003/IS	Clomicalm	Iceland	21.4.2008
EU/2/98/008/001/NO-004/NO	Neocolipor	Norway	21.4.2008
EU/2/98/008/001-004	Neocolipor	Liechtenstein	30.4.2008
EU/2/98/008/001-004/IS	Neocolipor	Iceland	9.4.2008

ANNEX III

List of extended marketing authorisations

The following marketing authorisations have been extended in the EEA EFTA States during the period **1 January-30 June 2008**:

EU-Number	Product	Country	Date of extension
EU/1/00/143/010/NO-011/NO	Kogenate Bayer	Norway	28.1.2008
EU/1/00/143/010-011	Kogenate Bayer	Liechtenstein	29.2.2008
EU/1/00/143/010-011/IS	Kogenate Bayer	Iceland	16.1.2008
EU/1/00/144/004	Helixate NexGen	Liechtenstein	29.2.2008
EU/1/00/144/004/IS	Helixate NexGen	Iceland	16.1.2008
EU/1/00/144/004/NO	Helixate NexGen	Norway	28.1.2008
EU/1/00/167/008	Prevenar	Liechtenstein	30.6.2008
EU/1/01/172/006	Kaletra	Liechtenstein	30.4.2008
EU/1/01/172/006/IS	Kaletra	Iceland	14.4.2008
EU/1/01/172/006/NO	Kaletra	Norway	14.4.2008
EU/1/01/195/016/NO-021/NO	Liprolog	Norway	29.1.2008
EU/1/01/195/016-021	Liprolog	Liechtenstein	29.2.2008
EU/1/01/200/002	Viread	Liechtenstein	30.4.2008
EU/1/01/200/002/NO	Viread	Norway	23.4.2008
EU/1/02/213/017-023	MicardisPlus	Liechtenstein	30.4.2008
EU/1/02/213/017/NO-023/NO	MicardisPlus	Norway	27.5.2008
EU/1/02/213/017-023/IS	MicardisPlus	Iceland	15.4.2008
EU/1/02/214/011-015	Kinzalcomb	Liechtenstein	30.4.2008
EU/1/02/215/015-021	Pritor Plus	Liechtenstein	30.4.2008
EU/1/02/218/012-029	Axura	Liechtenstein	30.6.2008
EU/1/02/218/012-029/IS	Axura	Iceland	28.5.2008
EU/1/02/218/016/NO-023/NO	Axura	Norway	27.5.2008
EU/1/02/219/016-021	Ebixa	Liechtenstein	30.4.2008
EU/1/02/219/022/NO-049/NO	Ebixa	Norway	21.5.2008
EU/1/02/219/022-049	Ebixa	Liechtenstein	30.6.2008
EU/1/02/219/022-049/IS	Ebixa	Iceland	27.5.2008
EU/1/03/260/019/NO-023/NO	Stalevo	Norway	27.5.2008
EU/1/03/260/019-023	Stalevo	Liechtenstein	30.4.2008

EU-Number	Product	Country	Date of extension
EU/1/03/260/019-023/IS	Stalevo	Iceland	20.5.2008
EU/1/03/267/008/NO-009/NO	Reyataz	Norway	16.5.2008
EU/1/03/267/008-009	Reyataz	Liechtenstein	30.4.2008
EU/1/03/267/008-009/IS	Reyataz	Iceland	22.4.2008
EU/1/03/268/004	Cholestagel	Liechtenstein	30.6.2008
EU/1/03/271/001-006/IS	Advate	Iceland	30.5.2008
EU/1/03/271/005/NO-006/NO	Advate	Norway	3.6.2008
EU/1/03/271/005-006	Advate	Liechtenstein	30.6.2008
EU/1/04/274/002	Velcade	Liechtenstein	30.4.2008
EU/1/04/274/002/NO	Velcade	Norway	21.4.2008
EU/1/04/279/036-043	Lyrica	Liechtenstein	29.2.2008
EU/1/04/283/008/NO-012/NO	Ariclaim	Norway	17.1.2008
EU/1/04/283/008-012/IS	Ariclaim	Iceland	15.2.2008
EU/1/04/298/003	Kivexa	Liechtenstein	30.4.2008
EU/1/04/298/003/NO	Kivexa	Norway	29.2.2008
EU/1/06/357/018-021	Gardasil	Liechtenstein	30.4.2008
EU/1/06/358/018-021	Silgard	Liechtenstein	30.4.2008
EU/1/06/366/017/NO-022/NO	Tandemact	Norway	22.4.2008
EU/1/06/366/017-022	Tandemact	Liechtenstein	30.4.2008
EU/1/06/366/017-022/IS	Tandemact	Iceland	16.4.2008
EU/1/06/368/058-087	Insulin Human Winthrop	Liechtenstein	29.2.2008
EU/1/06/370/025-033	Exforge	Liechtenstein	30.6.2008
EU/1/06/371/025-033	Dafiro	Liechtenstein	30.6.2008
EU/1/06/372/025-033	Copalia	Liechtenstein	30.6.2008
EU/1/06/373/025-033	Imprida	Liechtenstein	30.6.2008
EU/1/07/387/009/NO-010/NO	Advagraf	Norway	25.1.2008
EU/1/07/387/009-010	Advagraf	Liechtenstein	29.2.2008
EU/1/07/388/002	Sebivo	Liechtenstein	29.2.2008
EU/1/07/388/002/NO	Sebivo	Norway	28.2.2008
EU/1/96/006/004/NO-006/NO	NovoSeven	Norway	21.5.2008

EU-Number	Product	Country	Date of extension
EU/1/96/006/004-006	NovoSeven	Liechtenstein	30.4.2008
EU/1/96/006/004-006/IS	NovoSeven	Iceland	22.5.2008
EU/1/96/007/031/NO-038/NO	Humalog	Norway	31.1.2008
EU/1/96/007/031-038	Humalog	Liechtenstein	29.2.2008
EU/1/96/027/006/NO-007/NO	Hycamtin	Norway	9.4.2008
EU/1/96/027/006-007	Hycamtin	Liechtenstein	30.4.2008
EU/1/96/027/006-007/IS	Hycamtin	Iceland	26.3.2008
EU/1/97/030/085-139	Insuman	Liechtenstein	29.2.2008
EU/1/98/069/008/NO-010/NO	Plavix	Norway	22.4.2008
EU/1/98/069/008-010	Plavix	Liechtenstein	30.4.2008
EU/1/98/069/008-010/IS	Plavix	Iceland	17.4.2008
EU/1/98/070/008/NO-010/NO	Iscover	Norway	19.5.2008
EU/1/98/070/008-010	Iscover	Liechtenstein	30.4.2008
EU/1/98/070/008-010/IS	Iscover	Iceland	17.4.2008
EU/1/99/119/015	NovoRapid	Liechtenstein	30.4.2008
EU/1/99/119/015/NO	NovoRapid	Norway	25.2.2008
EU/2/04/044/007	Aivlosin	Liechtenstein	30.6.2008
EU/2/04/045/007/NO	Previcox	Norway	29.1.2008
EU/2/97/004/031-032	Metacam	Liechtenstein	30.6.2008

ANNEX IV

List of withdrawn marketing authorisations

The following marketing authorisations have been withdrawn in the EEA EFTA States during the period **1 January-30 June 2008**:

EU-Number	Product	Country	Date of withdrawal
EU/1/01/192/001/NO-005/NO	Levviavx	Norway	24.1.2008
EU/1/02/207/001-020	Quixidar	Liechtenstein	30.4.2008
EU/1/02/207/001-020/IS	Quixidar	Iceland	11.3.2008
EU/1/02/207/001-020/NO	Quixidar	Norway	2.4.2008
EU/1/99/098/001-002	Zenapax	Liechtenstein	30.6.2008
EU/2/00/025/001-004	Advasure	Liechtenstein	30.6.2008