Summary of Community decisions on marketing authorizations in respect of medicinal products from 15 June 2005 to 30 June 2005

(Published pursuant to Article 12 or Article 34 of Council Regulation (EEC) No 2309/93 ([1]))

(2005/C 186/02)

— Modification of a marketing authorization (Article 12 of Council Regulation (EEC) No 2309/93): Accepted

Date of the decision	Name of the medicinal product	Holder of the marketing authorization	Number of the entry in the Community Register	Date of notification
16.6.2005	Tracleer	Actelion Registration Ltd, BSI Building 13th Floor, 389 Chiswick High Road, London W4 4AL, United Kingdom	EU/1/02/220/001-005	20.6.2005
20.6.2005	Abilify	Otsuka Pharmaceutical Europe Ltd, Commonwealth House, 2 Chalkhill Road, Hammersmith, London W6 8DW, United Kingdom	EU/1/04/276/021-032	22.6.2005
24.6.2005	Vistide	Pfizer Enterprises SARL, 51, Avenue Kennedy — Rond Point du Kirchberg, L-1855 Luxembourg, Grand-Duché de Luxembourg.	EU/1/97/037/001	28.6.2005
28.6.2005	Taxotere	Aventis Pharma S.A., 20 avenue Raymond Aron, 92165 Antony Cedex, France	EU/1/95/002/001-002	30.6.2005
28.6.2005	Invirase	Roche Registration Ltd., 40, Broad-water Road, Welwyn Garden City, Hertfordshire, AL7 3AY, United Kingdom	EU/1/96/026/001-002	30.6.2005
28.6.2005	Osigraft	Howmedica International S. de R.L., Division of Stryker Corporation, Raheen Industrial Estate, Raheen, Limerick, Ireland	EU/1/01/179/001	4.7.2005
28.6.2005	Herceptin	Roche Registration Limited, 40 Broad-water Road, Welwyn Garden City, Hertfordshire AL7 3AY, United Kingdom	EU/1/00/145/001	30.6.2005
28.6.2005	Remicade	Centocor B.V., Einsteinweg 101, 2333 CB Leiden, Nederland	EU/1/99/116/001-003	30.6.2005
28.6.2005	Remicade	Centocor B.V., Einsteinweg 101, 2333 CB Leiden, Nederland	EU/1/99/116/001-003	30.6.2005

— Withdrawal of a marketing authorization (Article 12 of Council Regulation (EEC) No 2309/93)

Date of the decision	Name of the medicinal product	Holder of the marketing authorization	Number of the entry in the Community Register	Date of notification
24.6.2005	Valdyn	Pfizer Limited, Sandwich, Kent, CT13 9NJ, United Kingdom	EU/1/02/244/001-024	28.6.2005
24.6.2005	Rayzon	Pharmacia Europe EEIG, Sandwich, Kent, CT13 9NJ, United Kingdom	EU/1/02/210/001-008	29.6.2005

([1]) OJ L 214 of 24.8.1993, p. 1.

— **Modification of a marketing authorization (Article 34 of Council Regulation (EEC) No 2309/93): Accepted**

Date of the decision	Name of the medicinal product	Holder of the marketing authorization	Number of the entry in the Community Register	Date of notification
16.6.2005	Aivlosin	ECO Animal Health Ltd, 78 Coombe Road New Malden Surrey KT3 4QS United Kingdom	EU/2/04/044/001	20.6.2005
21.6.2005	Dexdomitor	Orion Corporation, Orionintie 1, FIN-02200 Espoo, Suomi	EU/2/02/033/001	23.6.2005
24.6.2005	Dexdomitor	Orion Corporation, Orionintie 1, FIN-02200 Espoo, Suomi	EU/2/02/033/001	28.6.2005
24.6.2005	DRAXXIN	PFIZER Ltd, Ramsgate Road, Sandwich, Kent CT 13 9NJ, United Kingdom	EU/2/03/041/001-005	28.6.2005
24.6.2005	Metacam	Boehringer Ingelheim Vetmedica GmbH, 55216 Ingelheim am Rhein, Deutschland	EU/2/97/004/001-011	28.6.2005
24.6.2005	Porcilis Pesti	Intervet International B.V., Wim de Körverstraat 35, 5831 AN Boxmeer, Nederland	EU/2/99/016/001-006	28.6.2005

Anyone wishing to consult the public assessment report on the medicinal products in question and the decisions relating thereto is invited to contact:

The European Medicines Agency
7, Westferry Circus, Canary Wharf
London E14 4HB
United Kingdom