Summary of Community decisions on marketing authorizations in respect of medicinal products from 1 December 2007 to 31 December 2007

(Decisions taken pursuant to Article 34 of Directive 2001/83/EC of the European Parliament and of the Council [1] or Article 38 of Directive 2001/82/EC of the European Parliament and of the Council [2])

(2008/C 19/05)

— Issuing, maintenance or modification of a national marketing authorisation

Date of the decision	Name(s) of the medicinal product	Holder(s) of the marketing authorization	Member State concerned	Date of notification
18.12.2007	Tracleer	Actelion Registration Ltd BSI Building 13th Floor 389 Chiswick High Road London W4 4AL United Kingdom	This Decision is addressed to the Member States	19.12.2007
18.12.2007	Silapo	STADA Arzneimittel AG Stadastraße 2-18 D-61118 Bad Vilbel	This Decision is addressed to the Member States	19.12.2007
18.12.2007	Retacrit	HOSPIRA Enterprises B.V. Taurusavenue 19-21b 2132 LS Hoofddorp Nederland	This Decision is addressed to the Member States	19.12.2007
19.12.2007	MabCampath	Genzyme Europe B.V. Gooimeer 10 1411 DD Naarden Nederland	This Decision is addressed to the Member States	20.12.2007

[1] OJ L 311, 28.11.2001, p. 67.
[2] OJ L 311, 28.11.2001, p. 1.