Commission Regulation (EC) No 1662/95 of 7 July 1995 laying down certain detailed arrangements for implementing the Community decision-making procedures in respect of marketing authorizations for products for human or veterinary use

This Regulation establishes, within the framework of decisions relating to
Matters shall be referred to the Committee by the chairman
Where the opinion of the Committee is obtained by written
Where, in the opinion of the Commission, written comments
put
Where a Member State has applied the emergency procedure
laid
Where the draft decision needs to be examined at a
Correspondence to members of the Committee, where the
Committee in
This Regulation shall enter into force on the third day
Signature

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EC) No 1662/95. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

(1) OJ No L 214, 24. 8. 1993, p. 1.

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View outstanding changes

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Changes and effects yet to be applied to :

Regulation revoked in part by S.I. 2019/775 Sch. 9 para. 1(c)