



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

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Inspections Human Medicines Pharmacovigilance Division
Committees and Inspections Department

Scientific recommendation on classification of advanced therapy medicinal products

Article 17 – Regulation (EC) No 1394/2007

Disclaimer: This document is a summary for public release of a scientific recommendation on classification of advanced therapy medicinal products. The original text adopted by the Committee for Advanced Therapies (CAT) has been redacted to delete commercially confidential information.

The present scientific recommendation refers exclusively to the case as presented to the European Medicines Agency (EMA) without prejudice to future evaluations by the Agency.

It is stressed that the scientific recommendation on advanced therapy classification does not amount to any endorsement of the plausibility of the product, including the mode of action or therapeutic indication(s) claimed by the applicant.

Brief description (or name when available) of the active substance(s)

Autologous tumour-infiltrating lymphocytes.

Brief description of the finished product

Suspension of autologous tumour-infiltrating T lymphocytes for infusion.

Proposed indication

Treatment of metastatic melanoma.

EMA/CAT conclusion

The procedure was finalised on 23 February 2017 for the following recommendation.

On the basis that the product:

- contains cells that have been subject to substantial manipulation;



- therapeutic effect relates to a view to treating patients with metastatic melanoma,

the EMA/CAT considers that the Product falls within the definition of a somatic cell therapy medicinal product as provided in Article 2(1) of Regulation (EC) 1394/2007.