
STATUTORY INSTRUMENTS

2014 No. 2883

HUMAN TISSUE

The Human Tissue (Quality and Safety for Human Application) (Amendment) Regulations 2014

Made - - - - 29th October 2014
Laid before Parliament 7th November 2014
Coming into force - - 15th December 2014

The Secretary of State is a Minister designated⁽¹⁾ for the purposes of section 2(2) of the European Communities Act 1972⁽²⁾ in relation to health protection measures regulating the use of material of human origin.

The Secretary of State makes these Regulations in exercise of the powers conferred by section 2(2) of the European Communities Act 1972.

Citation and commencement

1. These Regulations may be cited as the Human Tissue (Quality and Safety for Human Application) (Amendment) Regulations 2014 and come into force on 15th December 2014.

Amendment of the Human Tissue (Quality and Safety for Human Application) Regulations 2007

2. In regulation 4 of the Human Tissue (Quality and Safety for Human Application) Regulations 2007⁽³⁾ (references to Directives), in the definition of “the second Directive”, after “cells,” insert “as amended by Commission Directive 2012/39/EU⁽⁴⁾”.

(1) [S.I. 2004/3037](#).

(2) [1972 c.68](#).

(3) [S.I. 2007/1523](#). There are amendments not relevant to this instrument.

(4) OJ No L 327, 27.11.2012, p24.

Status: This is the original version (as it was originally made). This item of legislation is currently only available in its original format.

Signed by authority of the Secretary of State for Health.

29th October 2014

Jane Ellison
Parliamentary Under-Secretary of State,
Department of Health

EXPLANATORY NOTE

(This note is not part of the Regulations)

These Regulations amend the Human Tissue (Quality and Safety for Human Application) Regulations 2007 (“Principal Regulations”) in order to transpose Commission Directive 2012/39/EU of 26 November 2012 amending [Directive 2006/17/EC](#) as regards certain technical requirements for the testing of human tissues and cells (“the Amending Directive”).

Regulation 2 amends the definition of “the second Directive” ([Directive 2006/17/EC](#)) in regulation 4 of the Principal Regulations to include the changes brought about by the Amending Directive.

An impact assessment has not been produced for this instrument as no impact on the private or voluntary sectors is foreseen.