

Directions given under the Human Tissue (Quality and Safety for Human Application) Regulations 2007 applying the Human Tissue Act 2004

Directions on interim licence fee structure for procurement for human application licences for the financial year 2008/09

Ref 001/2008

These Directions are

General Directions given in relation to licences to procure tissues and / or cells for human application under Regulation 7(2) of the Human Tissue (Quality and Safety for Human Application) Regulations 2007.

Section of the Human Tissue Act 2004 (HT Act) as applied by the Human Tissue (Quality and Safety for Human Application) Regulations 2007 (the Regulations) providing for these Directions	Section 37 and Schedule 3 paragraph 2 (4) (f) of the HT Act as applied by Regulation 8 of the Regulations
These Directions come into force on	5 July 2008
These Directions remain in force	Until revoked

1. These Directions are made under Section 37 and Schedule 3 Paragraph 2 (4) (f) of the Human Tissue Act 2004 (HT Act) as applied by Regulation 8 of the Human Tissue (Quality and Safety for Human Application) Regulations 2007 (“the Regulations”) in order to set out the interim licence fee structure of the Human Tissue Authority (HTA) for licences for procurement of tissues and/or cells for human application for the financial year 2008/09.
2. The HTA’s interim licence fee structure is set out below and comprises such sums in respect of its costs in connection with superintending compliance with the terms of licences for the purposes of Schedule 3

Paragraph 2 (4) (f) of the HT Act as applied by Regulation 8 of the Regulations.

2008/09 – Interim Licence Fee Structure for licences for procurement for human application

Sector	Main Site	Satellite
Procurement Organisations (Procurers of tissues and/or cells for human application)	£2,900 pa	£1,000 pa

3. From the date of these Directions, procurement organisations procuring tissues and/or cells for human application are subject to these Directions and not Directions 001/2007.
4. These Directions are made by the HTA.

Dated the 5th day of July, 2008.

Signed:



Adrian McNeil
Chief Executive
Human Tissue Authority

EXPLANATORY NOTE

Schedule 3 Paragraph 2 (4) (f) of the Human Tissue Act, 2004 as applied by Regulation 8 of the Human Tissue (Quality and Safety for Human Application) Regulations 2007, provides as follows:

“ 2 (4) It shall be a condition of a licence -

(f) that there shall be paid to the Authority at such times as may be specified in directions sums of such amount as may be so specified in respect of its costs in connection with superintending compliance with the terms of licences. “