

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

Directions on licence fee structure for the financial year 2009/10

Ref 001/2009

These Directions are

General Directions

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| Section of the Human Tissue Act 2004 (HT Act), and as applied by the Human Tissue (Quality and Safety for Human Application) Regulations 2007 (the Regulations) for the human application sector, providing for these Directions | Section 37 and Schedule 3 paragraph 2 (4) (f) of the HT Act and as applied by Regulation 8 of the Regulations for the human application sector |
| These Directions come into force on | 1 April 2009 |
| These Directions remain in force | Until revoked |

1. These Directions revoke Directions 001/2007 and 001/2008.
2. These Directions are made under Section 37 and Schedule 3 Paragraph 2 (4) (f) of the Human Tissue Act 2004 (HT Act) and for the human application sector, those sections 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 licence fee structure of the Human Tissue Authority (HTA) for the financial year 2009/10.
3. The HTA’s 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 and for the human application sector, those sections as applied by Regulation 8 of the Regulations.

2009/10 – Licence Fee Structure

| Sector | Fee for main site (£) | Fee for satellite (£) |
|--|-----------------------|-----------------------|
| Research | 6,000 | 900 |
| Public display | 3,750 | 560 |
| Public display - exhibiting less than 20 items of relevant material | 1,000 | 560 |
| Post mortem | 8,000 | 2,000 |
| Anatomy | 6,500 | 975 |
| Human application - establishments storing, procuring, testing, processing, distributing, importing or exporting tissue and/or cells for human application | 11,000 | 3,800 |
| Human application (skin and bone) - establishments storing, procuring, testing, processing, and/or distributing, skin and/or bone for human application (for autologous use only) | 5,500 | 3,800 |

| | | |
|--|-------|-------|
| Human application (Procurement) - Procurers of tissue and/or cells for human application where they are not licensed for processing or storage of tissue and/or cells for human application. | 6,000 | 3,800 |
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4. These Directions are made by the HTA.

Dated the 1st day of April, 2009.

Signed: 

Adrian McNeil
Chief Executive
Human Tissue Authority

EXPLANATORY NOTE

Schedule 3 Paragraph 2 (4) (f) of the Human Tissue Act, 2004 and as applied by Regulation 8 of the Human Tissue (Quality and Safety for Human Application) Regulations 2007 for the human application sector, 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. “