

## SCHEDULE 1

Regulation 4

### Licensing

#### **Licence applications: general**

1.—(1) A person required by regulation 5 to hold a licence must make an application to the Licensing Authority in the form specified from time to time by the Licensing Authority.

(2) A person applying for a licence under sub-paragraph (1) must provide to the Licensing Authority—

- (a) such of the information described in paragraph 2 as the Licensing Authority may from time to time specify necessary to determine the licence application;
- (b) upon request in writing, any other information which the Licensing Authority requires for the purpose of considering the licence application; and
- (c) the fee specified in paragraph 4.

(3) A person issued a licence under these Regulations (“the licensee”) must apply to the Licensing Authority if the licensee seeks a material change to the licence in respect of any matter dealt with by that licence.

#### **Commencement Information**

**II** Sch. 1 para. 1 in force at 6.2.2018, see **reg. 1**

#### **Licence applications: indicative list of information**

2. The information referred to in paragraph 1(2) is information relating to—

- (a) responsibilities and organisational arrangements for protection and safety;
- (b) staff competences, including information and training;
- (c) design features of the radiological installation and of radiation sources;
- (d) anticipated occupational and public exposures in normal operation;
- (e) safety assessment of the activities and the facility in order to—
  - (i) identify ways in which potential exposures or accidental and unintended medical exposures could occur;
  - (ii) estimate, to the extent practicable, the probabilities and magnitude of potential exposures;
  - (iii) assess the quality and extent of protection and safety provisions, including engineering features, as well as administrative procedures;
  - (iv) define the operational limits and conditions of operation;
- (f) emergency procedures;
- (g) maintenance, testing, inspection and servicing so as to ensure that the radiation source and the facility continue to meet the design requirements, operational limits and conditions of operation throughout their lifetime;
- (h) management of radioactive waste and arrangements for the disposal of such waste, in accordance with applicable regulatory requirements;
- (i) management of disused sources;
- (j) quality assurance.

**Changes to legislation:** There are currently no known outstanding effects for the The Ionising Radiation (Medical Exposure) Regulations 2017, SCHEDULE 1. (See end of Document for details)

**Commencement Information**

**I2** Sch. 1 para. 2 in force at 6.2.2018, see [reg. 1](#)

**Licence applications: urgent cases**

**3.** The licensing authority may, on a case by case basis, relax any of the requirements relating to the making of an application for a licence in respect of a proposed urgent medical radiological exposure.

**Commencement Information**

**I3** Sch. 1 para. 3 in force at 6.2.2018, see [reg. 1](#)

**Licence applications: employer fees**

**4.—(1)** The fee payable by a person described in column 1 of Table 1 in respect of an application type specified in column 2 of that table is the corresponding amount in column 3.

(2) No fee is payable where the amount specified in column 2 is “0”.

**Table 1**

<i>Licence type (1)</i>	<i>Application type (2)</i>	<i>Fee (£) (3)</i>
Employer	New	250
	Amendment of an existing licence	200
	Renewal of an existing licence	200
	Notification	0
Practitioner	New	0
	Amendment of an existing licence	0
	Renewal of an existing licence	0
	Particular patient request	0

**Commencement Information**

**I4** Sch. 1 para. 4 in force at 6.2.2018, see [reg. 1](#)

**Review**

**5.—(1)** A person who is aggrieved (“an aggrieved person”) by—

- (a) a decision of the Licensing Authority—
  - (i) refusing to issue a licence;
  - (ii) imposing a limit of time upon a licence; or

- (iii) revoking a licence; or
  - (b) the terms of any conditions attached to a licence by the Licensing Authority, may ask the Licensing Authority for a review.
- (2) Any aggrieved person seeking a review must—
- (a) within 28 days of the date that the person was notified of the decision, or the terms, which caused them to become an aggrieved person request the Licensing Authority to undertake a review described in paragraph (1) ; and
  - (b) must particularise in writing the reasons for seeking the review.
- (3) The Licensing Authority must undertake a review, and provide the results of that review in writing to the aggrieved person.

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**Commencement Information**

**I5** Sch. 1 para. 5 in force at 6.2.2018, see [reg. 1](#)

**Destination of fees**

6. A fee payable under these Regulations is payable to the Secretary of State.

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**Commencement Information**

**I6** Sch. 1 para. 6 in force at 6.2.2018, see [reg. 1](#)

**Changes to legislation:**

There are currently no known outstanding effects for the The Ionising Radiation (Medical Exposure) Regulations 2017, SCHEDULE 1.