

## SCHEDULE 2

Applications in respect of pharmaceutical lists and the procedures to be followed

### PART 4

Determination and deferral of applications

#### **Additional matters for consideration in relation to applications for inclusion in a pharmaceutical list**

**23.**—(1) In the case of a routine or excepted application by a person (A) for inclusion in a pharmaceutical list who is not already included in it, [<sup>F1</sup>NHS England] must, prior to determining the application—

- (a) check with the NHS BSA whether A, and if A is a body corporate whether any director or superintendent of A, has any record of, or is under investigation for, fraud;
  - (b) check with the Secretary of State whether the Secretary of State holds any information about A, and if A is a body corporate about any director or superintendent of A, that is relevant to its consideration of whether—
    - (i) the application should be refused or deferred under regulations 33 or 34, or
    - (ii) conditions should be imposed under regulation 35;
  - (c) take up references from, and check the references provided by, the referees whose details A is required to provide pursuant to paragraph 3(8).
- (2) In such a case, having considered whether—
- (a) the application should be refused or deferred under regulations 33 or 34; or
  - (b) conditions should be imposed under regulation 35,

if it is minded to impose conditions under regulation 35, it must notify A at least 7 days in advance of determining that it is to impose such conditions and consider any representations (which may be at an oral hearing) that A makes prior to the determination with regard to the notification.

#### **Textual Amendments**

- F1** Words in Regulations substituted (6.11.2023) by [The Health and Care Act 2022 \(Further Consequential Amendments\) \(No. 2\) Regulations 2023 \(S.I. 2023/1071\)](#), reg. 1(1), **Sch. para. 1**

**Changes to legislation:**

There are currently no known outstanding effects for the The National Health Service (Pharmaceutical and Local Pharmaceutical Services) Regulations 2013, Paragraph 23.