SCHEDULE 1

AMENDMENTS TO SUBORDINATE LEGISLATION

Plant Protection Products (Fees and Charges) Regulations 2011

15.—(1) Schedule 2 is amended as follows.

(2) In the Schedule heading, for "fee" substitute "fees and standalone MRL application fees".

(3) After the Schedule heading insert the paragraph heading "Fees for import tolerances".

(4) The existing content of the Schedule (after the Schedule heading) becomes paragraph 1.

(5) In that paragraph—

(a) in the first sentence, for "product-related applications" substitute "import tolerances";

(b) in the table, before item 1 insert—

"A1	Preliminary consideration of an application to determine whether the application can proceed further	229
A2	Co-ordination of applications	1,872"

(c) in the notes following the table—

(i) for note (1) substitute—

"(1) This category is mainly for active substances not currently approved in respect of the part of the United Kingdom to which the application relates. In certain cases it may also include active substances still being reviewed if toxicological endpoints have not yet been agreed and accepted in respect of that part of the United Kingdom.";

(ii) in note (2)—

(aa) for "plant protection products" substitute "active substances";

(bb) for "at a European level" substitute "and accepted in respect of the part of the United Kingdom to which the application relates";

(iii) in note (3)—

(aa) for "plant protection products" substitute "active substances";

(bb) for "at European level" substitute "and accepted in respect of the part of the United Kingdom to which the application relates";

(iv) after note (3) insert—

"Fees for multiple import tolerances for the same active substance are calculated on a modular basis with a charge applied for each crop."

(6) After that paragraph insert—

"**Fees for standalone MRL applications**

2. Fees for standalone MRL applications are in accordance with the following table.

Item	*Category*	*Fee (£)*
1	Preliminary consideration of an application to determine whether the application can proceed further	229
2	Co-ordination of applications	1,872

Item	*Category*	*Fee (£)*
3	Full human health description(1)	16,224
4	Metabolism and residues evaluation(2)	6,760
5	Residues evaluation(3)	2,028”

Notes

(1) This category is mainly for active substances not currently approved in respect of the part of the United Kingdom to which the application relates. In certain cases it may also include active substances still being reviewed if toxicological endpoints have not yet been agreed and accepted in respect of that part of the United Kingdom.

(2) This category is for active substances where toxicological endpoints have already been agreed and accepted in respect of the part of the United Kingdom to which the application relates but the residue definition has only been established for crop groups unrelated to the intended use.

(3) This category is for active substances where relevant toxicological endpoints and residue definition have already been agreed and accepted in respect of the part of the United Kingdom to which the application relates.

Fees for multiple standalone applications for the same active substance are calculated on a modular basis with a charge applied for each crop or combination of maximum residue levels.