SCHEDULE 8

EC DECLARATION OF CONFORMITY TO TYPE PROCEDURE
(CORRESPONDING TO ANNEX 5 OF THE DIRECTIVE)

PART I

QUALITY SYSTEM

1.—(1) The manufacturer shall make an application in writing for evaluation of his quality system to a notified body.

(2) The application shall be signed by or on behalf of the manufacturer and shall include—

(a) the information and undertakings which are required by paragraph 2 of Schedule 5 to be included in an application under paragraph 1 of that Schedule, other than the information required by paragraph 2(1)(e) of that Schedule so far as it relates to research and development and design;

(b) the technical documentation relating to the approved type and a copy of the EC type-examination certificate.

2.—(1) Application of the quality system must ensure that the products conform to the type described in the EC type-examination certificate.

(2) Subject to sub-paragraph (3) below, paragraphs 3(3) and (4)(a), (b), (d) and (e), 4 and 5 of Schedule 5 shall apply for the purposes of this Schedule as they apply for the purpoes of that Schedule, but as though any reference to paragraph 3(2) to (4) of that Schedule were a reference to sub-paragraph (1) of this paragraph and paragraphs 3(3) and (4)(a), (b), (d) and (e) of that Schedule.

(3) Paragraph 3(4)(b)(ii) of Schedule 5 shall not apply in respect of the desired quality of design.