S C H E D U L E S

SCHEDULE 24

Packaging information requirements

PART 1

Outer and immediate packaging

5. A statement of the active substances in the product, expressed qualitatively and quantitatively per dosage unit or according to the form of administration for a given volume or weight, using their common names.

Changes to legislation:
There are currently no known outstanding effects for the The Human Medicines Regulations 2012, Paragraph 5.