

LIST OF QUALITY GUIDELINES ACCEPTED FROM THE GUIDELINES FOR HUMAN USE⁷

DEVELOPMENT PHARMACEUTICS AND PROCESS VALIDATION

LIMITATIONS TO THE USE OF ETHYLENE OXIDE IN THE MANUFACTURE OF
MEDICINAL PRODUCTS

THE USE OF IONISING RADIATION IN THE MANUFACTURE OF MEDICINAL
PRODUCTS

CHEMISTRY OF ACTIVE SUBSTANCES

REQUIREMENTS IN RELATION TO ACTIVE SUBSTANCES

EUROPEAN DRUG MASTER FILE PROCEDURE FOR ACTIVE SUBSTANCES

PLASTIC PRIMARY PACKAGING MATERIALS

SPECIFICATIONS AND CONTROL TESTS ON THE FINISHED PRODUCT

STABILITY TESTING ON ACTIVE INGREDIENTS AND FINISHED PRODUCTS

QUALITY OF HERBAL REMEDIES

⁷ Volume 3 of The Rules Governing Medicinal Products in the European Union, 1998 edition.