ROYAL PHARMACEUTICAL SOCIETY

PROGRAMME

The 15th Joint Qualified Person Symposium – The QP in a new world

09.00am **Registration, tea and coffee**

Welcome & introduction

MHRA. Updates to GMP – including experiences with Annex 16 and the Falsified Medicines Directive requirements for serialisation

Discussion

Falsified Medicines Directive – Safety measures for manufacturers

Serialisation challenges - infrastructure

Serialisation and tamper evidence for operations

Questions/Feedback

12.45pm **Lunch**

QP Certification and Release - Transitioning from Traditional Dosage Forms to Advanced Therapeutic Medicinal Products

Clinical Trials Regulation update

MHRA enforcement group investigations and the illegal trading of controlled drugs

Discussion

16:00pm **Closing comments**

Please note the Programme may be subject to changes