(c) Each Party shall transmit to the other Party, without delay, information on any withdrawal of the certificate of a confirmed test facility if the facility has been found to be non-compliant with the principles of GLP.

6. (a) Each Party may request the other Party, by indicating in writing a reasoned doubt on whether a study was conducted in accordance with the principles of GLP, to conduct further inspections or study audits on a confirmed test facility, in accordance with the applicable laws, regulations and administrative provisions of that other Party.

(b) The requested Party shall inform the requesting Party of the results of the inspections or study audits, or provide an explanation of why such an inspection or study audit has not been carried out.

(c) The requesting Party shall not be obliged to accept the data generated by the test facility concerned from the date on which the request is made, until the results of the further inspection or study audit conducted by the Competent Authority of the requested Party have reconfirmed the compliance of the test facility with the principles of GLP.

(d) If, in exceptional cases, doubts persist, and the requesting Party can justify a specific concern, that Party may contest the compliance of the test facility concerned in accordance with Article 7 of this Protocol.

PART B

SECTION I: THE APPLICABLE LAWS, REGULATIONS AND ADMINISTRATIVE PROVISIONS STIPULATING THE COVERAGE OF CHEMICALS SUBJECT TO TESTING IN ACCORDANCE WITH THE PRINCIPLES OF GLP

THE UNITED KINGDOM	JAPAN
1. Medical Products: The Human Medicines Regulations 2012 (SI 2012/1916) and amendments thereto 2. Veterinary Medicinal Products: The Veterinary Medicines Regulations 2013 (SI 2013/2033) and amendments thereto 3. Plant Protection Products: (a) Plant Protection Products Regulations 2011 and amendments thereto (b) Plant Protection Products Regulations (Northern Ireland) 2011 and amendments thereto 4. Biocides: Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (EU Exit) Regulations 2019 and amendments thereto	1. Pharmaceuticals: (a) Law concerning Securing Quality, Efficacy and Safety of Products including Pharmaceuticals and Medical Devices (Law No. 145 of 1960) and amendments thereto (b) Regulation for Enforcement of the Law concerning Securing Quality, Efficacy and Safety of Products including Pharmaceuticals and Medical Devices (Ministerial Ordinance of Ministry of Health and Welfare No. 1 of 1961) and amendments thereto 2. Veterinary Drugs: (a) Law concerning Securing Quality, Efficacy and Safety of Products including Pharmaceuticals and Medical Devices (Law No. 145 of 1960) and amendments thereto