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Good pharmacovigilance practice - MHRA Inspectorate
Good pharmacovigilance practices (GVP) are a set of measures drawn up to facilitate the performance of pharmacovigilance in the European Union (EU). GVP apply to marketing-authorisation holders, the European Medicines Agency (EMA) and medicines regulatory authorities in EU Member States. They cover medicines authorised centrally via the Agency as well as medicines authorised at national level.

Good pharmacovigilance practices | European Medicines Agency
Good pharmacovigilance practice for medicines (GPvP) Guidance on what pharmacovigilance is and compliance issues from previous inspections. Published 18 December 2014

Good pharmacovigilance practice for medicines (GPvP) - GOV.UK
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Abbreviations and a Glossary of common terms. For more details on the contents, check out the MHRA Q&As on the Guide.

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MHRA's GXP data integrity guide published - MHRA Inspectorate
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Mhra (Author of Good Pharmacovigilance Practice Guide)
Guidance on good clinical practice has been produced by the International conference on harmonisation of technical requirements for registration of pharmaceuticals for human use (ICH). You can also...

Good clinical practice for clinical trials - GOV.UK
Good Manufacturing Practice. Regulations and Guidelines; Links. Regulatory Authorities; Associated Bodies and Organisations; Pharmacopoeias; Discussion Forum; COVID-19 Webinar - GMP Specific; Good Pharmacovigilance Practice. Regulations and Guidelines; PV Hot Topics; Links; Discussion Forum; COVID-19 Webinar - PV Specific; Medical Devices ...

MHRA: New rules for January 2021 | Research Quality ...
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