The road to medicinal products traceability

This poster stems from the GS1 Healthcare document “Regulatory roadmap: traceability of medicinal products”, which aims to provide detailed, yet non-exhaustive, directions to support regulators before, during and after the drafting of policies in regard to the traceability of pharmaceuticals.

The reason for its creation is to help address the growing number of requests from regulators for support in aligning with the global framework of medicines identification, marking, and traceability, and to provide them with a neutral, international non-for-profit standards organization involved with the support of the GS1 Healthcare Public Policy Work Team.

See the full document here: https://www.gs1.org/library/public-policy/GS1_Healthcare-ROAD-MAP_FINAL.pdf

About GS1 Healthcare

GS1 Healthcare is a global, voluntary user community bringing together all healthcare supply chain stakeholders, including manufacturers, distributors, healthcare providers, solution providers, regulatory bodies and industry associations. The mission of GS1 Healthcare is to lead the Healthcare sector to the successful development and implementation of global standards by bringing together experts in Healthcare to enhance patient safety and supply chain efficiencies.

www.gs1.org/healthcare

1 Analysis of the local healthcare market and definition of the regulatory objectives:
   • patient safety and/or
   • payment monitoring and/or
   • supply chain efficiency

2 Choice of the relevant traceability model
   Eg. Track and Trace; Point of Dispense verification system

3 Definition of the timeframe

4 Definition of the product scope

5 Define data carriers (barcodes), identification and data exchange using GS1 Global standards

© Copyright GS1 AISBL. All Rights Reserved.