PHARMACEUTICAL REGULATIONS TIMELINE

UZBEKISTAN

Government Authority/System Operator

State Tax Committee, Ministry of Health & CRPT TURON LLC

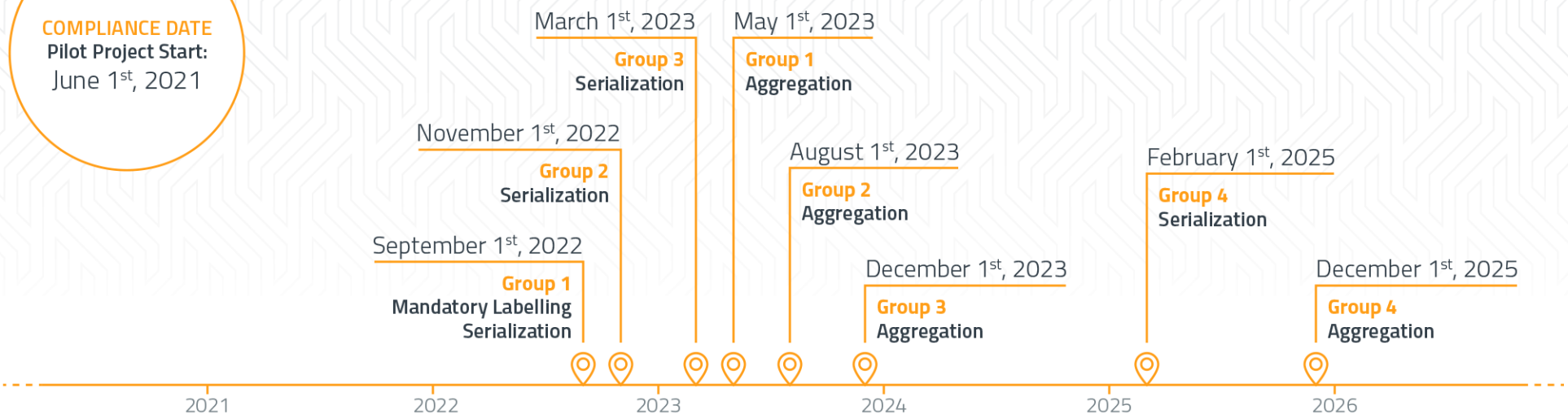
Track and Trace System

ASL BELGISI - National information system for digital labeling and traceability of products

Requirement Summary

- Serialization requirements mandate that the product must bear four data points in a GS1 Data Matrix code, which are as follows: GTIN, Serial Number, Crypto, and Crypto key. Stickers are permitted for serialization purposes.
- In addition, aggregation is mandatory, up to the case level, with pallet level being optional and limited to two levels within scope.
- Data reporting is an essential aspect of the process.

COMPLIANCE DATE
Pilot Project Start:
June 1st, 2021



Group 1: **Medicinal products with secondary** (outer) **packaging** (except for orphan)

Group 2: **Medicines with primary** (inner) **packaging** (provided there is no secondary (outer) packaging) (except for orphan)

Group 3: **Medicines and medical devices for orphan diseases** (according to the list approved by the Ministry of Health)
Medicines included in the register of medicines with foreign registrations, the results of which are recognized in Uzbekistan

Group 4: **Medical products according to the list determined by the Ministry of Health**

Source: https://crpt-turon.uz/m-products/pharma
NOTE: The regulatory timeline may be subject to change.