

HONEYWELL ACT | PHARMA FAQs



HOW DOES HONEYWELL HELP ADDRESS PHARMACEUTICAL TRACEABILITY REGULATIONS IN UAE?

Honeywell has developed Honeywell ACT | Pharma, a comprehensive track and trace SaaS solution, to help pharmaceutical supply chain stakeholders to better address the requirements from applicable pharmaceutical traceability regulations in the UAE.

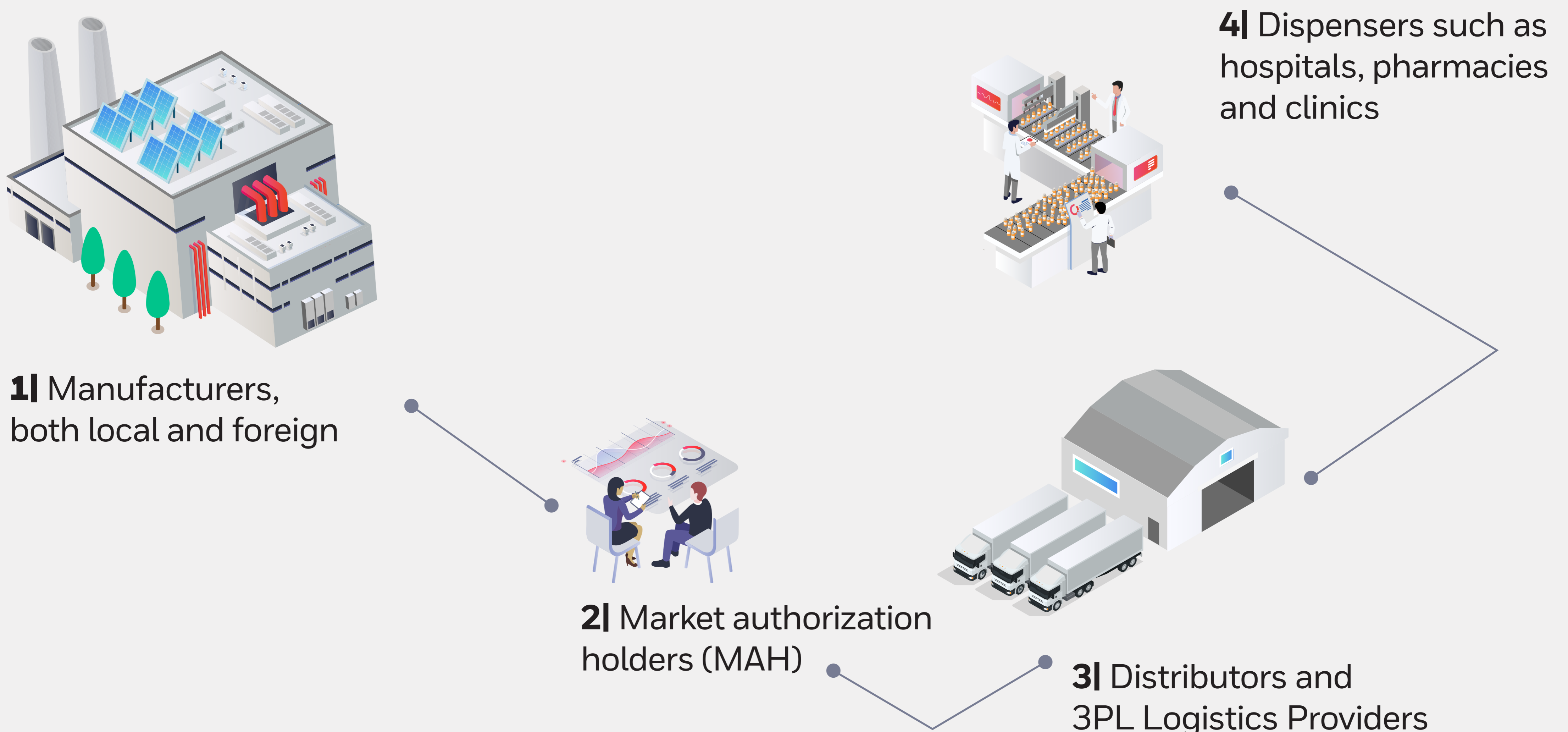


WHAT IS TATMEEN?

Tatmeen is a repository system initiated by the UAE Ministry of Health and Prevention (MoHAP) as part of the recently enacted pharmaceutical serialization and traceability regulations in UAE. These regulations aim at improving visibility, reliability, safety and security of the pharmaceutical supply chain, in order to reduce counterfeits and sub-standard medicines from circulation in UAE.

WHO ARE THE KEY PHARMACEUTICAL SUPPLY CHAIN STAKEHOLDERS UNDER UAE PHARMACEUTICAL TRACEABILITY REGULATIONS?

Tatmeen classifies the key stakeholders as:



WHAT IS REQUIRED FROM THESE STAKEHOLDERS BY THE UAE PHARMACEUTICAL TRACEABILITY REGULATIONS?



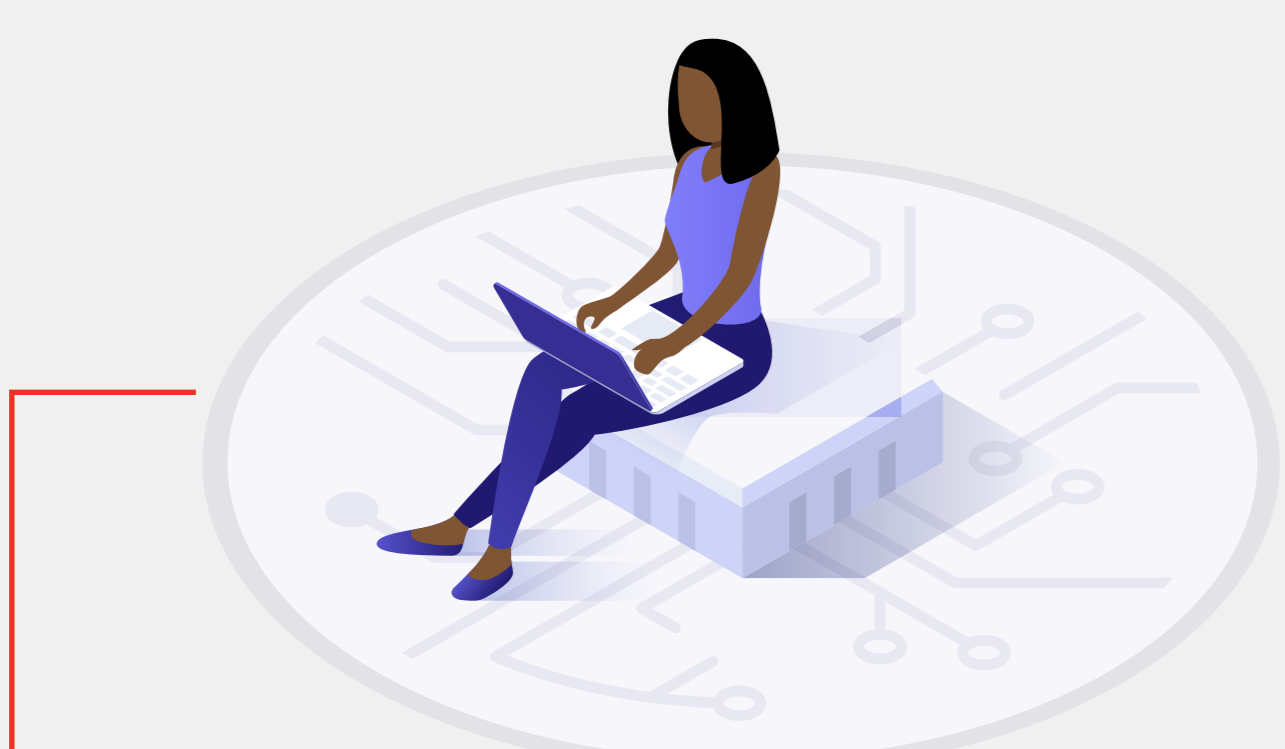
UAE classified key stakeholders in the pharma supply chain (manufacturers, market authorization holders, distributors, dispensers) will need to report certain supply chain events (packing, shipping, receiving and other events) as they happen to the Tatmeen repository system.

WHAT DOES THE HONEYWELL ACT | PHARMA SOLUTION INCLUDE?

Honeywell ACT Pharma includes a SaaS offering that can capture serialization data and product movements and status. In addition, the software enables enhanced supply chain tracking and visibility.

Honeywell ACT | Pharma provides 3 ways to integrate data into the Honeywell Track and Trace cloud:

MOBILE APPLICATION to capture supply chain events in the field



WEB PORTAL for configuration, audit trail and dashboards



REST API for integration with the system of record (e.g ERP, WMS, MES, MDM, etc..)

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