PROJECT FACTSHEET

Funded by the European Union

Strengthening of the Medicines and Medical Devices Agency of Moldova as regulatory agency in the field of medicines, medical devices and pharmaceutical activity

Period of implementation: 11.01.2017 - 10.03.2019

EaP countries:
Moldova

EU contribution: € 1 100 000

Implementing organisation(s):
STATE MEDICINES CONTROL AGENCY

Social media account links:

Project website: amed.md/en/content/twinning-0

Project description:
The overall objective of the EU-funded Twinning Project “Strengthening of the Medicines and Medical Devices Agency of the Republic of Moldova as regulatory agency in the field of medicines, medical devices and pharmaceutical activity” is full and correct implementation of the EU acquis in the area of medicinal products and medical devices and preparation of the Medicines and Medical Devices Agency of the Republic of Moldova for joining the EU regulatory agencies network as an equal partner.
The main purpose this EU-funded project is to strengthen the functioning of the MMDA with regards to medicinal products manufacturing, marketing, pharmacovigilance, distribution and pricing and medical devices in scope of market supervision, vigilance and registration as well as to clinical trials and pharmaceutical activity.
The completion of this EU-funded Project should deliver the MMDA with capacities at the same level as peer institutions in the EU Member States and should allow patients to benefit from safer, better quality and more effective medicinal products and medical devices.

Expected results:
- Approximation of the legal framework and strengthening of MMDA’s institutional and organisational capacities.
- Strengthening of MMDA’s regulatory functioning with respect to medicinal products.
- Strengthening of MMDA’s regulatory functioning with respect to medical devices.
- Improvement of rational use of medicines and medical devices in Moldova.
- Transition of MMDA to full compliance with the EU requirements.