Digitalna transformacija regulatornih poslova za lekove

Razvoj RIMS registra humanih lekova
i dalja harmonizacija i standardizacija u skladu sa EU

03.04.2025. godine
PKS, Resavska 13-15, sala 1

8:30 – 9:00 **Registracija učesnika**

9:00 – 10:00 **Razvoj RIMS registra humanih lekova**
Marko Erić, Agencija za lekove i medicinska sredstva Srbije

10:00 – 10:20 **Pauza**

10:20 – 10:50 **Praksa unosa podataka o lekovima sa primerima**
Tanja Nađić, Agencija za lekove i medicinska sredstva Srbije
Јелена Терзић, Agencija za lekove i medicinska sredstva Srbije

10:50 – 11:20 **Prezentacija softverskog rešenja**
Zoran Dobrosavljević, Abstract

11:20 – 12:00 **Diskusija**

Korisni akronimi

RIMS	Regulatory Information Management System
FHIR	Fast Healthcare Interoperability Resources
ISO	International Organization for Standardization
IDMP	Identification of Medicinal Products
SPOR	Data management services for substances, products, organisations and referentials
SMS	Substance Management Services
PMS	Product Management Services
OMS	Organisation Management Services
ORG ID	SPOR OMS Organisation identity
LOC ID	SPOR OMS Location Identity
RMS	Referentials Management Services
PLM	Product Lifecycle Management
ePI	Electronic Product Information
eAF	Electronic Application Forms