Proposal for possible working groups within the working structure of the Euripid Stakeholder Dialogue Platform

Working Groups

Transparency Working Group

The Transparency Working Group aims at investigating the possibilities of increasing transparency of pricing of medicinal products via the EURIPID Collaboration and website.

Task	Method	Expected outcome/deliverable
Enhancing the access to the EURIPID database	To investigate if there is an interest among the stakeholders to get access to the EURIPID database.	Yes or No answer.
	If there is an interest then to investigate what kind of legal, organisation and financial changes would be necessary to be implemented compared to the current conditions.	A proposal to the Board of Participants of the EURIPID Collaboration on the legal, organisational and financial issues.
Enhancing pricing transparency	Investigating the options to share "real prices" of medicinal products between the Member States of the EU.	Feasibility study with list of pre-requisites.

EPR Monitoring Working Group

The EPR Monitoring Working Group aims at developing methodology for the monitoring of harmonisation of EPR activities of the EU countries in line with the recommendations of the EURIPID Guidance Document on External Reference Pricing.

Task	Method	Expected outcome/deliverable
ERP activity monitoring	To develop a methodology of monitoring the EPR activity and	ERP monitoring tool

	the implementation of the recommendations of the EURIPID ERP Guidance Document in the EU Member States.	
Update of the recommendations of the ERP Guidance Document (if necessary)	Revision of the recommendations of the ERP Guidance Document based on the experiences.	Updated ERP Guidance Document.

Patients' Access Working Group

The Patients' Access Working Group aims at investigating the possibilities of developing methodology to measure patients' access to medicines within the EU by using the data and features of the EURIPID database and website.

Task	**Method**	**Expected outcome/deliverable**
Measuring patients' access to medicines in the EU	Developing a methodology to measure patients' access	Methodology description
Quality improvement of sales volume information	To investigate the possibilities of the inclusion of volume information from the databases established under the Falsified medicines directive	Feasibility study.