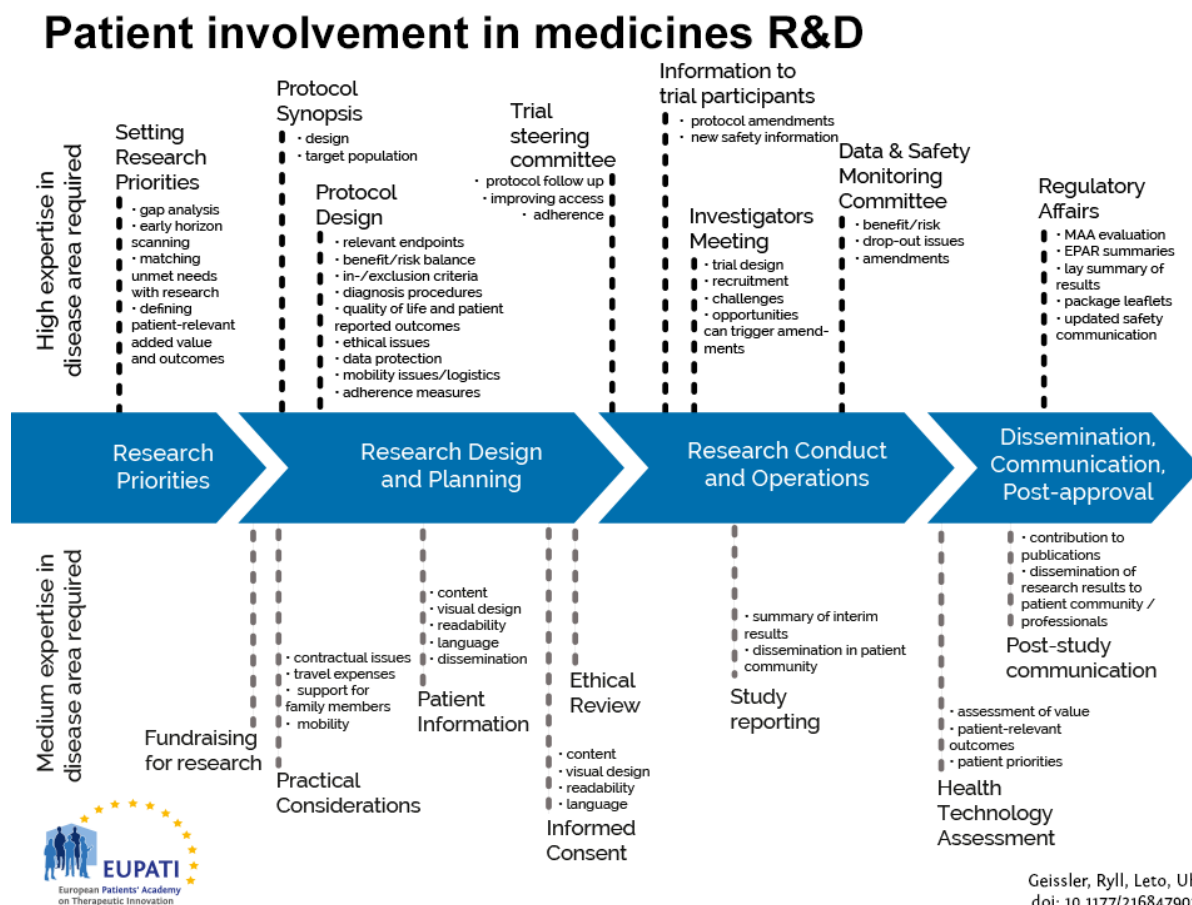


# Mini-course – Information & consent in trials

## Introduction

This EUPATI Mini-course starter kit is designed for patient involvement in product information, informed consent and patient information to trial participants.

EUPATI Mini-course starter kits have been derived from content found in the EUPATI toolbox and EUPATI Patient Expert Training Course. The starter kits are thought to address roles that patients play in medicines development for example those shown in the figure below.



Patients can be involved across the process of medicines

R&D. This diagram created by Geissler, Ryll, Leto, and Uhlenhopp identifies some existing areas in which patients are involved in the process. It distinguishes between the level of expertise in a disease area that is required and the different areas where involvement can take place.

The starter kits provide you with links to relevant background reading in the toolbox and associated PowerPoint slide decks and media in order to prepare a single or multi-day training on the subject. Each of the starter kits contains a selection of PPT slides which you may use to educate patients/advocates about the “basics” in that area, e.g. in a two-hour to one-day seminar.

The starter kits are based on existing content from the EUPATI Toolbox, plus additional links to add-on Toolbox material. None of the “starter kits” are “ready-made course” modules – they are a ready-to-reuse resource for an experienced trainer to prepare and execute a course. You will need to edit them and put them into context.

Before you begin please download and review the ‘Manual for Trainers’.

- **Presentation: Manual for Trainers**

Size: 722,143 bytes, Format: .pptx

A manual for trainers describing how to use the EUPATI mini-course starter kits to create trainings on patient involvement.

**Product information, informed**

# consent and patient information to trial participants

This starter kit provides background reading, slides, a video, and quizzes to create training for patients who intend to become involved in product information, informed consent and patient information to trial participants.

## Core reading

Risk communication in medicines

Information on medicinal products

Participants rights responsibilities organisations

Recording and reporting of clinical trial results

Clinical study results publication and application

## Optional

There is no optional content in the setting research priorities starter kit.

## Presentations

- Risk-Communication-in-Medicines

Size: 495,341 bytes, Format: .pptx

A presentation describing risk communication process in medicines, which can be adapted for own use.

- Presentation: Information on Medicinal Products

Size: 547,126 bytes, Format: .pptx

A presentation describing the information in medicinal

products, which can be adapted for own use.

- **Presentation: Participants' Rights, Responsibilities and Role of Patient Organisations**  
Size: 387,072 bytes, Format: .pptx  
A presentation describing the clinical trial participants' rights, responsibilities and role of patient organisations, which can be adapted for own use.
- **Presentation: Recording and Reporting Clinical Trial Results**  
Size: 372,913 bytes, Format: .pptx  
A presentation describing the recording and reporting of clinical trial results, which can be adapted for own use.
- **Presentation: Clinical Study Results Publication**  
Size: 449,072 bytes, Format: .pptx  
A presentation describing the clinical study results publication, which can be adapted for own use.

## Quizzes

Quizzes will be provided for each presentation in the future on:

Risk communication in medicines

Information on medicinal products

Participants rights responsibilities organisations

Recording and reporting of clinical trial results

Clinical study results publication and application

## Videos

Informed consent for vulnerable populations (Webinar)

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