

Objectives

Training session for patients and consumers involved in EMA activities, 25 November 2014

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General objectives:

To gain an understanding of the centralised procedure for how medicines are evaluated in Europe. To see where patients are consulted in these process and to learn more about where you can also be involved.

Specific objectives:

- Distinguish between Scientific Advice meetings and Scientific Advisory Groups
 - Understand the role of patients in these processes
- Identify information destined to the public that is reviewed by patients
 - How to review and impact of this review
- Navigate the website improve your understanding of how to find information on a medicine? Regulatory decision?
- Summarise the purpose of Pharmacovigilance and the role of patients
- Understand logistical aspects such as using EudraLink and how to declare your interests when attending a meeting at EMA