DSFE 2022 annual meeting: Post-authorization safety studies (PASS): can routinely collected data be used to identify side effects of drugs?

Date: 1 November 2022.

Location: Room 5.S.B.C, Danish Cancer Society Research Center, Strandboulevarden 49, DK-2100 Copenhagen.

Sign-up deadline: 10 October 2022.

Abstract submission: Abstracts should be ≤300 words and include the sections: introduction, methods, results, and conclusion. Abstracts should be written in English and may include one figure (the figure legend is not included in the word count). Abstracts should be sent to info@dsfe.dk before 26 September 2022. Three abstracts will be chosen for a 10-minute oral presentation. Submitters will receive response on or before 6 October 2022.

Time	Topic	Presenter
09:30	Arrival	
10:30	Opening remarks	Mette Bliddal, DSFE Chair
10:45	Purpose and impact of PASS - Goals and data collection approaches - Types of routinely collected data across countries - Regulatory role and impact	Susana Perez-Gutthann
11:30	Development of PASS - Methodological considerations - Possibilities and concerns using routinely collecting data - Examples	Espen Jimenez Solem
12:15	Lunch	
13:15	Oral abstract presentations	
13:45	General assembly	
14:15	Hypothesis-free screening of adverse drug reaction in big data - Advantages and disadvantages - Supplementing traditional side-effect surveillance - Examples	Jesper Hallas and Lars Christian Lund
15:00	Coffee	
15:30	Around the table discussion	
16:00	Closing remarks	