



8 December 2015 EMA/334164/2015 Draft for public consultation

Guideline on good pharmacovigilance practices (GVP)

Annex II – Templates: Communication Plan for Direct Healthcare Professional Communication (CP DHPC)

Draft finalised by the Agency in collaboration with Member States	17 November 2015
Draft agreed by European Risk Management Strategy Facilitation Group (ERMS FG)	24 November 2015
Draft adopted by Executive Director	8 December 2015
Start of public consultation	15 December 2015
End of consultation (deadline for comments)	29 February 2016
Date for coming into effect	Q2 2016

Comments should be provided using this <u>template</u>. The completed comments form should be sent to <u>gvp@ema.europa.eu</u>.



DHPC COMMUNICATION PLAN			
Medicinal product(s)/active substance(s)			
Marketing authorisation holder(s)	In cases where the DHPC concerns several marketing authorisation holders of the same active substance or is part of a class review, it is strongly encouraged that a single consistent message is sent to healthcare professionals in each EU Member State.		
	All concerned marketing authorisation holders in each Member State are strongly encouraged to collaborate, so that a single DHPC is prepared and circulated in each Member State. The letter circulated in each Member State should cover all active substance-containing products authorised in that Member State.		
	It is encouraged that the originator marketing authorisation holder (where available) in each Member State acts as the contact point for the national competent authority, on behalf of the other concerned marketing authorisation holders in the same Member State. If no originator product is marketed in the Member State, it is encouraged that the concerned generic companies acts as contact point for the competent authority.		
Safety concern and purpose of the communication	Consider using the title of the DHPC to describe the safety concern		
DHPC recipients	List all recipients of the DHPC in this section, e.g. general practitioners, specialists, pharmacists, nurses, professional societies, national associations.		
Member States where the DHPC will be distributed			
Timetable Delete steps	which are not applicable Date		
DHPC and communication plan (in English) agreed by PRAC			
DHPC and communication plan (in English) agreed by CHMP/CMDh			
Cubarication of translated DUDCs to the national compatent			

Timetable Delete steps which are not applicable	Date
DHPC and communication plan (in English) agreed by PRAC	
DHPC and communication plan (in English) agreed by CHMP/CMDh	
Submission of translated DHPCs to the national competent authorities for review	
Agreement of translations by national competent authorities	
Dissemination of DHPC	