

# Quality Expert Working Group: Subgroups

## VICH GL60: Good Manufacturing Practice for Active Pharmaceutical Ingredients Used in Veterinary Medicinal Products:

This document (Guide) is to provide guidance regarding GMP for the manufacturing of active pharmaceutical ingredients (APIs) used in Veterinary Medicinal Products under an appropriate system for managing quality. It is also intended to help ensure that APIs meet the requirements for quality and purity that they purport or are represented to possess.

### VICH GL60 Subgroup Members

**Topic Leader: Mai Huynh (US FDA)**



M. Huynh  
(US FDA)



M. Kerrigan  
(US FDA)



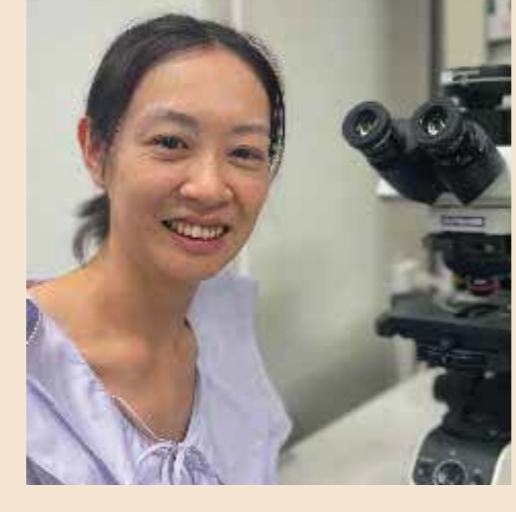
H. Fournel  
(EU AnimalhealthEurope)

J. Todd  
(UK VMD)

G. Edmunds  
(APVMA)



G. Verdier  
(EU Anses)



Y. Hosoda  
(JMAFF)



M. Folger  
(EU AnimalhealthEurope)

N. Henry  
(Health Canada)

B. Pies  
(US AHI)

C. Doyle  
(US AHI)

O. Moriyama  
(JVPA)

L. Labelle  
(CAHI)

I. Jarvis  
(CAHI)

## VICH GL61: Pharmaceutical Development:

This guideline describes the suggested contents for the Pharmaceutical Development which provides an opportunity to present the knowledge gained through the application of scientific approaches and quality risk management to the development of a product and its manufacturing processes. Adopting the principles in this guideline is optional for manufacturers.

### VICH GL61 Subgroup Members

**Topic Leader: Mai Huynh (US FDA)**



M. Huynh  
(US FDA)



S. Bowman  
(US FDA)



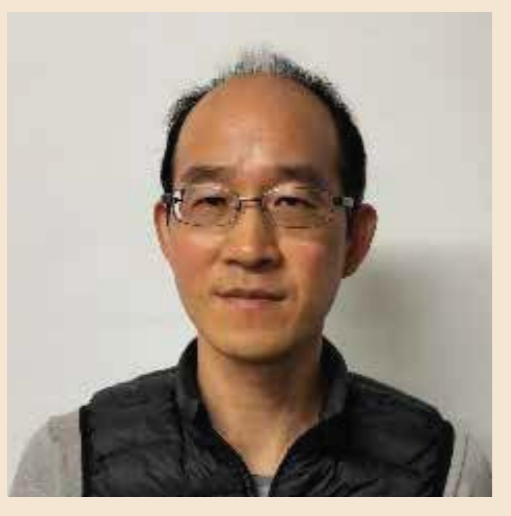
C. Janich  
(EU BVL)



P. Macours  
(EU Anses)



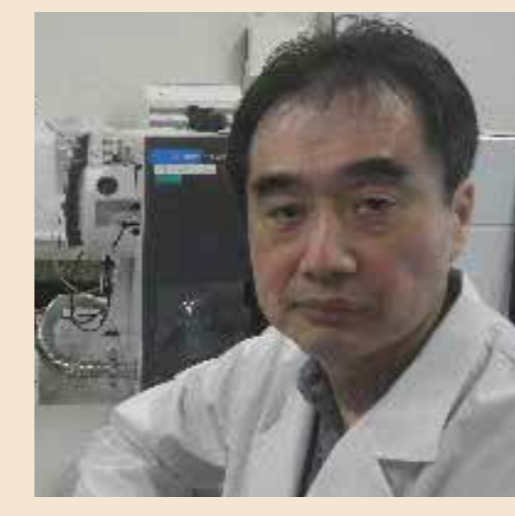
M. Ohashi  
(JVPA)



R. Teng  
(APVMA)



D. Blum  
(US AHI)



K. Eguchi  
(JMAFF)



J. Benoliel  
(Canada VDD)



G. Clarke  
(UK VMD)



M. Folger  
(EU AnimalhealthEurope)

V. Neron De Surgy  
(EU AnimalhealthEurope)

D. Katerere  
(South Africa)



Amsterdam  
November 2024