

Summit compounding overview day 1 “what have we learned sofar“

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The Hague, Friday, September 24

Scope and definitions:

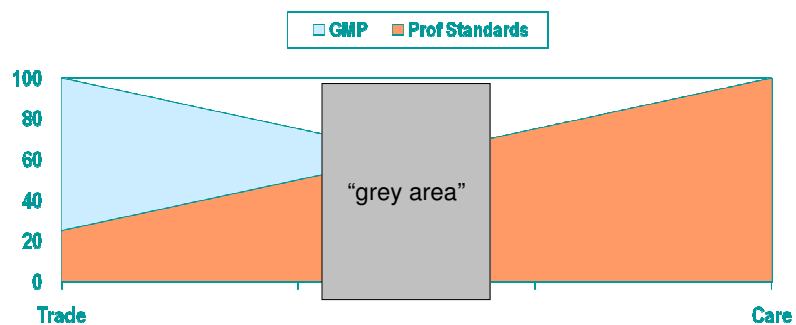
“why are we on earth as pharmacists?”

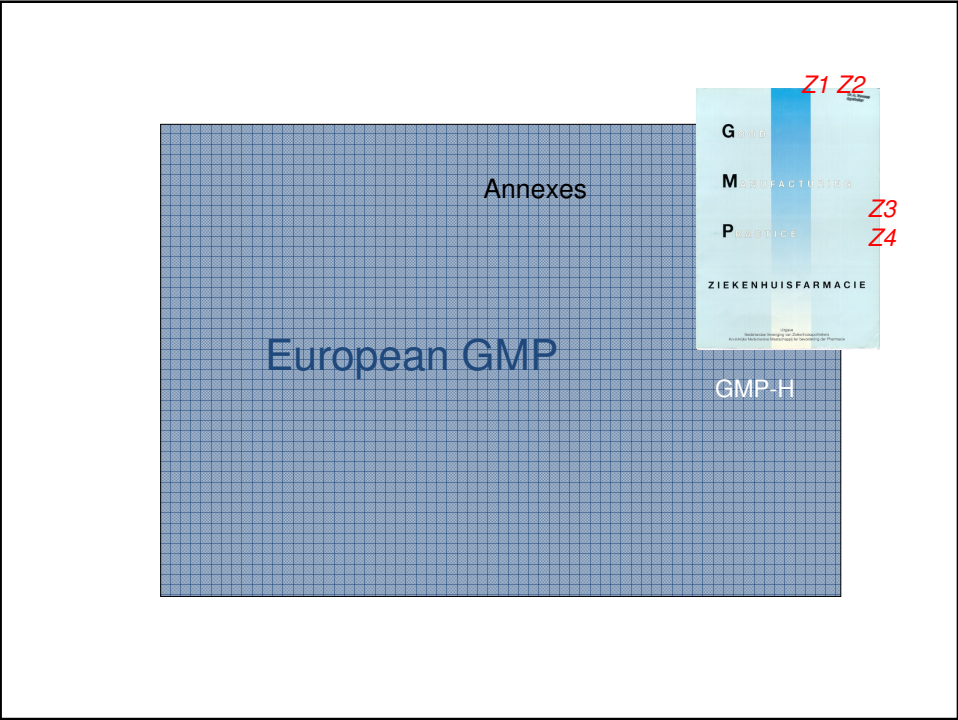
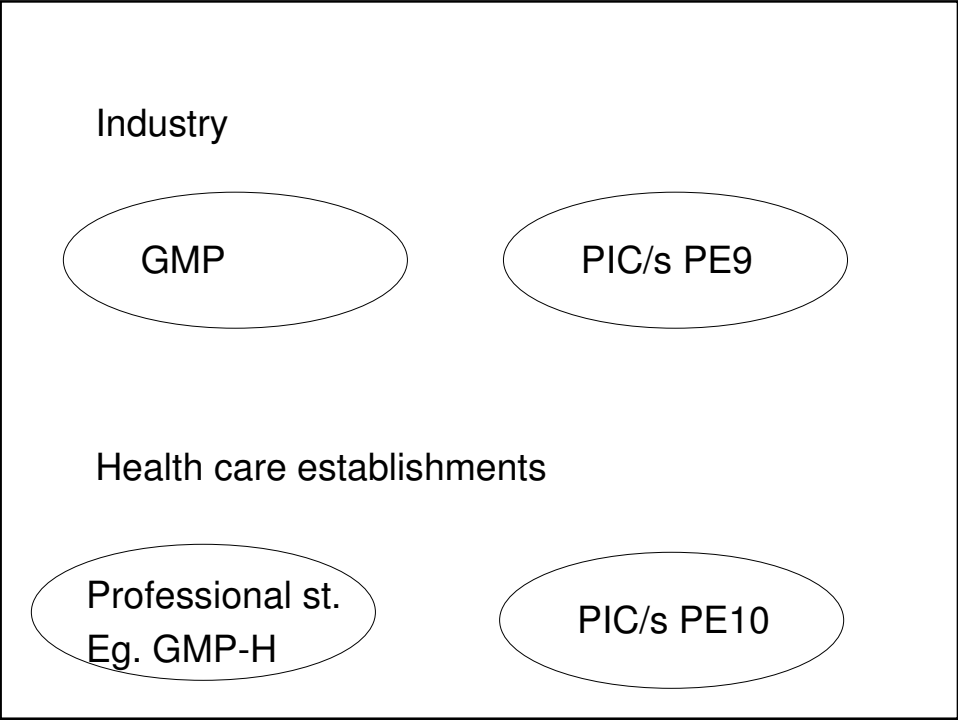
- Why are not all medicines available?
 - Orphan drugs
 - Drug shortages
 - Neglected patients (old, young, bad swallowers)
- There is simply no other drug or the preparation is better than a licensed medicine
- Risk of a possibly defective design or preparation process
- Balancing the risks; but ask for enough money!
- Added value

Quality and safety: how to reach a quality standard in European hospital pharmacies

- Survey
- Harmonization of standards
- Legal and professional standards

Care versus Trade





Site master file:

- General information: WHAT
- Personnel: WHO
- Premises and equipment: WHERE
- Documentation: HOW

- Basic document
- Design step by step

Business plan:

- What to consider: trends in therapy, customers, costs, legislation, competency.
- Proposal (eg facilities, equipment, personnel, Q system, validation, logistics and Procurements)
- Consider “what if”: risk management

Premises and equipment:

- How do you validate and when do you start with validation
- V- model
- VMP Documentation
- Design principles and fundamentals
- Example of lay outs
- Discussion! (legislation, cost and risks)

TPN case

- State of the art facility
- Still an incident might occur
- Professional Standards!
- Transparency!

The group work

- You all work hard but does it work!
- Are you all organized (president and secretary)
- Presentation!

Preview on day 2

- We start at 8:30(!) with “Product design”
- Continue with your topic and add the new subjects
- Keep in mind that we expect also, a **poster** from every group for the next **EAHP congress**, March 2011 (deadline October, 15)
 - Sign out (Points!) and
 - **Now: cocktail!**