5th PCNE Working Conference 21-24 February 2007 **Overcoming the barriers in pharmaceutical care research**

Workshop 4 Interprofessional collaboration in pharmaceutical care research

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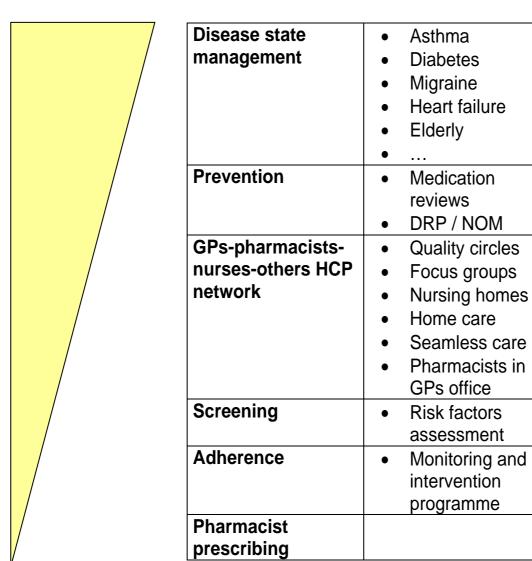
Overcoming the barriers to interprofessional collaboration in pharmaceutical care research

Aims of the workshop 4...

Through debate, discussion and risk analysis of three case studies:

 prepare a PCNE guide for risk management of interprofessional issues that can arise in pharmaceutical care research

Mapping of the actual interprofessional pharmaceutical care research



A Common Risk Management Process



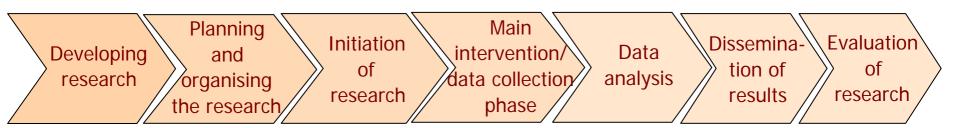
Result of the SWOT analysis on interprofessional collaboration in pharmaceutical care research

From a pharmacist-centred research to

a primary care-centred research ->

- Build a win-win situation for the primary care team
- Research to demonstrate the added value of the pharmacist for the primary care team and vice-versa
- Reinforce new collaborative models, particularly in integrated care and adherence

Description of the global Pharmaceutical care research process :



Risk management grid

| Impact | Risk Management Actions | | | | | | | |
|------------------|--|------------------------------------|--------------------------------------|--|--|--|--|--|
| 3 Significant | Considerable management required | Must manage and monitor risks | Extensive management essential | | | | | |
| Moderate 2 | Risks may be worth accepting with monitoring | Management effort worthwhile | Management 6 effort required | | | | | |
| Minor 1 | Accept risks | Accept, but monitor risks | Manage and monitor risks | | | | | |
| | Low 1 | Medium 2 Likelihood | High 3 | | | | | |

Adapted and modified from http://www.tbs-sct.gc.ca/pubs_pol/dcgpubs/RiskManagement/rmf-cgr01-1_e.asp#Key%20Concepts

| PROCESS | Planning and Initiation of research organising the research phase Planning and Initiation of research organism the research organism the research phase Planning Alam Initiation of Init | | | | | | | | | |
|--|--|---|--|------------------------------|---------------------------|--|---|--|--|--|
| | research | Conflicts of | patient | delivering the | recommenda- | transfer into | | | | |
| | stakeholders | interest | enrollment | intervention | tions | practice | Timely delivery | | | |
| | (3,3 = 9) selection of | (3,3 = 9) | (3,3 = 9) | (3,3 = 9) | (2,2=4) | (3,3 = 9) | (1,1 = 1) | | | |
| OJECT MANAGEMENT ACTIVITIES (impact of risk; likelihood of risk) | design, methods and outcomes (3.3 = 9) | communication strategy (3,3 = 9) | baseline data collection (3,1 = 3) | timely delivery (2,3 = 6) | perspectives (2,2 = 4) | lobbying (3,2 = 6) | impact factor | | | |
| | research team members (2,2 = 4) | non financial resources, incl. goodwill (3,2 = 6) | sites recruitment (3,1 = 3) | study monitoring | statistics | public com- munication (3,1 = 3) | publications | | | |
| act of ris | strategic development (3,1 = 3) | successful pilot / feasibility study (3,2 = 6) | training (3,1 = 3) | data collection | prospective modeling | scientific publication | fundings | | | |
| TIVITIES (imp | formulating research question (3,1 = 3) | design of study protocol / materials (2,2 = 4) | | | results presentation | conferences | other performance indicators | | | |
| ENT ACT | research funding (2,1 = 2) | environmental scan (2,1 = 2) | | | limitations | education | successful transfer into practice | | | |
| EM | literature phase | budget | | | | | PhD | | | |
| PROJECT MANAG | concept development | ethical approval criteria for study sites inclusion Development of education database timeline / milestones risk assessment | | | | | | | | |
| | | milestones risk assessment statistical design | | | | | | | | |