

Pharmaceutical care Hand-out

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Online lectures

Topic	Online Lecture	Time
Introduction to Pharma.Care Management	https://youtu.be/vYGMHLJE_qw	12
Phases and steps Pharmaceutical Care	https://youtu.be/LyBUaVPNnO8	13
Medicines Optimisation Principles	https://youtu.be/q_MNFXokloc	15
Drug Related Problems	https://youtu.be/F0Mh1JfWmks	19
PCNE system	https://youtu.be/OKXtH85zOD4	21
Assessment of Appropriate prescribing	https://youtu.be/k6_3r1NM3wE	15
Beers, STOP/START, PRISCUS	https://youtu.be/fTtQLyPBR5E	15
1 st Part of Side effects	https://youtu.be/niAlCzwyHLA	9
More about adverse events	https://youtu.be/8s_8C-dYF7I	24
Pharmacovigilance	https://youtu.be/9Nv6s0n91vE	21
Adherence Part 1	https://youtu.be/l6U3h0Xx5c4	25
Adherence Part 2	https://youtu.be/7PhNAFPAEG8	21
Believes about medications	https://youtu.be/NWM7gEXex3A	16
Quality of life	https://youtu.be/G3-tyiPPI9M	15
D-D interaction	https://youtu.be/HXKzxx5UgBs	24



Second: Systems Medication (Drug) Related Problems

Be aware you should know:

هـ تـكـيـفـ هـنـ بـلـدـ الـكـلـيـلـ (ـعـتـلـعـكـافـعـاـ فـيـ خـفـقـنـسـ الـبـلـدـ)
هـ تـكـيـفـ هـنـ بـلـدـ الـكـلـيـلـ (ـعـتـلـعـكـافـعـاـ فـيـ خـفـقـنـسـ الـبـلـدـ)
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1) It is not simple the Adverse drug reactions, Drug-related problems (DRP) are events or circumstances involving drug therapy that actually or potentially interfere with desired health outcomes.

2) DRP and its classifications keep on consciously updated, and they may be varied:

Change over years, time / Based on country and area / Based on Patient criteria (Inpatient vs Outpatients, DRP Classification) حسب المرض أو بحسب المرض أو حسب المرض أو حسب المرض
Chronic vs Acute / based on purpose (for description, for management, for statistic, for economy ...etc)

3) Here we will have a quick general overview of systems in USA, Spain, Netherlands, Sweden and Jordan.

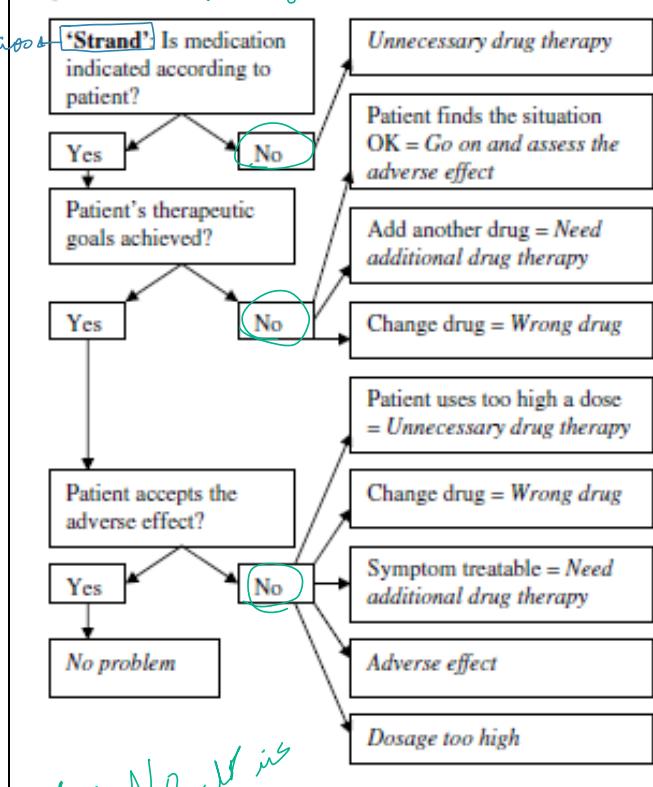
PLUS, the European System

النظام الأوروبي لـ DRP
نـسـمـنـ الـنـكـرـ لـ DRP

1) Cipolle/Morley/Strand classification

أـنـهـ مـنـ الـأـنـوـيـنـ وـعـدـ

patient ↳ drug ↳ Strand system



Community pharmacies in the US to evaluate pharmacists' activities in their daily provision of pharmaceutical care. This Classification can only be employed when the event has already been experienced by the patient.

DRPs were classified as follows: (core)

- Unnecessary therapy (Indication Or dose)
- Need for additional therapy
- Wrong drug
- Dosage is too low (refer to i)
- Need additional drug therapy
- Adverse drug reaction
- Dose is too high
- Adherence problem

وـ قـدـمـهـ مـنـ الـأـنـوـيـنـ وـعـدـ
أـنـهـ مـنـ الـأـنـوـيـنـ وـعـدـ

إـنـهـ طـوـرـواـ مـفـهـومـاـ يـقـرـبـ مـنـ الـفـكـرـ
الـأـسـاسـيـةـ /ـ الـجـوـهـرـيـةـ

2) American Society of Hospital Pharmacists (ASHP) classification

In this classification, the DRPs were classified as follows:

- i. Medication with no indication
- ii. Condition for which no drug is prescribed
- iii. Medication prescribed inappropriately for a particular condition
- iv. Inappropriate dose, dosage form, schedule, route of administration, or method of administration
- v. Therapeutic duplication
- vi. Prescribing of medication to which the patient is **allergic**
- vii. Actual and potential adverse drug events
- viii. Actual and potential drug-drug, drug-disease, drug-nutrient, and drug-laboratory test interactions that are clinically significant
- ix. Interference with medical therapy by **social or recreational drug use**
- x. Failure to receive the full benefit of prescribed therapy
- xi. Problems are arising from the **financial impact of therapy**
- xii. **Lack of understanding** of the medication
- xiii. Failure of the patient to adhere to the regimen.

العنودات بالخط الفائق
فردو (العنودات الفائق)
لحد الارض
System

3) Granada consensus

In 1998, a group of Spanish experts reached a consensus on the definition and analysis of DRPs. In this classification the DRPs were classified as follows:

TABLE I – Classification of 'Drug Related Problems' (DRP)

Necessity	Problem	three domains (categories) or non quantitative	
		① قسم	② قسم
① الاجراء (Necessity)	DRP 1: The patient suffers from a health problem as a result of not taking the medicine that he needs.		
② الكفاءة (Effectiveness)	DRP 2: The patient suffers from a health problem as a result of taking a medicine that he does not need.		
③ الامان (Safety)	DRP 3: The patient has a health problem resulting from a non-quantitative ineffectiveness of a medicine. DRP 4: The patient has a health problem resulting from a quantitative ineffectiveness of a medicine. DRP 5: The patient suffers from a health problem as a consequence of a <u>non-quantitative safety problem</u> of a medicine. مثلاً مرضيPenicillin جاف	non quantitative	non quantitative
	DRP 6: The patient suffers from a health problem as a consequence of a <u>quantitative safety problem</u> of a medicine. مثلاً مرضيPenicillin جاف	(2+) متعدد وغير وغير وغير	

Major issues for the drug:

المسائل / القضايا الرئيسية الخاصة
بالدواء

و غالباً تقصد بها في المصطلح:

الأثار الجانبية المهمة

مشكلات السلامة

التدخلات الدوائية

مشكلات الامتصاص أو التوازن الحيواني

الشممية أو ضيق المجال العلاجي

لكن تترجم حرفة فقط:

مشكلات رئيسية =
Major issues =
بالنسبة للدواء =
Major issues for the drug

For example, a patient went to the doctor because of temporary back pain. The doctor prescribed ibuprofen, intended to be taken until the back pain improves. However, the patient keeps renewing the prescription along with other medications he takes for his chronic diseases.

4) Hanlon approach (University of Pittsburgh)

It is not just a system for classifying DRPs, but also includes MAI.

Hanlon et al. have developed a method for assessing the appropriateness of medication based on the medication appropriateness index (MAI).

In this classification, the DRPs were classified as follows:

- i. indications
- ii. effectiveness
- iii. Correct directions
- iv. Practical directions
- v. Drug – Drug interaction
- vi. Drug disease-condition interaction

الاتجاهات
الفعالية



- vii. Duplication
- viii. Duration of therapy
- ix. Economic justification
- x. Improper drug selection

5) National Coordinating Council \rightarrow wrong *الخاطئ* *الغير صحيح*

for Medication Error Reporting and Prevention (NCC-MERP) taxonomy of medication errors.

In this classification, the DREs were classified as follows:

- i. The medication is in control of the health care professional, patient, or consumer.
- ii. Dose omission
- iii. Improper dose
- iv. Wrong strength/concentration
- v. Wrong drug
- vi. Wrong dosage form
- vii. Wrong technique (includes inappropriate **crushing** of tablets) *مثلاً أخذ الصوب* \rightarrow *التي ترمي به قبض المساند* (sublingual)
- viii. Wrong route of administration *أدوية غير ملائمة لتناولها* *تناولها خاطئاً وبلعها* (swallow)
- ix. Wrong date (probably relating to administration) *في الماء*
- x. Wrong duration *أخطاء في مدة الدورة*
- xi. Wrong time *أخطاء في موعد الدورة*
- xii. Wrong patient**
- xiii. **Monitoring** error (includes contraindicated drugs)
- xiv. **Deteriorated** drug error (dispensing drug that has expired)
- xv. Other.

6) SHB-SEP classification

The Health Base Foundation developed this system in The Netherlands for use in **pharmacy software's** based on the medical **Subjective/ Objective/Evaluation/Plan structure**; however, the S and O codes have been combined into one problem description.

The main problem categories comprise both a patient- and pharmacy-oriented perspective.

The system is still being revised regularly, but each updated version is not sequentially numbered to facilitate differentiation from previous versions.

- i. Patient initiative doubts or insufficient understanding (also second opinion)
- ii. Question about drug use (dosage/advice/way of use)
- iii. Worries about complications/adverse reactions
- iv. Self-care advice**
- v. Advice on medical aids** \rightarrow *Community pharmacies*
- vi. Information request (general/disease/complaint/disorder)
- vii. Pharmacy team initiative administration
- viii. Alterations in prescription (not based on medication-surveillance signal)
- ix. Evaluation as result of a consultation by invitation
- x. Evaluation without patient consultation.

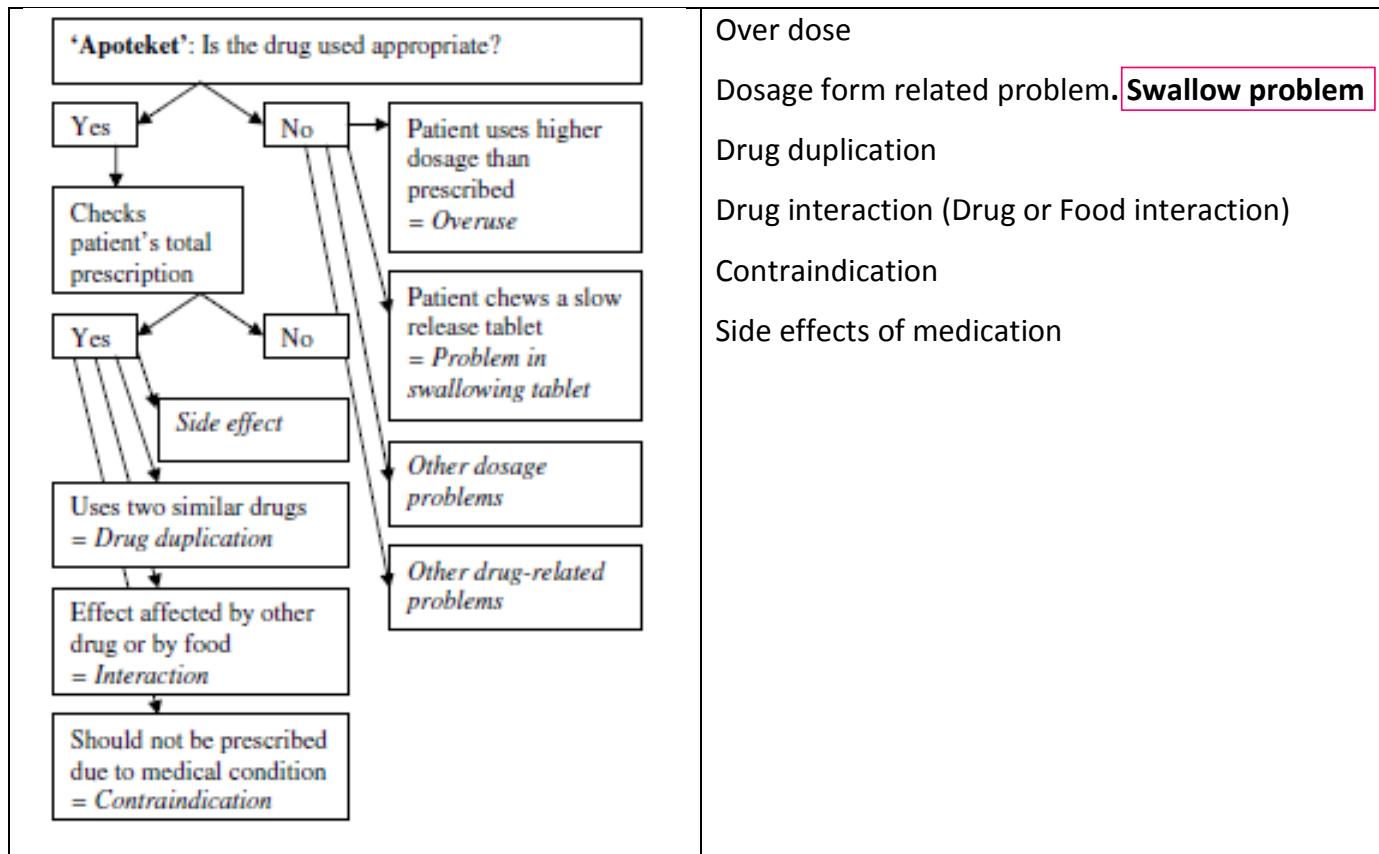


فهي الأدوية التي تصرف بغير بطاقة أو تصرف مخالفة للبيانات

7) Apoteket Classification System

Apoteket (definite singular apoteket) is pharmacy in Swedish language.

Apoteket AB has developed a classification system that Swedish pharmacies can use when counselling patients.



8) Pharmaceutical Care Network Europe (PCNE) system

دوائر
الدواء
والبيئة
والصحة

The original classification was created in 1999 by pharmacy practice researchers during a working conference of the PCNE in an effort to develop a standardized classification system that is suitable and comparable for international studies. Last version V8 June 2017. Updated to be in 2019

It has the following Domains

Domain P: Problem

1. Treatment effectiveness There is a (potential) problem with the (lack of) effect of the therapy	P1.1 No effect of drug treatment/ therapy failure P1.2 Effect of drug treatment not optimal P1.3 Untreated symptoms or indication
2. Treatment safety Patient suffers, or could suffer, from an adverse drug event	P2.1 Adverse drug event (possibly) occurring
3. Others	P3.1 Problem with cost-effectiveness of the treatment P3.2 Unnecessary drug-treatment P3.3 <i>Unclear problem/complaint. Further clarification necessary (please use as escape only)</i>

See the other Domains:

Domain C: Cause of Problem (Prescribing, Dispensing, Use)

Domain I: Planned Interventions

Domain A: Intervention Acceptance

Domain O: Status of ADP

	Primary Domain	Code V8.01	Cause
Prescribing	1. Drug selection The cause of the (potential) DRP is related to the selection of the drug	C1.1 C1.2 C1.3 C1.4 C1.5 C1.6 C1.7	Inappropriate drug according to guidelines/formulary Inappropriate drug (within guidelines but otherwise contra-indicated) No indication for drug Inappropriate combination of drugs or drugs and herbal medication Inappropriate duplication of therapeutic group or active ingredient No drug treatment in spite of existing indication Too many drugs prescribed for indication
	2. Drug form The cause of the DRP is related to the selection of the drug form	C2.1	Inappropriate drug form (for this patient)
	3. Dose selection The cause of the DRP is related to the selection of the dose or dosage	C3.1 C3.2 C3.3 C3.4 C3.5	Drug dose too low Drug dose too high Dosage regimen not frequent enough Dosage regimen too frequent Dose timing instructions wrong, unclear or missing
	4. Treatment duration The cause of the DRP is related to the duration of treatment	C4.1 C4.2	Duration of treatment too short Duration of treatment too long
	5. Dispensing The cause of the DRP is related to the logistics of the prescribing and dispensing process	C5.1 C5.2 C5.3 C5.4	Prescribed drug not available Necessary information not provided Wrong drug, strength or dosage advised (OTC) Wrong drug or strength dispensed
	6. Drug use process The cause of the DRP is related to the way the patient gets the drug administered by a health professional or carer, despite proper dosage instructions (on the label)	C6.1 C6.2 C6.3 C6.4 C6.5	Inappropriate timing of administration and/or dosing intervals Drug under-administered Drug over-administered Drug not administered at all Wrong drug administered
	7. Patient related The cause of the DRP is related to the patient and his behaviour (intentional or non-intentional)	C7.1 C7.2 C7.3 C7.4 C7.5 C7.6 C7.7 C7.8 C7.9	Patient uses/takes less drug than prescribed or does not take the drug at all Patient uses/takes more drug than prescribed Patient abuses drug (unregulated overuse) Patient uses unnecessary drug Patient takes food that interacts Patient stores drug inappropriately Inappropriate timing or dosing intervals Patient administers/uses the drug in a wrong way Patient unable to use drug/form as directed
Use	8. Other	C8.1 C8.2 C8.3	No or inappropriate outcome monitoring (incl. TDM) Other cause; specify No obvious cause

See a detailed table version

N.B. One problem can lead to more interventions

Primary Domain	Code V8.01	Intervention
No intervention	I0.1	No Intervention
1. At prescriber level	I1.1 I1.2 I1.3 I1.4	Prescriber informed only Prescriber asked for information Intervention proposed to prescriber Intervention discussed with prescriber
2. At patient level	I2.1 I2.2 I2.3 I2.4	Patient (drug) counselling Written information provided (only) Patient referred to prescriber Spoken to family member/caregiver
3. At drug level	I3.1 I3.2 I3.3 I3.4 I3.5 I3.6	Drug changed to Dosage changed to Formulation changed to Instructions for use changed to Drug stopped New drug started
4. Other intervention or activity	I4.1 I4.2	Other intervention (specify) Side effect reported to authorities

Acceptance of the Intervention proposals

N.B. One level of acceptance per intervention proposal

Primary domain	Code V8.01	Implementation
1. Intervention accepted (by prescriber or patient)	A1.1 A1.2 A1.3 A1.4	Intervention accepted and fully implemented Intervention accepted, partially implemented Intervention accepted but not implemented Intervention accepted, implementation unknown
2. Intervention not accepted (by prescriber or patient)	A2.1 A2.2 A2.3 A2.4	Intervention not accepted: not feasible Intervention not accepted: no agreement Intervention not accepted: other reason (specify) Intervention not accepted: unknown reason
3. Other (no information on acceptance)	A3.1 A3.2	Intervention proposed, acceptance unknown Intervention not proposed

Status of the DRP

N.B. This domain depicts the outcome of the intervention. One problem (or the combination of interventions) can only lead to one level of solving the problem

Primary Domain	Code V8.01	Outcome of intervention
0. Not known	O0.1	Problem status unknown
1. Solved	O1.1	Problem totally solved
2. Partially solved	O2.1	Problem partially solved
3. Not solved	O3.1	Problem not solved, lack of cooperation of patient
	O3.2	Problem not solved, lack of cooperation of prescriber
	O3.3	Problem not solved, intervention not effective
	O3.4	No need or possibility to solve problem