Table 1: description of Drug Related Problems

- **□ Identification** of DRP is based on the analysis of a drug prescription and according to available clinical and paraclinical data on the patient.
- Only one choice: If the patient's drug regimen reveals several problems, fill up as many intervention form as of problems.

⇒ The question: does this patient develop or is he susceptible to develop a clinical symptom linked to a specific drug or is there a drug problem requiring an intervention to avoid unnecessary mobilization of resources?

resources?		
	DRP	DESCRIPTION
1.1	Non conformity to	- Non conformity of the drug choice compared to the Formulary:
	guidelines or contra-	An equivalent is available on the formulary.
	indication	- Non conformity of the drug choice compared to guidelines:
		An other drug has a better benefit / risk ratio or a better cost / efficacy ratio according to
		current guidelines.
		- There is a physio-pathologic contra-indication for the present drug:
		ie, the patient is asthmatic and is being prescribed beta-blockers.
1.2	Untreated indication	- Valid indication without a drug.
		- A new symptom is not being treated.
		- A drug is missing after transfer.
		- The patient is missing a pre-medication or a prophylactic treatment.
		- A synergic or corrective drug should be associated.
1.3	Subtherapeutic dosage	- Dose too low for this specific patient (daily dose).
1.3	Submerapeutic dosage	A A A
		- Length of the treatment too short
1.4	0 1	(ie, antibiotic prescription of 5 days instead of 10 days)
1.4	Overdosage	- Supra-therapeutic posology:
		dose too high for this specific patient
		there is a risk for accumulation of the drug
		- Duplicate prescription: a same active substance is being prescribed several times (ie,
		oral acetaminophen and oral association of dextropropoxyphen/acetaminophen)
1.5	Drug without indication	- No justified indication for the drug.
		- The drug is being prescribed for a too long period without any risks (ie, antibiotic
		prescribed for 15 days)
		- Therapeutic redundancy: prescription of two different molecules from the same
		therapeutic class
1.6	Drug interaction	A drug interferes with another drug and can lead to a non adapted pharmacological
		impact (over or under expressed)
		- Level according to the French Red Book Vidal©
		- Interaction published but not integrated into Vidal©
		(specify the bibliographic references)
1.7	Adverse drug reaction	The patient presents an adverse drug reaction while on the right posology.
		It can be a clinical, biological, or kinetic effect.
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1.8	Improper administration	The chosen drug is correct but the mode of administration is not adapted:
		- Other route more effective or less costly for the same efficacy
		- the method for administration is not adequate (reconstitution, dilution, manipulation,
		length of administration)
		- Inappropriate drug form
		- incomplete formulation (dosage missing)
		- Inappropriate timing of administration and/or repartition of doses
1.9	Failure to receive drug	- Physico-chemical incompatibility between several injectable drugs: there is a risk for
		precipitation between drugs during perfusion.
		- Compliance problem.
1.10	Drug monitoring	The patient doesn't benefit from a suitable or sufficient follow-up: lab tests, kinetics,
		symptoms (glycemia, EKG, blood pressure, blood concentration of specific drugs)