Protocol for data collection on drugs use in inpatients

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Abstract

Forms used for data collection in the study entitled: “Risk factors for adverse drug reactions in pediatric inpatients: a cohort study”

Forms 1 and 2 must be used in the first pharmaceutical consultation after hospital admission of the patient:

i) form 1 is aimed at obtaining information on the prior history of ADR of the patient; and

ii) form 2 is aimed at obtaining information on the prior history of use of the drug.

Forms 3-6 must be used for the daily recording of information observed in medical prescriptions.

i) form 3 is used to record potential drug interactions;

ii) form 4 for registration of off-label drugs prescribed;

iii) form 5 for the daily recording of the use of drugs; and

iv) form 6 for daily monitoring of prescription errors**.**

Furthermore, form 7 must be used for the registration of abnormal laboratory tests or other suspected adverse events.

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Guidelines

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i) form 1 is aimed at obtaining information on the prior history of ADR of the patient; and

ii) form 2 is aimed at obtaining information on the prior history of use of the drug.

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iii) form 5 for the daily recording of the use of drugs; and

iv) form 6 for daily monitoring of prescription errors**.**

Furthermore, form 7 must be used for the registration of abnormal laboratory tests or other suspected adverse events.

Protocol

Warnings

Because this protocol deals with forms, no safet warnings are required.