Evaluation to improve the quality of medication preparation and administration in pediatric and adult intensive care units

Theresa Hermanspann1,2
Eva van der Linden2
Mark Schoberer2
Christina Fitzner3,4
Thorsten Orlikowsky2
Gernot Marx2
Albrecht Eisernt1,6

1Hospital Pharmacy, RWTH Aachen University Hospital, Aachen, Germany; 2Department of Pediatric and Adolescent Medicine, Section of Neonatology, RWTH Aachen University Hospital, Aachen, Germany; 3Department of Medical Statistics, Medical Faculty RWTH Aachen University, Aachen, Germany; 4Department of Anaesthesiology, RWTH Aachen University Hospital, Aachen, Germany; 5Department of Intensive Care Medicine, RWTH Aachen University Hospital, Aachen, Germany; 6Institute of Pharmacology and Toxicology, Medical Faculty RWTH Aachen University, Aachen, Germany

Purpose: To determine the type, frequency, and factors associated with medication preparation and administration errors in adult intensive care units (ICUs) and neonatal ICUs (NICUs)/pediatric ICUs (PICUs).

Patients and methods: We conducted a prospective direct observation study in an adult ICU and NICU/PICU in a tertiary university hospital. Between June 2012 and June 2013, a clinical pharmacist and medical student observed the nursing care staff on weekdays during the preparation and administration of intravenous drugs. We analyzed the frequency and type of preparation and administration errors and factors associated with errors.

Results: Six hundred and three preparations in the adult ICU and 281 in the NICU/PICU were observed. Three hundred and eighty-five errors occurred in the adult ICU and 38 in the NICU/PICU. There were 5,040 and 2,514 error opportunities, with overall error rates of 7.6% and 1.5%, respectively. The total opportunities for error meant each single step of preparation and administration that was relevant for the drug. Most errors applied to the category “uniform mixing” (adult ICU: n=227, 59%; NICU/PICU: n=14, 37%). The multivariate logistic regression results showed a significantly different influence of the “preparation type” for the adult ICU compared with the NICU/PICU with regard to the occurrence of an error. Preparations for adult patients of the LCD type (liquid concentrate with diluent into syringe or infusion bag) were more often associated with errors than the P (powder in a glass vial that must be reconstituted and diluted if necessary), P=0.012, and LC (liquid concentrate into syringe), P=0.002 type.

Conclusion: “Uniform mixing” was the most erroneous preparation step in intravenous drug preparations in two ICUs. Improvement of nurse training and the preparation of prefilled syringes in the pharmacy might reduce errors and improve the quality and safety of drug therapy.

Keywords: intravenous medication, medication errors, medication safety, patient safety, quality improvement

Introduction
Drug therapy for hospital inpatients is error prone. Errors can cause adverse events and patient harm. MEs can occur during the prescription, preparation, and administration processes.1 Critically ill patients in ICUs usually need numerous intravenous medications, and the error risk is multiplied by the number of calculation and preparation steps. Therefore, the preparation and administration performed by the nursing staff in the ward are complex and time-consuming.2 The administration step of a drug is the last barrier before an error could affect a patient. Error detection is less likely if the same person performs preparation and administration, which is a common practice in ICUs. This makes ICU patients especially vulnerable to preparation and administration error.
errors for intravenous drugs. Studies show error rates for at least one error with preparations of intravenous drugs of 48.4%–97.7%. NICU/PICU patients have an even higher risk of MEs because of the use of off-label and unlicensed drugs and the lack of weight-adapted drug doses and strengths for use in this patient group. The preparation of pediatric intravenous drugs often includes dilution steps from the more concentrated original solutions that are used for adults. Studies with pediatric inpatients alone show error rates of 8.0%–62.7% for the preparation and administration of all types of drugs.

Interventions such as nurse training and central preparation of drugs in the pharmacy reduce the error rates for the preparation and administration of intravenous drugs.

First, we aimed to determine the type and frequency of MEs at the preparation and administration processes in two ICUs in our hospital. The secondary objective was to identify factors that could influence the error rate.

**Patients and methods**

We conducted an observational study in the NICU/PICU and one in the adult ICU at a university hospital. Both studies have been approved by the local ethics committee (EK 151/12 and EK 175/13) and by the local data protection officer of the medical faculty of RWTH Aachen University. Additionally, the director of nursing of the wards provided their informed consent. Furthermore, before conducting the study, the observer introduced herself to every nurse and asked for permission for direct observation. Only after the oral informed consent by nurse, the observer started data collection.

A clinical pharmacist and medical student directly and disguisedly observed the nursing care staff during the preparation and administration of intravenous drugs on the wards. The observation was disguised because the nursing care staff was not aware of the goal of the study. The nursing care staff was informed about an observation on the ICUs and disguisedly observed the nursing care staff during the observation period. Data collection was preceded by 3 pilot-days during which the observers became familiar with the processes and medications on the ward. Between September 2012 and June 2013, the medical student observed the nursing care staff of the NICU/PICU. During this period, the clinical pharmacist supervised the medical student while being available for questions and discussions. Drug prescription in the adult ICU was done in the CPOE system Philips CS770 ICCA, Philips Medical Systems, Andover, MA, USA. After preparing the medication according to the physicians’ prescription on the ward and administering it to the patient, nurses electronically confirmed each administration of medication in the ICCA.

NICU/PICU physicians prescribed medication using the CPOE VISITE 2000, Pharma GmbH, Oberhaching, Germany. Prescriptions were printed out for the nurse to transcribe the medication in the paper-based patient record. The nurse prepared the medication in a separate room with an LAF, administered it to the patient, and confirmed the administration in the patient record.

**Data collection**

Data collection on both wards was performed on weekdays during the day at time periods when usually most preparations were performed. Drug rounds took place in the morning, at lunchtime, and in the evening. The observers attended at least one drug round per day. The sampling was performed by time period and not by patient. A standardized data collection form was used to evaluate the steps of the preparation and administration processes for intravenous drugs, including drug name, solvent, diluent, reconstitution, mixing, dose, labeling, time and route of administration, and administration rate. Errors were classified in one category of ME (Table 1). We chose this classification in accordance with a compilation of the American Society of Hospital Pharmacists in the guidelines on preventing MEs in hospitals with a focus on errors in different preparation steps. Each prepared drug was assigned into the “type of preparation” (Table 2) and “therapeutic drug class” that was defined by the first level of WHO ATC class. If an ME was likely to result in a serious adverse event, the observer intervened in the preparation or administration process. Otherwise, there was no intervention by the observer when an error occurred.

**Definitions**

According to the NCC MERP, MEs are defined as “any preventable event that may cause or lead to inappropri-
In our study, inappropriate medication use was defined as any instance in which the preparation or administration of injectable drugs varied from the physician’s prescription, the SMPC of the drug, hospital policies and procedures, national/international guidelines, internal guidelines, or the good manufacturing practice. Aspects of hygiene (hand disinfection, workplace surface disinfection, and disinfection of vial membrane) were not considered. Additionally, we did not assess the clinical appropriateness of the prescription.

### Data analysis
We calculated the overall error rate as percentage (with a 95% CI) for both ICU and NICU/PICU by dividing the overall number of errors by the number of total opportunities for error for all observed preparations. The total opportunities for error meant each single step of preparation and administration that was relevant for the drug. Additionally, we calculated the error rate as a percentage (with a 95% CI) for both the ICU and NICU/PICU. The error rate in percentage was the number of preparations with at least one error, in relation to the number of all observed preparations multiplied by 100.

Furthermore, we calculated the error rate as a percentage (with a 95% CI) for every category of ME by dividing the number of errors per single category by the number of all observed preparations with relevance for the category. A generalized linear mixed model with a binary distribution (PROC GLIMMIX) was used to investigate the possible influence of “ward” adjusted for “preparation type” and “therapeutic drug class” for the primary outcome to make an error within a drug preparation. The intercept was assumed to be random. We assessed a 5% significance level for each model and adjusted CIs and P-values for multiple testing by the Scheffé method.21 Significant interaction effects were included in the model. Statistical analyses were performed using SAS (version 9.4; SAS Institute Inc., Cary, NC, USA) and R (version 3.3.0).22

### Results
The pharmacist and medical student observed 884 preparations: 603 preparations in the adult ICU and 281 preparations in the NICU/PICU (Figure 1). In two cases in the adult ICU, the observer intervened in the preparation process because of a wrong dose. The first error was related to a tenfold insulin dose and the second one to a tenfold penicillin dose. In the NICU/PICU, there was no intervention. On the adult ICU, 340 preparations had at least one error (error rate 56%, 95% CI: 52–60). There were 385 errors out of 5,040 error opportunities, with an overall error rate of 7.6% (95% CI: 6.9–8.4). In the NICU/PICU, 36 preparations had at least one error (error rate 13%, 95% CI: 8.9–17). There were 38 out of 2,514 error opportunities, with an overall error rate of 1.5% (95% CI: 1.0–2.0). We assigned each preparation into a type of preparation and a therapeutic drug class (Figure 1).

![Table 1 Categories of MEs in the preparation and administration processes](https://www.dovepress.com/)

| Category of ME                  | Definition                                                                 |
|---------------------------------|---------------------------------------------------------------------------|
| 1. Medication                   | Preparation and administration of a drug that differed from the physician’s prescription |
| 2. Type of solution             | Preparation of a drug with the wrong solvent or diluent                    |
| 3. Volume of solution           | Wrong volume of solvent or diluent                                        |
| 4. Uniform mixing               | Insufficient mixing of two solutions                                       |
| 5. Reconstituted                | Insufficient reconstitution of a powder with the solvent                   |
| 6. Dose                         | Dose of a drug that differed >5% from the physician’s prescription         |
| 7. Labeling                     | Incorrect labeling with omission of the drug name and/or concentration     |
| 8. Timing error                 | Time period of >2 hours between preparation and administration of the drug |
| 9. Route of administration      | Wrong route of administration                                             |
| 10. Infusion rate               | Wrong infusion rate                                                       |

**Abbreviation:** ME, medication error.

### Table 2 Type of preparation

| Type of preparation | Definition                                                                 |
|---------------------|---------------------------------------------------------------------------|
| P                   | Powder in a glass vial that must be reconstituted and diluted if necessary |
| LC                  | Liquid concentrate into syringe                                           |
| LCD                 | Liquid concentrate with diluent into syringe or infusion bag              |

**Table 1** Categories of MEs in the preparation and administration processes

**Table 2** Type of preparation
Most errors in the adult ICU occurred in the categories “uniform mixing” (n=227 in 351 preparations, 65% error rate; 95% CI: 60–70) and “labeling” (n=109 in 603 preparations, 18% error rate; 95% CI: 15–21). In the NICU/PICU, most errors were related to uniform mixing (n=14 in 226 preparations, 6.2% error rate; 95% CI: 3.1–9.3) and the infusion rate (n=9 in 281 preparations, 3.2% error rate; 95% CI: 1.1–5.3) (Table 3). Results of the multivariate generalized linear mixed model showed a significant interaction between the “preparation type” and “ward” on the occurrence of an error (P=0.0456). The “preparation types” differed significantly for the adult ICU, but not for the NICU/PICU, whereas the therapeutic drug class had no influence on the occurrence of an error in the multivariate model (P=0.149). Preparations for adult patients of the LCD type were more often associated with errors than the P (OR =2.7, 95% CI: 1.2–6.1, P=0.012) and LC (OR =1.8, 95% CI: 1.2–2.7, P=0.002) type (Figure 2). The results for the NICU/PICU did not show a significant influence of the “preparation type” (Figure 2) on the occurrence of an error.

Discussion

The error rates in our studies for the preparation and administration of intravenous drugs for an adult ICU and NICU/PICU were 7.6% and 1.5%. It is difficult to compare our results with other studies because they were performed in different settings (medical discipline and type of ward) with differences in the methods and definitions.

The results of studies in adult patients for all types of medication showed error rates for preparations with at least one error ranging from 3.3% to 44.6%.3,4,23–25 Within these
studies, intravenous medication often has much higher error rates than other types of medications. Nguyen et al reported an error rate of 72.2% in intravenous vs 11.8% in oral preparations. Studies regarding the preparation and administration of intravenous drugs alone showed error rates as high as 48%–97%. Our results for the error rate in the adult ICU (56%) for preparations with at least one error divided by the total number of preparations fall well within the lower range.

For the pediatric patient group, we found no studies on intravenous drug preparations alone. The studies by Chedoe et al and Chua et al on all types of medication showed that the intravenous route contributed significantly to the rate of drug administration errors. Cousins et al reported a 79% error rate for preparations that were not mixed properly in a German hospital. In a study by Chedoe et al, “not mixed” was the third most frequent error category in the pre- and postintervention period.

The second error group for adult patients was “labeling” (18%). To increase the safety in the administration process, national and international organizations and institutions recommended standardized colored syringe drug labeling according to the International Organization for Standardization standard 26825. These labels were already used in the

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**Table 3 Frequencies of error categories**

| Category of medication error | Number of preparations (errors) in the adult ICU | Error rate in the adult ICU (%) | Number of preparations (errors) in the NICU/PICU | Error rate in the NICU/PICU (%) |
|-----------------------------|--------------------------------------------------|---------------------------------|-----------------------------------------------|-------------------------------|
| Medication                  | 603 (1)                                           | 0.2                             | 281 (0)                                       | 0                             |
| Type of solution            | 465 (0)                                           | 0                               | 253 (1)                                       | 0.40                          |
| Volume of solution          | 465 (22)                                          | 4.7                             | 253 (4)                                       | 1.6                           |
| Uniform mixing              | 351 (227)                                         | 65                              | 226 (14)                                      | 6.2                           |
| Reconstituted               | 144 (8)                                           | 5.6                             | 96 (0)                                        | 0                             |
| Dose                        | 603 (2)                                           | 0.3                             | 281 (0)                                       | 0                             |
| Labeling                    | 603 (109)                                         | 18                              | 281 (3)                                       | 1.1                           |
| Timing error                | 603 (0)                                           | 0                               | 281 (7)                                       | 2.5                           |
| Route of administration     | 603 (0)                                           | 0                               | 281 (0)                                       | 0                             |
| Infusion rate               | 603 (16)                                          | 2.7                             | 281 (9)                                       | 3.2                           |

Abbreviations: ICU, intensive care unit; NICU, neonatal intensive care unit; PICU, pediatric intensive care unit.

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**Figure 2** Comparison of the preparation types (P, LC, and LCD) with regard to the occurrence of an error in the adult ICU and NICU/PICU.

Note: Preparation type 1 vs 2: corresponding to the preparation types LC vs P, LCD vs P, and LCD vs LC.

Abbreviations: CI, confidence interval; ICU, intensive care unit; LC, liquid concentrate into syringe; LCD, liquid concentrate with diluent into syringe or infusion bag; NICU, neonatal intensive care unit; P, powder in a glass vial that must be reconstituted and diluted if necessary; PICU, pediatric intensive care unit.

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ME. Relating to our local hospital guidelines, the sufficient mixing is defined as shaking at least three times. Sufficient mixing is necessary for a homogeneous distribution of the drug in the diluent and consequently for a uniform dosing. This is particularly important for continuous infusions with an infusion pump. It is difficult to compare this result with those in previous studies because different definitions for preparation errors were used. “Uniform mixing” is often part of the category “incorrect drug preparation” without differentiating between the individual preparation steps. Cousins et al reported a 79% error rate for preparations that were not mixed properly in a German hospital. In a study by Chedoe et al, “not mixed” was the third most frequent error category in the pre- and postintervention period.
two ICUs before conducting the study. Nevertheless, there is still a lack of some preprinted labels, especially for antinfectives. In a study by Cousins et al, the name of the drug was missing in 19% of preparations, and the dose was missing in 29% of preparations.7 Despite available standardized drug labels, nearly one-third of labeling errors in our study (29%) concerned the drug propofol. We assume that the nurses may have omitted the labels with the thought that the characteristic lipid emulsion of propofol could make labeling redundant.

The second most frequent error category in the NICU/PICU was the “infusion rate”. All nine errors concerned a lower infusion rate with >5% deviation from the SMPC. This is explained by the fact that most intravenous infusions in our NICU/PICU are given short infusions of either 30 or 60 minutes. Analgesics are usually given faster, especially in cases of acute pain, to achieve a faster onset of action. The incidents were classified as errors because they meet the error criteria defined in this study. However, it is debatable whether they constitute an error in a wider medical sense. Other studies also reported administration errors that were characterized by rapidly administered bolus doses.7,28

Multivariate analysis showed that the type of preparation had a significant influence on the occurrence of an error in the adult ICU. The preparation type “liquid concentrate with diluent into syringe or infusion bag” was more often associated with an error than the other types of preparations (“powder in a glass vial that must be reconstituted and diluted if necessary” and “liquid concentrate into syringe”). One reason could be the twofold dilution in this type of preparation. The therapeutic drug class had no influence on the occurrence of an error in both ICUs.

Apart from the preparation type, the nurse–patient ratio, the number of different drugs, and the place of drug preparation may influence the error rate. In our adult ICU, the number of patients per nurse was on average higher than in the NICU/PICU (1.9 vs 1.4 patients per nurse). Studies showed that the occurrence of MEs was associated with the number of patients per nurse.10,24 Additionally, the number of different drugs in the observation period was much higher in the adult ICU than in the NICU/PICU (64 vs 45 drugs). These factors may explain the lower error rate in our pediatric patient group. Nevertheless, the preparation of intravenous drugs for pediatric patients typically requires more calculation and dilution steps, which increases the risk of error. In addition to human and organizational factors, the environment has a fundamental role in the management of drug therapy. Environmental factors include noises, interruptions, and disorder.29 Drug preparation in the adult ICU was performed on a dedicated bedside table on the ward, while the preparation in the NICU/PICU was performed in a separate room with an LAF. Both settings were affected by interruptions and the lack of preparation space. Strategies to reduce interruptions and distractions are needed.

The results of our studies allow for the development of interventions to improve our preparation and administration process. Taxis and Barber showed that one cause of MEs is lack of knowledge of preparation and administration procedures.28 Studies demonstrated that nurse education is one type of intervention to reduce MEs.18,30,31 Providing nurse training and information about the most frequent errors, “uniform mixing”, “labeling”, and “infusion rate”, in our study may help to reduce these error types. In a study by Chua et al, an intervention between two observation periods showed a reduction in the administration errors from 44.3% to 28.6%.19 The intervention involved a presentation for the nursing staff that involved sharing the error results of the first observation period. The intervention increased awareness of preparation and administration errors.

Another possible strategy is the preparation of ready-to-use syringes in the pharmacy. De Giorgi et al showed that ready-to-use syringes have a good cost–efficacy ratio as a safety tool for the preparation and administration processes.32 Another study found that prefilled syringes are more expensive than ampules, but prefilled syringes could reduce MEs.23

Limitations
Within the last years, a CIRS was introduced in both wards. Errors associated with the preparation and administration of drugs were not recorded during our study period. In the time period after our study until 2016, recorded errors in the CIRS resulted in an implementation of the four-eyes principle during the preparation of drugs by two nurses in all neonatal and pediatric wards. Incident report review like CIRS is another type of method to detect medications errors. However, its detection rate is lower than that of observational studies.34 Nevertheless, these implemented measures could have influenced the error rate within the last 5 years. Presenting of CIRS reports could increase the awareness of preparation and administration errors in both wards and therefore reducing the error rate. Additionally, the four-eyes principle could decrease the error rate in the NICU/PICU. Regardless of the implemented measures within the last 5 years, the resulting error rates of our study revealed the processes that are most error prone during the preparation and administration of intravenous drugs in our study. These results allow developing interventions to improve these error prone processes.
The two ICUs in which the studies were conducted are difficult to compare. Both wards differ regarding the patient characteristics, sample size, medication, and structural and organizational setting.

We only analyzed the preparation and administration process of intravenous drugs. Further studies are needed to evaluate the preparation and administration of other types of medication as well as the prescribing, transcribing, and dispensing process with the aim to improve medication safety.

We did not judge the clinical relevance of each error. Most errors likely will not cause severe patient harm. To assess the severity of each error, an independent expert panel of physicians, pharmacists, and nurses is needed.

We chose the disguised observation technique in our study. This method gives more reliable and objective results than spontaneous reporting or chart review, but it is time-consuming. One disadvantage of this method is the influence on the nurses’ behavior (Hawthorne effect). The error rate could increase if the observer caused nervousness among the nurses or the error rate could decrease if nurses were more watchful during the observation. Additionally, the observers’ behavior could influence the nurses too. A discrete and nonjudgmental behavior has less influence on the nurses. This would play an important role, if the observer decided to intervene to avoid an error, which could result into a serious adverse event. Although there might be a small effect on the nurses’ behavior, studies have shown that the observer does not have a significant influence on the error rate.

The two observers could not be on the wards at all times and could not observe two preparations at the same time; therefore, some preparations were likely missed. Furthermore, we observed the wards during time periods when most medications were prepared, including the time periods with the highest workload for the nurses, which could increase the error rate. Both observers had the same standardized data collection form to ensure that the same error definition was applied. They conducted a pilot study to achieve familiarity with the preparation and administration processes. This approach resulted in a high accordance in the judgment of errors between these two observers. Nevertheless, there might be interobserver variability but the influence on the error rate is rather low.

Conclusion
We found that the type of preparation had a significant influence on the occurrence of an error in the adult ICU. Preparations with more than one dilution step were more often associated with errors than other types of preparations. In both wards, the most erroneous ME category was “uniform mixing”. Nurse training may be a feasible intervention to reduce errors. Additionally, the preparation of prefilled syringes in the pharmacy and improvements in the nurse–patient ratio and preparation location might reduce the error rate and thus improve the quality and safety of drug therapy.

Abbreviations
ATC, Anatomical Therapeutic Chemical; CI, confidence interval; CIRS, critical incidence reporting system; CPOE, computerized physician order entry; ICCA, IntelliSpace Critical Care and Anesthesia; ICU, intensive care unit; LAF, laminar air flow; LC, liquid concentrate into syringe; LCD, liquid concentrate with diluent into syringe or infusion bag; ME, medication error; NCC MERP, National Coordinating Council for Medication Error Reporting and Prevention; NICU, neonatal intensive care unit; P, powder in a glass vial that must be reconstituted and diluted if necessary; PICU, pediatric intensive care unit; SMPC, summary of product characteristics.

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Author contributions
TH designed and conducted the analysis. She analyzed and interpreted the data and drafted the first manuscript. EvdL conducted the analysis. She interpreted parts of the data and reviewed the manuscript. MS assisted with the analysis and interpretation of the data and revised the manuscript. CF interpreted the data while conducting the statistical tests and reviewed the manuscript. TO contributed to the conception of the analysis and reviewed the manuscript. GM contributed to the conception of the analysis and reviewed the manuscript. AE contributed to the conception and interpretation of the analysis and reviewed the manuscript. All authors contributed toward data analysis, drafting and critically revising the paper, gave final approval of the version to be published, and agreed to be accountable for all aspects of the work.

Disclosure
No benefits in the form of an honorarium, grant, or other form of payment contributed to the manuscript. The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest and report no conflicts of interest in this work.

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