Effects of bar code-assisted medication administration (BCMA) on frequency, type and severity of medication administration errors: a review of the literature

Jeroen Hassink,1 Mark Jansen,1 Pieter Helmons2

ABSTRACT
Bar code-assisted medication administration (BCMA) is increasingly being adopted as an additional tool in the prevention of medication administration errors. This literature review summarises the evidence behind the effects of BCMA technology on medication safety. Although most studies show an error-reducing effect of BCMA technology, compliance with the new technology after its implementation and the long-term effects on error reduction are often not assessed. Most importantly, the effect of medication error reduction on patient outcomes is limited.

Introduction
The medication distribution process is an important source of medication errors. Medication error rates reported in the literature vary widely depending on the methodologies and definitions used. A recent review summarised the prevalence of medication errors as 5.7% of administrations (range 0.038–56.1%, n = 51 studies), 1.07 errors per 100 patient-days (range 0.35–12, n = 9) or 6% of patients hospitalised (range 0.93–24%, n = 7).1 Most errors originate in the medication administration process (median 55%, range 9–90.7%).1 With few barriers to prevent them from occurring, only 2% of medication administration errors are intercepted at the patient bedside.2 Bar code-assisted medication administration (BCMA) is increasingly adopted as an additional barrier in the prevention of medication administration errors. In 2009, 27.9% of hospitals in the USA had implemented BCMA,3 which increased to 50.2% in 2011.4

BCMA technology is developed to improve compliance with checking the ‘five rights’ of medication administration: right patient, right route, right drug, right dose and right time. The right patient is identified by matching the unique bar code on the patient wristband to the patient information in the electronic medication administration record (eMAR). The right drug, right dose, right dosage form and right time are checked by matching the bar code on every unit- or multidose medication to the information in the eMAR. In a 2009 position statement, the American Society of Health-System Pharmacists encouraged health systems to adopt BCMA technology to improve patient safety and the accuracy of medication administration and documentation.5 Most studies evaluating the effect of BCMA on medication administration errors have been conducted in the USA. However, this technology is also used in European countries including Denmark, Italy and the Netherlands6 and, in 2006, the Council of Europe Expert Group on Safe Medication Practices also encouraged the use of electronic systems to improve the safety of medication administration.7 In June 2010 the general assembly of the European Association of Hospital Pharmacists called for the implementation of bar-coded single dose-packed drugs in national and European regulations.8

While BCMA as a tool in the prevention of medication administration errors makes intuitive sense, there is limited evidence demonstrating the effect of this intervention on medication administration errors and patient outcomes. In addition, increased workload is a commonly voiced concern by nursing staff as the use of bar coding technology can potentially result in a longer duration of medication administration. This review of the literature focuses on (1) the effect of BCMA on frequency, type and severity of medication administration errors and (2) the effect of BCMA technology on the duration of the medication administration process.

Methods
Study selection
In May 2012 a PubMed search was performed to select studies investigating at least one of the following topics: the effect of BCMA on the rate or severity of medication administration errors or studies evaluating the effect of BCMA on the duration of administering medication. Only studies with a prospective design and in which observational techniques were used to measure medication errors and/or administration time were included. The detailed search criteria and selection procedure of the 10 articles included in this study are shown in figure 1.9–18 We also reviewed the reference lists of the selected articles. This revealed the full text article19 of an abstract we had already selected.15 In addition, we included a study that met the above-mentioned criteria and was published in this journal and a Dutch pharmaceutical journal, not indexed in PubMed.20 21 As a result, a total of 11 studies were included9–14 16–21

Settings and intervention
The studies were conducted on wards with different levels of care and in organisations with varying medication use processes (table 1). Implementation of BCMA was accompanied by the implementation of an eMAR in all studies. In two studies the introduction of BCMA was accompanied...
by additional interventions such as simultaneously implementing bedside assortment picking (table 1). In all studies, error rates were calculated using the same formula: total errors divided by the sum of observed administrations and omissions. In the study by Franklin et al, bar code technology was used to stock the automated dispensing cabinet and assure the correct identity of the medication. At the bedside, BCMA was then used to assure the correct identity of the patient.

### Results

#### Error frequency

Error rates before and after implementation of BCMA are summarised in table 2.

As wrong time errors are generally considered to be less severe, the results are reported as total errors and errors excluding wrong time errors. Baseline error rate varied between 5.8% and 25.3% if time errors were included and between 1.6% and 27.3% when time errors were excluded. Most studies show a 30–50% reduction in medication administration errors after implementation of BCMA when time errors are excluded. However, implementation of BCMA does not result in a consistent reduction when time errors are included.

#### Error type

The type and number of error categories varied between studies. Error categories that were assessed in at least three studies and are expected to be reduced by BCMA are omissions, wrong drug errors, unauthorised drug errors, wrong dosage form errors and extra dose errors. Only one study did not find a reduction in unauthorised drug errors and omissions and wrong drug and wrong dose errors even increased. Wrong dose errors also increased in the ICU setting in the study by Helmons et al. Wrong dosage form errors and extra dose errors increased in the study by Ros et al.

Wrong route errors are not expected to be influenced by BCMA and wrong time errors only partially. Reduction of these errors was inconsistent among studies. Most studies were underpowered to identify statistically significant differences within individual categories.

Overall it seems that wrong time errors are the most frequently occurring.

#### Error severity

Adverse drug events (ADEs) are defined as an injury resulting from the use of a medicine or omission of an intended medicine. This definition includes adverse drug reactions and harm from medication incidents. As a result, medication errors resulting in harm are considered ADEs. An error that could potentially lead to harm is a potential ADE. One study assessed the severity of observed ADEs and two studies categorised the potential severity of observed administration errors (table 3).

Morris et al found that BCMA reduced the risk of preventable ADEs by 47% and Poon et al showed a 50.8% reduction in potential ADEs. In this latter study the reduction in many of the potential ADEs could be attributed to improved medication administration documentation. Franklin et al did not find a reduction in error severity.
<table>
<thead>
<tr>
<th>Study</th>
<th>Type of ward</th>
<th>Hospital</th>
<th>Setting pre-intervention</th>
<th>Setting post-intervention</th>
<th>Other points of interest</th>
<th>Observation period</th>
</tr>
</thead>
<tbody>
<tr>
<td>Paoletti et al.</td>
<td>Cardiac (telemetry) Medical-surgical</td>
<td>20-bed cardiac ward 36-bed medical Surgical ward in a general hospital, Lancaster, USA</td>
<td>Decentralised cabinet distribution system Handwritten order Handwritten paper MAR</td>
<td>Decentralised cabinet distribution system Handwritten order eMAR with pharmacist order entry BCMA</td>
<td>1.5 year surveillance data</td>
<td>ND</td>
</tr>
<tr>
<td>Poon et al.</td>
<td>ICU Medical Surgical</td>
<td>35 units in a 735-bed tertiary academic medical centre, Boston, USA</td>
<td>CPOE MAR transcribed by nurses</td>
<td>CPOE eMAR BCMA</td>
<td>Transcription errors Severity classification of potential ADEs 2 year surveillance data (in supplement)</td>
<td>2-4 weeks before and 4-8 weeks afterwards. 4 h observation of staff nurses on 35 observed units</td>
</tr>
<tr>
<td>Franklin et al.</td>
<td>Surgical</td>
<td>28-bed ward in a teaching hospital, London, UK</td>
<td>Stock cupboards and two drug trolleys Drug prescription on paper MAR</td>
<td>ADC and two electronic drug trolleys CPOE bar code scan used to confirm drug identity when loading medication into drawer ADC and for patient identification eMAR with manual confirmation of administration</td>
<td>Prescribing errors Staff time spent on medication tasks Potential severity assessment of observed errors</td>
<td>3-6 months before and 6-12 months after. Sample of 56 drug rounds before and 55 after (including nights and weekends) during a 2-week period</td>
</tr>
<tr>
<td>Helmons et al.</td>
<td>ICU (medical- surgical) Medical-surgical</td>
<td>13- and 20-bed ICU 22-, 26-bed medical surgical ward in a 386-bed academic teaching hospital, San Diego, USA</td>
<td>Unit-based ADCs CPOE Printed paper MAR manually updated</td>
<td>Unit-based ADCs CPOE eMAR BCMA</td>
<td>Medication administration accuracy Time spent on medication tasks</td>
<td>1 month before and 3 months after implementation. During week and weekend days focus on medication round 09:00</td>
</tr>
<tr>
<td>DeYoung et al.</td>
<td>ICU</td>
<td>38-bed medical ICU in a 744-bed community teaching hospital, Grand Rapids, USA</td>
<td>ND</td>
<td>Handwritten or preprinted orders eMAR with pharmacist order entry BCMA</td>
<td>–</td>
<td>1 month before and 4 months after. 24 h a day during 4 days</td>
</tr>
<tr>
<td>Morris</td>
<td>NICU</td>
<td>36-bed ward in a children's hospital, Iowa City, USA</td>
<td>Handwritten orders entered by pharmacist in pharmacy information system Paper MAR on which orders were transcribed and administrations recorded</td>
<td>Handwritten orders entered by pharmacist in pharmacy information system eMAR bidirectionally interfaced with pharmacy information system BCMA</td>
<td>Severity assessment of observed preventable ADEs</td>
<td>19 consecutive weeks before implement-ation and 1 month after implementation during 31 weeks</td>
</tr>
<tr>
<td>Ros et al., Wesselink et al.</td>
<td>Neurological</td>
<td>42-bed ward, community teaching hospital, Apeldoorn, The Netherlands</td>
<td>Dispensing to the ward from pharmacy by drug trolley CPOE eMAR with manual confirmation of administration</td>
<td>BAP cart</td>
<td>Time spent on medication tasks</td>
<td>1 year and 8 months before and 3 months after. Three daily medication rounds during 21 days</td>
</tr>
</tbody>
</table>

ADC, automated dispensing cabinet; ADE, adverse drug events; BAP, bedside assortment picking; BCMA, bar code-assisted medication administration; CPOE, computerised physician order entry; eMAR, electronic medication administration record; ICU, intensive care unit; MAR, medication administration record; ND, not determined.
Duration of medication administration

The general idea that the use of BCMA technology is time-consuming for nursing staff is considered a barrier to implementation. Seven studies addressed this topic (table 4). 11 12 16-19 21

Two studies 11 12 evaluated the time spent by nursing staff to complete the medication administration task and three 16 17 21 studies measured the duration of each administration. Two studies determined the percentage of total nursing time spent on medication administration by using either the time and motion method 15 or the work sampling method. 18 No increase in medication administration time was found.
Poon et al 12 also found that, after implementation of BCMA, the time spent on direct patient care activities varied among the different study settings (table 1)—for example, medication in an ICU is generally administered intravenously in an area with a higher nurse-to-patient ratio. Indeed, observation of different types of medication errors than observations performed by BCMA technology. 10-12 21

Poon et al 12 and Dwibebi et al 20 also found that, after implementation of BCMA, the time spent on direct patient care activities increased.

Discussion

The effect of BCMA on the medication error rate is variable among the studies included in this review. BCMA technology seems to decrease the incidence of medication administration errors when time errors are excluded. However, the studies included in this review are heterogeneous.

First, the number and types of administration errors included in the studies vary. In some studies error categories that are not reduced by BCMA are included (eg, technique errors, wrong route errors). This influences the baseline error rate and dilutes the overall effect size of BCMA technology. 10-12 21 Second, the study setting has an effect on the baseline prevalence of medication errors and therefore on the potential effect after implementation of BCMA. As an example, medication in an ICU is generally administered intravenously in an area with a higher nurse-to-patient ratio. Indeed, observation of medication administration in an ICU setting resulted in the detection of different types of medication errors than observations performed on a general medicine ward. 21 Furthermore, medication use processes varied among the different study settings (table 1)—for example, dispensing of drugs by the pharmacy, use of traditional ward stock or use of automatic dispensing cabinets.

There is also a difference between studies in the time of observation (eg, continuous observation or observing specific medication rounds). As the time of the medication administration round is a determinant for medication errors, 24 25 the moment of observation could influence the baseline error rate.

In two studies, 11 20 the intervention was comprised of more than BCMA and an eMAR. It is therefore not possible to contribute the
error reductions either to BCMA technology or to the other intervention (e.g., automated dispensing cabinet) in these studies. The degree of implementation of the technology is of importance to the results. Shortcomings in design, implementation and workflow integration encourage workarounds. The current study results might therefore reflect the impact of the technology in the context of its implementation rather than the impact of the technology itself. Not all studies evaluated user compliance with the new technology. As a result, workarounds could have influenced the effect of BCMA on medication administration errors. Helmons et al and Paoletti et al reported on the compliance rate which was around 90%. Poon et al reported that 20% of the drugs administered using bar code eMAR technology were given without the bar code scanning step during the study period. However, no studies evaluated which errors detected in the study were the result of non-compliance. Although the goal of BCMA is to enhance medication safety, studies that evaluate the prevention of potential harm after implementation of BCMA are limited. Only two studies showed a reduction in the severity of potential ADEs. These limited data support the beneficial effects of BCMA and eMAR on patient outcomes. Evidence on the long-term effect and safety of BCMA is also limited. However, this information is important as workarounds evolve over time. The duration of the positive effects of BCMA on medication administration errors varied from 1 month to 12 months after implementation. Paoletti et al and Poon et al reported data on long-term medication administration error warnings after BCMA implementation. In both studies the number of warnings remained constant during periods of 1.5 and 2 years after implementation of BCMA, respectively, suggesting a long-term effect of this technology in the detection of medication errors.

BCMA did not increase the time spent on medication administration. This is a reassuring finding as nursing staff are concerned about the time-consuming aspects of BCMA technology. The successful implementation of BCMA is the culmination of judicious planning, design, testing, training and support that occurred before, during and after BCMA deployment. The degree of implementation of BCMA technology is therefore an important variable in studies evaluating the effect of BCMA.

This review of the literature generally found a positive effect of BCMA on decreasing medication errors without increasing medication administration time. However, these results are difficult to interpret because of the variability in study design, intervention and reporting of outcome measures and confounders. We have created a study design and reporting checklist as a guide for future research in this area (figure 2), although we realise that conducting a study that meets all of these criteria will not be easy.

### Table 4 Results of studies evaluating the influence of BCMA on time spent on medication administration related tasks

<table>
<thead>
<tr>
<th>Outcome measure</th>
<th>Baseline</th>
<th>Post-BCMA</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Franklin et al²³</td>
<td>Mean (range) duration of each drug round (min)</td>
<td>50 (15-105)</td>
<td>40 (16-78)</td>
</tr>
<tr>
<td>Helmons et al²²</td>
<td>Median (range) duration of a medication administration round on the general medicine ward (min)</td>
<td>10 (1-30)</td>
<td>10 (1-50)</td>
</tr>
<tr>
<td>Wesselink et al²¹</td>
<td>Median (range) duration of a medication administration round on the ICU (min)</td>
<td>12 (1-58)</td>
<td>13.5 (1-53)</td>
</tr>
<tr>
<td>Wesselink et al²¹</td>
<td>Mean duration of administration per drug (min)</td>
<td>1.294</td>
<td>1.198</td>
</tr>
<tr>
<td>Poon et al¹⁹</td>
<td>Percentage of time spent on administering medication</td>
<td>26.9%</td>
<td>24.9%</td>
</tr>
<tr>
<td>Poon et al¹⁹</td>
<td>Percentage of time spent on direct patient care</td>
<td>26.1%</td>
<td>29.9%</td>
</tr>
<tr>
<td>Dwibedi et al²⁶</td>
<td>Mean duration of administration activity (s)</td>
<td>59.8</td>
<td>45.5</td>
</tr>
<tr>
<td>Tsai et al²⁷</td>
<td>Mean duration of time spent on direct patient care (s)</td>
<td>47.4</td>
<td>182.3</td>
</tr>
<tr>
<td>Huang et al²⁸</td>
<td>Mean working time for oral medication administration (s)</td>
<td>36.49</td>
<td>18.42</td>
</tr>
<tr>
<td>Huang et al²⁸</td>
<td>Percentage of time spent on medication-related tasks</td>
<td>Paper group</td>
<td>BCMA group</td>
</tr>
<tr>
<td>Huang et al²⁸</td>
<td>Percentage of time spent on direct patient care</td>
<td>25.0%</td>
<td>17.4%</td>
</tr>
</tbody>
</table>

*In this study a cross-sectional design rather than a before-after design was used. BCMA, bar code-assisted medication administration; ND, not determined in original publication.
Key messages

- Current studies generally support the potential of bar-code-assisted medication administration (BCMA) technology to reduce medication administration errors.
- Current studies do not indicate an increase in nursing time spent on medication administration when using BCMA technology.

Conclusions

The results of this review generally support the medication administration error reducing potential of BCMA technology up to 1 year after implementation without indications of increasing nursing time spent on medication administration. However, current studies do not always mention user compliance and degree of implementation, factors narrowly related to the effectiveness of BCMA technology and necessary to ascertain the maximum achievable effectiveness. Future research should focus on the long-term effects of BCMA on medication error reduction, the causes of errors after BCMA implementation, the effects on nursing workflow and the harm prevented by this technology.

Correction notice This article has been corrected since it was first published Online First. ‘Dr’ has been removed from the beginning of Jeroen Jules Margriet Hassink’s name in the ‘Correspondence to’ section of the article.

Contributors JJMH revised the manuscript based on the comments of the reviewers. MMPMJ and PJH thoroughly revised the new version of the manuscript.

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References