

Warfarin prescription and administration: reducing the delay, improving the safety

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Abstract

Warfarin is frequently administered to hospital patients. The prescription and administration of this medication are particularly susceptible to error. Factors contributing to this include the narrow therapeutic index, patient-specific target range, and the need for regular INR monitoring. NICE guidelines state that warfarin should be given at the same time every day and the Bristol Royal Infirmary guidelines are warfarin to be given at 14:00. The 14:00 dosing ensures standardisation of administration; poor adherence to this recommendation may cause patient harm. We noticed that many warfarin doses were often given outside of maximal staffing hours and it was often left to the on call doctor to prescribe warfarin at erratic and inconsistent times.

Our primary aim was to reduce the number of adverse outcomes associated with warfarin prescription and administration. We targeted two system measures: the proportion of warfarin administrations occurring within an hour of the 14:00 prescription and the proportion of INR results outside target range. We employed the model for improvement and carried out our project across seven acute medical wards. Baseline data showed that only 24% of doses were being given within an hour of the recommended time and 64% of doses were being given after 17:00 during minimal staffing hours. We successfully introduced a warfarin box within our trust which demonstrated an improvement in warfarin administration from 24% of patients receiving their warfarin within an hour of 14:00 to 49% and this was subsequently associated with a reduction in INRs above target range (23% to 9%).

Problem

Warfarin is a commonly prescribed medication and continues to be an important therapy in the prevention of arterial and venous thrombosis, and furthermore in the prevention of stroke related to atrial fibrillation[1-4]. Due to the therapeutic rise in international normalised ratio (INR), the predominant risk of warfarin is bleeding, which has the potential to cause significant morbidity or mortality if not monitored closely[5]. However, for many patients the increased risk of clot formation or stroke means that the risk of harm from warfarin therapy is acceptable and outweighed by the benefits.

The guidance for warfarin administration at University Hospitals Bristol has changed from 18:00 to 14:00 daily[6]. Warfarin is a "critical" medication, meaning that each dose should be administered within 60 minutes of the prescribed time on the drug chart.

Our project aimed to improve the prescription and administration of warfarin so that we were meeting our trust guideline of 14:00. Firstly, we aimed to increase the proportion of warfarin doses administered correctly as a process measure (ie within 60 minutes of the prescription time) and secondly, as an outcome measure, to reduce the number of "INR over 5" incidents occurring monthly.

Frequently, patients do not receive their warfarin within 60 minutes of the time prescribed. This introduces potential safety issues:

- Reducing or increasing the interval between doses may result in a less stable INR pattern

- A less stable INR pattern may increase the likelihood for INRs to move outside of the target range

- If warfarin is not prescribed before 5pm, the on-call doctor will be called to write the prescription: involving a busy doctor who will likely be less familiar with the patient than their regular doctor. This increases the risk of prescription errors, inaccurate dosing.

Background

Warfarin is the most commonly used oral anticoagulant and it is a frequently prescribed medication in the hospital setting. Given its narrow therapeutic index and need for regular INR monitoring it is often susceptible to medication incidents. NICE guidelines state that warfarin should be given at the same time every day and the Bristol Royal Infirmary guidelines are warfarin to be given at 14:00[3,4]. Traditionally warfarin is given at 18:00 as per community dosing times, however our trust found the 14:00 dosing ensures standardisation of administration and ensures that doses are given at a time of maximal staffing to ensure prescriptions aren't missed. Furthermore a previous quality improvement project carried out at another trust showed that changing the time of administration from 18:00 to 14:00 was associated with a reduction in the number of warfarin doses given out of hours[7].

Warfarin doses are prescribed individually, requiring a doctor's signature for each one. In the hospital setting warfarin dosing may require continuous adjustment to achieve a stable INR within the target range, for example due to catabolic illness status and new medications prescribed. This means that often only one to three

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doses of warfarin are prescribed at a time for in-patients.

Dosing is based on the INR result, generated by the hospital laboratory technicians from a "clotting tube" blood sample (result usually received within a couple of hours of arriving at the lab, unless sent in an "urgent" bag).

Baseline measurement

We recorded baseline data from two buildings in our hospital; four wards in the "new" hospital and three wards in the "old" hospital.

Baseline data was collected twice weekly, over a month period. We recorded the following each day for all patients taking warfarin: administration time of warfarin, the time the INR result was available on the hospital computer system (if one had been requested), and the INR result (with a reference target range for each patient). Data for patients on a warfarin loading regimen was not identified and therefore sub therapeutic INRs could not be reliably analysed.

Data was collated onto a run chart to compare times of warfarin administration each day (including a daily average). Tabulated results and the run chart are attached. Of 111 warfarin dosing events recorded, 27 (24%) were given within an hour of 14:00. Forty (36%) doses were administered before 17:00. Of 100 INR events reported, 23% were above patient-specific target range. 89% INR reports were available before 14:00.

See supplementary file: ds3554.pdf - "Baseline data warfarin project for BMJ"

Design

First intervention: education of prescribers and administrators

In our trust the principal prescribers are junior doctors. Prescribing staff were targeted through focused teaching sessions and posters on wards. Administration is carried out by nursing staff. Nurses were targeted through ward-based patient safety briefs and posters on wards (co-ordinated by charge nurses). Posters highlighted the following key safety issues:

1. Send INR with morning bloods
2. Doctors to dose warfarin in the morning
3. Give warfarin at 14:00
4. Make warfarin patients clear to the team.

Second intervention: warfarin boxes

To intervene at a system level, we introduced "warfarin boxes" (figure 1) in a prominent location on each ward. Warfarin charts were to be placed in the box following the 12:00 nursing drug round to act as a visual prompt for doctors to prescribe in advance of 14:00. Nursing staff could then retrieve all charts from one clearly identified location.

Strategy

PDSA cycle 1: educational intervention

Education about timely warfarin administration focused on prescribing staff (junior doctors) and administration staff (ward staff nurses). No improvement in administration times demonstrated, small reduction in INR reports above patient-specific target range.

PDSA cycle 2: warfarin box

Intervening at a system level, "warfarin boxes" were introduced on test wards. Improvements were seen in daily average administration times and proportion of doses given "on-time", in addition to a reduction in the proportion of INR reports above patient-specific target range.

See supplementary file: ds3556.pdf - "PDSA Cycles 1 and 2 Warfarin QI"

Post-measurement

Baseline: 111 dose administrations recorded. Twenty-seven (24%) were given within an hour of 14:00 prescription, and 40 (36%) were given before 17:00. Number of INRs recorded that were above patient specific target range: 23 (23%).

After first intervention: 78 dose administrations recorded. Ten (13%) were given within an hour of 14:00 prescription. Twenty-one (23%) were given before 17:00. Number of INRs recorded that were above patient specific target range: 19 (19%).

After second intervention: 102 administrations recorded. Fifty (49%) given within an hour of 14:00 prescription, and 64 (63%) given before 17:00. Number of INRs recorded that were above patient specific target range: nine (9%).

Please see the attachment for tabulated results and run chart showing daily and period average times of warfarin administration. Also included is figure 1: warfarin box.

See supplementary file: ds4588.pdf - "Warfarin project upload file"

Lessons and limitations

One of our biggest problems was the failure of the first intervention to have an impact on the time of warfarin administration, although it did have an effect on reducing the variability of dosing times. The first intervention may not have worked because it failed to capture all of the relevant people involved in the process.

We educated junior doctors during junior doctor teaching sessions. However, rotas do not always allow the doctors to attend specific teaching sessions due on-call commitments, study leave or annual leave. Furthermore, our sessions with nursing staff may not have captured all nurses due to shift patterns (not co-ordinated with our education sessions). As the sessions were discrete events, they did

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not promote continuity of the project. It was our experience that education was not sustainable for this project; since doctors rotate every fourth months, the importance of warfarin timing and administration would require regular education sessions (which would be time consuming and labour intensive). Building an intervention like the warfarin box into the regular ward process meant that regular staff could promote a system that was already in place for new doctors when they arrived on the ward.

Regarding the process measure 'INRs outside patient-specific target range': we were unable to identify INRs that were erroneously below target range, meaning that we could only account for INRs erroneously above target range. This was because our data collection did not account for patients who had a subtherapeutic INR secondary to being loaded on warfarin. In this instance a patient being below target range is not necessarily a fault of warfarin administration but is considered part of the normal loading process. INRs below and above target range can cause harm to patients; an INR below target range is associated with an increased clotting risk, while an INR above target range is associated with increased bleeding risk. We therefore missed an opportunity to identify patients at a higher clotting risk due to having INRs below target range.

In some cases we found that delayed administration was due to patients' expectation to receive their warfarin dose at 18:00 (as in the community) and in some cases they even requested to have it at this time despite the nursing staff offering it at 14:00. This acted as a barrier to our intervention.

A few of our project medical wards had only small numbers of patients taking warfarin. In some cases this was due to an increase in patients being on the newer oral anticoagulants. As a result, two out of the seven wards reported difficulty in implementing the warfarin box as it was not needed on a daily basis. The infrequent need for and use of the box made it more difficult to maintain awareness of the issues associated with warfarin prescribing and administration.

Nursing staff often felt it was hard to push the issue of a warfarin prescription to a doctor who may be busy with medically unwell patients. They felt that warfarin prescription was often low on a doctor's list of priorities, especially when they were busy during the middle of the day.

Conclusion

This quality improvement project successfully targeted consistent prescription and administration of warfarin at the trust recommended time of 14:00.

The first intervention focused on education and although it did not produce an improvement in the daily mean time of administration, there was a reduction in the variability of the daily mean times (as demonstrated in the run chart).

Through introduction of a "warfarin box" we were able to improve the number of doses given within an hour of the recommended time

and the daily average of administration time. There was a subsequent decrease in the number of INRs above target range which we envisage may aid in reducing the number of associated adverse events.

References

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Declaration of interests

No interests to declare.

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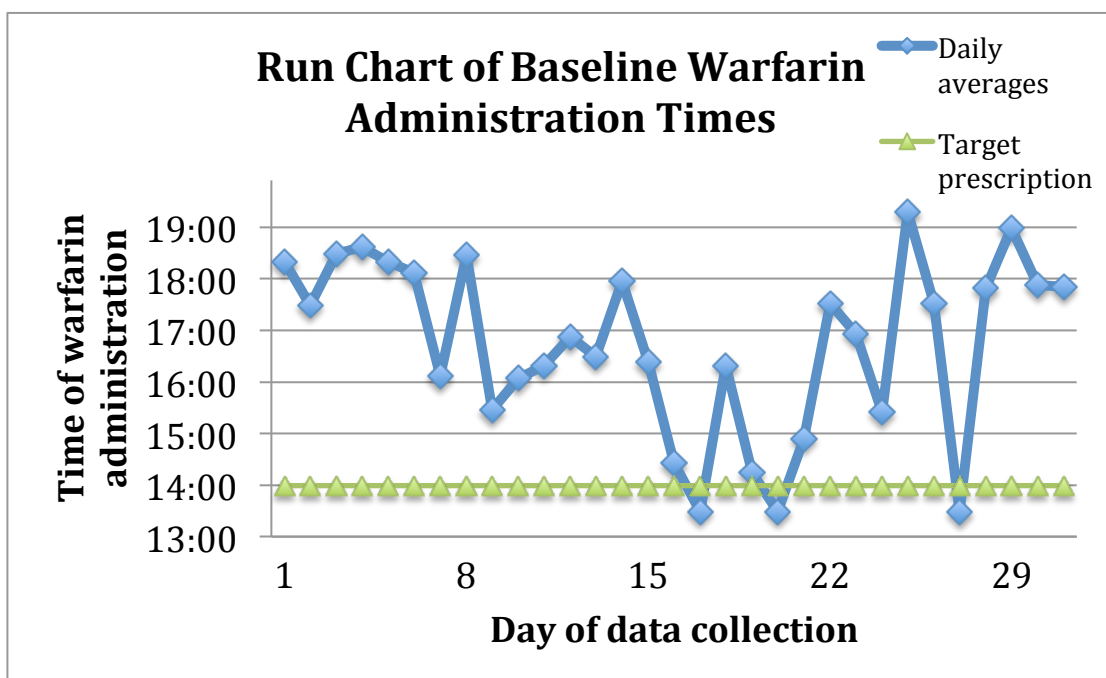
Data collection: baseline

Figure 1: Table of baseline results

	Baseline
Number of doses of warfarin administered	111
Number of doses given at 14:00*	27 (24%)
Number of doses given before 17:00	40 (36%)
Number of INR events reported	100
Number of INR results above patient-specific range	23 (23%)
Number of INR reports available by 14:00	89 (89%)

*As a 'critical' medication, warfarin doses must be administered within 60 minutes of the prescription time. Therefore any doses 13:00-15:00 were included in this category.

Figure 2: Run chart of baseline results



PDSA Cycle 1**Aim:** what are you trying to accomplish?

Our primary aim was to reduce the number of adverse outcomes associated with warfarin prescription and administration. We targeted two system measures:

1. The proportion of warfarin administrations occurring within an hour of the 14:00 prescription (aim to increase)
2. The proportion of INR results outside patient-specific target range (aim to reduce).

Plan: what will your test be?

We will provide education on warfarin prescribing and administering to all staff involved: junior doctors for prescription, nursing staff for administration.
We will aim to engage junior doctors at Trust based teaching sessions.
We will aim to engage nursing staff on each of our test wards.
We will put up posters highlighting the key safety points involved in prescription and administration of warfarin.

Prediction: what do you think will happen as a result of your test?

We hope the proportion of patients administered warfarin within one hour of 14:00 will increase.
We hope the proportion of INR results outside of patient-specific target range will decrease.

Do: what happened when you carried out your test?

Medical and nursing staff were grateful of clarification of the 14:00 dosing.
Staff seemed engaged in pursuing this safety initiative.

Study: how did the results of your test compare with predictions?

There was no increase in the proportion of the patients receiving warfarin within one hour of the 14:00 prescription.
Percentage of INR results above patient-specific target range reduced from 23% to 19%.

Act: how will you change your previous test in light of what you have learned?

We will aim to intervene at a system level to improve timely prescription and administration.
We will talk to medical and nursing staff to discover their perceived delays in this process.

PDSA Cycle 2

Aim: what are you trying to accomplish?

Our primary aim was to reduce the number of adverse outcomes associated with warfarin prescription and administration. We targeted two system measures:

3. The proportion of warfarin administrations occurring within an hour of the 14:00 prescription (aim to increase)
4. The proportion of INR results outside patient-specific target range (aim to reduce).

Plan: what will your test be?

We will produce a Warfarin Box to be placed on each ward in a prominent location (to be decided by the ward team).

The idea is that warfarin charts will be placed in the box after the 12:00 nursing drug round, as a visual prompt for doctors to prescribe in advance of 14:00. Nursing staff can then retrieve all the charts from one clearly identified location.

Prediction: what do you think will happen as a result of your test?

We hope the proportion of patients administered warfarin within one hour of 14:00 will increase. We hope the proportion of INR results outside of patient-specific target range will decrease.

Do: what happened when you carried out your test?

We visited the test wards as a group of three or four (two or three doctors, one lead anticoagulation pharmacist). We discussed with each Ward team (nursing staff and junior doctors) the rationale behind the Warfarin Box, and its intended benefits.

The enthusiasm for the Warfarin Box trial varied; some wards were extremely keen and believed it would 'solve the problem'. Others believed it would be another box taking up space on a busy desk.

Study: how did the results of your test compare with predictions?

Increase in proportion of patients receiving warfarin within one hour of 14:00 (49% vs 24% at baseline).

The run chart demonstrated an improvement in average administration time from 16:56 at baseline to 15:50 after this intervention.

Reduction in proportion of INR results above patient-specific target range (9% vs 23%).

Act: how will you change your previous test in light of what you have learned?

We will go back to the test wards to discuss with nursing staff and junior doctors about the factors they considered key to the improvement demonstrated. Dependent on their views regarding the usefulness of the Warfarin Box, we will consider rolling out this intervention onto other wards.

Table of results: baseline, first intervention, second intervention

	Baseline	Post – first intervention	Post – second intervention
Total number of warfarin doses administered	111	78	102
Number of doses given at 14:00* <i>(figure as a percentage of total doses given)</i>	27 (24%)	10 (13%)	50 (49%)
Number of doses given before 17:00 <i>(figure as a percentage of total doses given)</i>	40 (36%)	21 (23%)	64 (63%)
Number of INR tests reported	100	101	95
INR above range <i>(figure as a percentage of total doses given)</i>	23 (23%)	19 (19%)	9 (9%)
INR report by 14:00 <i>(figure as a percentage of total doses given)</i>	89 (89%)	93 (93%)	92 (97%)

*according to Trust guidelines, a critical medicine must be given within 60 minutes of the prescription time. Any doses given between 13:00-15:00 were therefore included here.

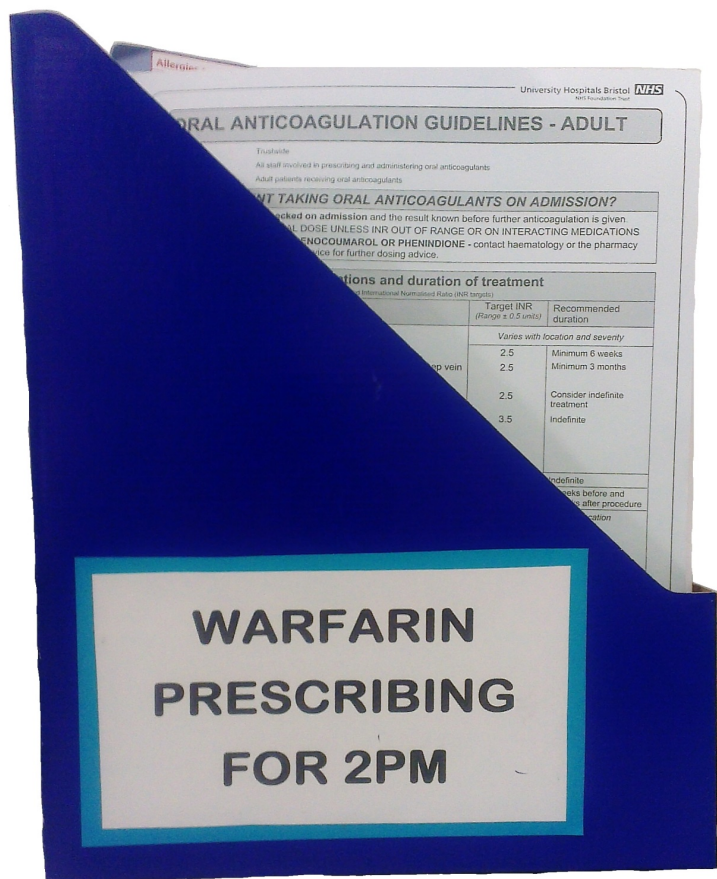


Fig. 1: Warfarin Box

Run Chart of Warfarin Administration Times

