

Usefulness of the traceability of pharmaceutical analysis in patient's medical record



CP-102

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BACKGROUND:

Pharmaceutical analysis (PA) is one of the most important activity of clinical pharmacist as a real benefit on care security. Each day all prescription should be analyzed. But because of lack of time or dedicated staff it is hard to achieve it in good condition. That's why we wanted to show the interest to trace a complete analysis (CA) (that is at least a level 2 analysis) in the patient record (PR) in terms of time saved on the daily analysis (DA) while increasing its quality.

OBJECTIVES:

- To show that traceability of CA saves time on DA
- To determine when all patients will have a CA in their medical record and the time totally won

METHOD:

Only EHPAD* patients are included.

- 1) Achievement of 30 random CA and measurement of time spent on each prescription.
 - 2) DA by 2 different means and measurement of time spent on each prescription:
 - 30 DA of PR without traceability
 - 30 DA of PR including CA traced
 - The sum and the average were calculated on Excel.
- 3) Results extrapolation to estimate when all PR will include a CA traced and the time saved.
 - * Health structure devoted to old and dependant people

The 3 levels of pharmaceutical analysis: (SFPC's definition)

- **Level 1:** Prescription review (posology, contraindications, main drug-drug interactions)
- Level 2: Therapeutic's review (included biology for dose adjustement)
 - **Level 3:** For new or unstable patients (to follow up therapeutic objectives, drug monitoring, compliance...)

RESULTS:

Is daily analysis time shorter when a CA is recorded?

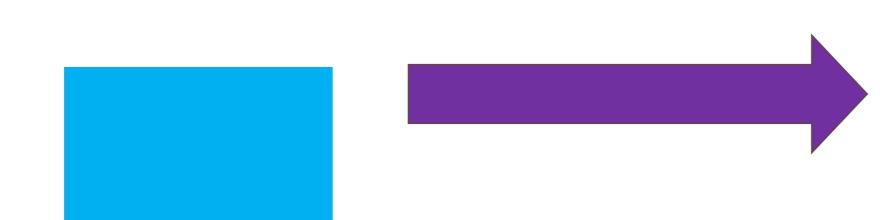
Total: 105,39mins
Average: 3,51mins/prescription

Without traceability:

With traceability:
Total: 34,32mins

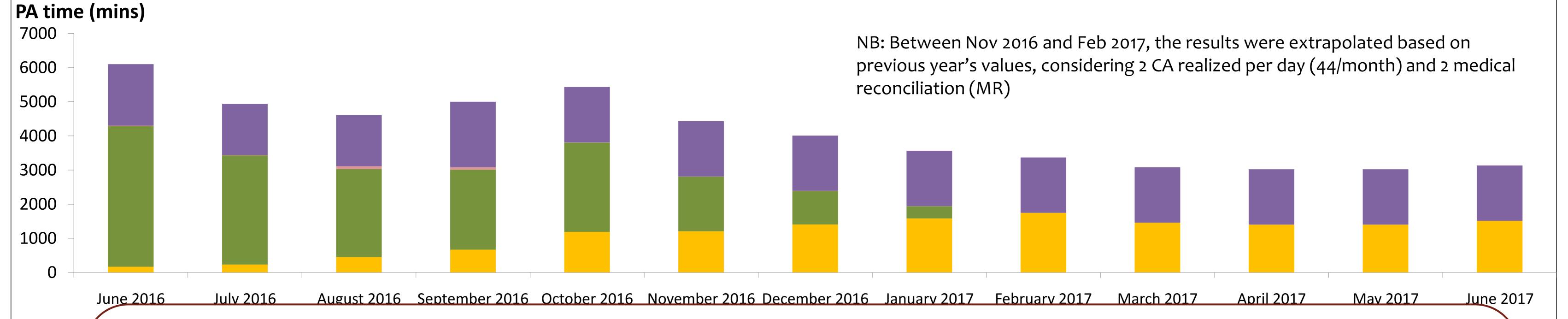
Average: 1,14min/prescription

- For PR without CA: DA requires on average 3,51mins per prescription
- For PR with CA: DA requires on average 1,14mins per prescription
- On average it takes 9,2 minutes [5,27-14,27] to make a CA for prescription of approximatively 10,5 lines [5-19].



Three times less spent on DA when a CA is drawn.

And what about the time when all PR will include a CA or a MR? And the total time saved?



- In yellow, the time spent for **DA when a CA is traced**. It becomes constant from march 2017, when all PR have a CA or a MR.
- In green, the time spent for **DA** when there is no **CA** traced. It decreases from Nov 2016 when 70% of PR have a CA or a MR.
 - In pink (August and Sept), the time not spent on PA, because of a lack of time (holidays time)
 - In purple, the time spent on medical reconciliation (MR) for all new patients (on average 24 per month).

We clearly see that the overall analysis time decreases as soon as CA are included into PR and remains constant when all PR have a CA. The time spent for MR remains stable (on average 1hour including the interview with the doctor and the traceability of the MR)

Finally, the PA time is divided by 2,7 (4118 vs 1511mins) and the total time is divided by 2.

CONCLUSION and DISCUSSION:

This study shows that the traceability of the CA optimizes the DA time and quality. Indeed, the decrease in DA time is explained by the systematic search for the CA, which identifies the particular points to watch. The DA is more focused on critical points of the prescription, which explains its better quality and the time saved.

This time will be used to **develop clinical pharmacy** in the hospital, such as the establishment of output medication reconciliation, medication review with the doctor, pharmaceutical interviews with patients (compliance, therapeutic education, drug information...)

However, it is important to note that the implementation of CA requires **complete computerization** and an easy access to medical information necessary for pharmaceutical validation. The time required to achieve the CA can be planned over several weeks or months.