How to use pharmacy dispensing data to measure adherence and identify nonadherence with oral hypoglycaemic drugs

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Background

- Adherence calculations often based on information of automated databases
- Require a number of methodological choices.
- Methodological framework available to assess adherence with oral hypoglycaemic agents (OHA) by health insurance claims data.
- Pharmacy dispensing data are useful to identify non-adherent (NA) patients for pharmaceutical care and need additional methodological categories.



Objective

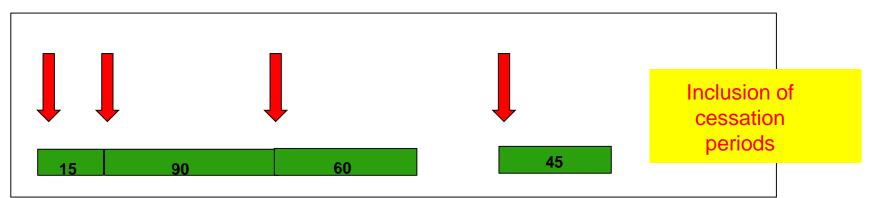
To examine the influences of different parameter values within a framework expanded for the use of dispensing data in estimating OHA adherence.



Data were used from the Dutch Foundation of Pharmaceutical Statistics, SFK.

- Detailed information of dispensed drugs:
 - ATC codes
 - Amount dispensed
 - Prescribed Daily Dose (PDD)
- Patients
 - Anonymous code
 - Sex
 - Year of birth
- Optional: pharmacies with shared population and the same anonymous codes can register as clusters





Adherence between dispensings:

- = <u>days covered by drug use from the first and the one but last dispensing</u> days between the start of the first and the start of the last dispensing
- $= \frac{15 + 90 + 60}{15 + 90 + 90}$
- =165/195 =85%

Adherence until a hypothetical end date:

- = <u>days covered by drug use from all dispensings in the study period</u> days from the first dispensing until the eind of the study period
- = <u>15+90+60+45</u>
 - 15+90+90+160
- =210/355 =59%

Dichotomous measure 80%

'adherent'

'non adherent'



- Patient inclusion by diagnosis, drug prescribing or dispensing of an OHA drug (ATC class A10B)
- 2. Patient type as inclusion criterion: starters, prevalent users or all OHA users
- 3. Minimum number of dispensings as inclusion criterion: at least several dispensings, >1 OHA dispensing during study period
- 4. Length of observation period: some months, 1 year (July 2013 July 2014)
- 5. Calculation of the average medication availability (AMA): from dispensings, calculation by the number of dispensed drugs and the prescribed daily dose
- 6. Time interval under observation: within dispensings or until the end of the study period as fixed date (including periods of drug cessation)
- 7. Adherence measure: assumption of switching between OHA drug classes or concomitant use of several OHA drug classes
- 8. Stockpiling: not or well considered



- 9. Dealing with adherence >100%: no or well truncation
- Number of analyzed medication classes: only one OHA or all OHA classes in use
- 11. Patients with absence periods (e.g. hospital stay, holidays):exclusion
- 12. Threshold to calculate the mean rate of adherent patients (MRAP): MPRAP60/80/90: percentage of patients with an AMA>60%/ >80% / >90%,
- 13. 'Drop in' patients (with only 1 dispensing of any drug in a pharmacy): excluded
- 14. 'Non actual patients' (without a dispensing of any drug in the last four months of the study period): **excluded**
- 15. Insulin users beside OHA use: included
- 16. Source of dispensing data from pharmacy clusters or individual pharmacies



The following adherence measures were calculated for OHA use in the study period between July 2013 and July 2014:

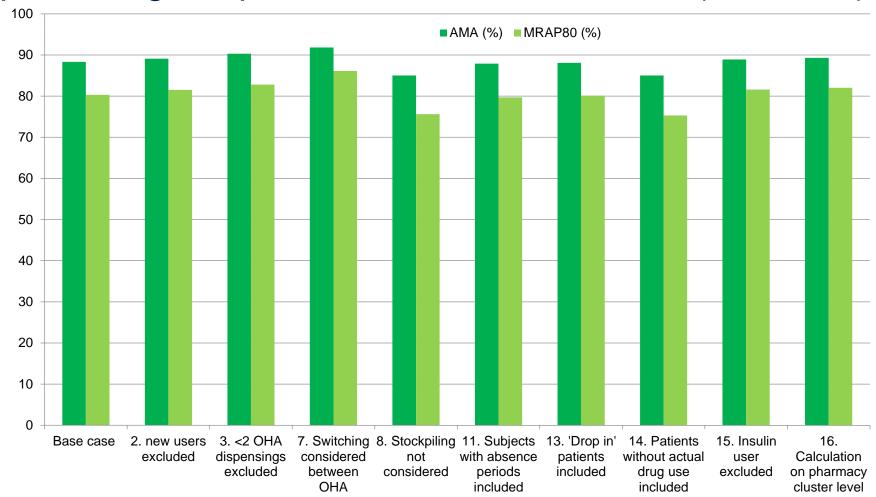
- 1. the average medication availability (AMA)
- 2. a) the percentage of patients with an AMA ≥ 80% (MRAP80)
 - b) the absolute mean number of NA patients per pharmacy.
- Consequences from variation on parameter values for the adherence measures were compared to a base case scenario.



Results

Data were available for 604,500 OHA users in 1,737 (88%) Dutch community pharmacies.

Results: Average medication availability (AMA) and percentage of patients with an AMA>80% (MRAP80)



Base case: AMA 88.3%

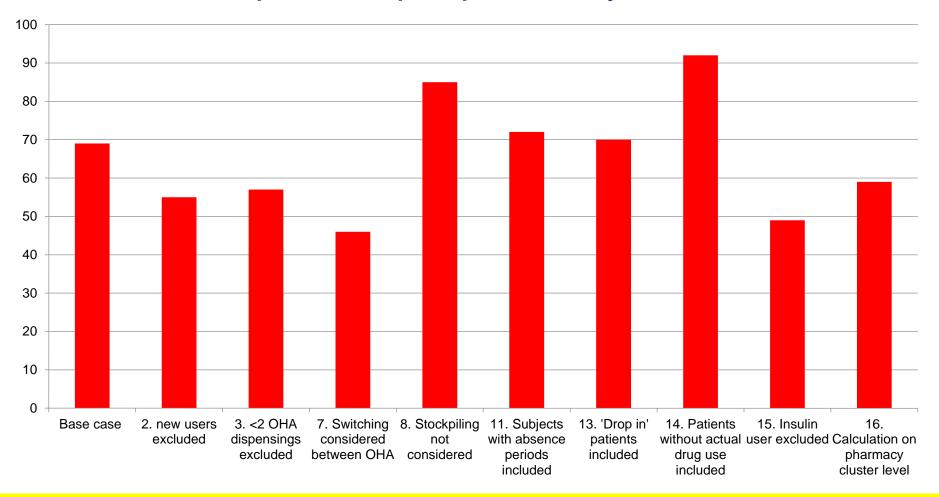
MRAP80: 80,3%

Range of variation

AMA: 85.5% and 91.8%

MRAP80: 75.3% and 86.1%

Results: the mean number of non-adherent patients per pharmacy



Base case:
Mean number of NA patients / pharmacy: 69

Range of variation: 49 - 92

Discussion

- Adherence scores in percentages relative robust to variation of parameter values.
- Due to high number of OHA users substantial differences in absolute numbers of NA patients per pharmacy.
- Validation of patients identified as non-adherent is needed in clinical practice.

Discussion: base case choices

Include:

- all patients, starters as well as prevalent users
- insuline users
- periods of drug cessation (not only use between dispensing periods)

Exclude:

- 'drop in' patients
- 'non actual patients'

Consider concomitant use of several OHA drug classes instead of switching

If possible, use all patients' dispensings, also from several pharmacies.

