Defining medication adherence in individual patients

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Background: The classification of patients as adherent or non-adherent to medications is typically based on an arbitrary threshold for the proportion of prescribed doses taken. Here, we define a patient as pharmacokinetically adherent if the serum drug levels resulting from his/her pattern of medication-taking behavior remained within the therapeutic range.

Methods: We used pharmacokinetic modeling to calculate serum drug levels in patients whose patterns of dosing were recorded by a medication event monitoring system. Medication event monitoring system data were from a previously published study of seven psoriasis patients prescribed 40 mg subcutaneous adalimumab at 14-day intervals for 1 year. Daily serum concentrations of adalimumab were calculated and compared with a known therapeutic threshold.

Results: None of the seven patients took adalimumab precisely every 14 days. Three patients who took adalimumab at intervals of 6–26 days could be classified as pharmacokinetically adherent, because their daily adalimumab serum concentration never fell below the therapeutic threshold. The four other patients, who took adalimumab at intervals of 7–93 days, could be classified as pharmacokinetically non-adherent, because their adalimumab serum concentration fell below the therapeutic threshold on 3.5%–71.3% of days.

Conclusion: Patients with varying patterns of adalimumab dosing could be classified as pharmacokinetically adherent or non-adherent according to whether or not their serum drug concentrations remained within the therapeutic range.

Keywords: pharmacokinetic adherence, drug therapy/utilization, drug administration schedule, patient compliance, adalimumab, pharmacokinetics

Introduction

Medication adherence is typically defined as a ratio of the number of drug doses taken to the number of doses prescribed over a given time period, eg, as the medication possession ratio (MPR).¹ The canonical threshold for adherence is 80%, based on Haynes' definition of adherence to antihypertensive medication as taking $\geq 80\%$ of pills.² This and similar thresholds are arbitrary.³ The use of such arbitrary categories of good and poor adherence is almost invariably unsupported by research documenting the appropriateness of the cutoff for a specific medication class or disease.⁴ In addition, the MPR is a unidimensional concept that does not differentiate between different patterns of non-adherence at any given MPR value. Hence, the MPR does not make use of the detailed information on timing of doses that is provided by a medication event monitoring system (MEMS).

Relationships between patterns of adherence – ie, distributions of time intervals between doses – and drug exposure (ie, serum levels) can be analyzed by pharmacokinetic modeling.⁵ Maclean et al used pharmacokinetic modeling to compute the serum drug levels resulting from different patterns of adherence and compared them to the therapeutic range of the drug.⁶ These authors proposed that patients with

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determining if the final serum concentration measurement for each day was below 4.9 mg/L. If so, then that day was categorized as below threshold. Otherwise, that day was categorized as above threshold. This calculation disregarded data on days 0–6 to account for the time it took to achieve the therapeutic serum concentration after the initial dose.

Results

Serum concentrations of adalimumab are plotted against time in days for patients 1–7 in Figure 1. Three patients (numbers 3, 6, and 7) maintained serum concentrations above the therapeutic threshold of 4.9 mg/L for the entire observation period (Figure 1A). Serum levels fell below the threshold at least some days for patients 1, 2, 4, and 5 (Figure 1B). One patient (number 2) spent 71.3% of the days below the threshold, and two other patients, numbers 1 and 4, fell below the threshold on 23.2% and 31.8% of the

days, respectively (Table 1). Patient number 5 fell below the therapeutic threshold only briefly, for 3.5% of the days at the end of the observation period (Table 1).

Discussion

The method described here – pharmacokinetic modeling of serum drug levels resulting from a patient's particular dosing pattern – leads to three advances. First, it provides an objective means of defining a threshold for adherence – ie, a therapeutic drug serum level – whereas adherence has typically been defined in terms of an arbitrary cutoff – eg, an MPR. Second, it enables the detailed timing of doses provided by MEMS to be interpreted in terms of the clinical effects. Third, it enables identification of patients whose pattern of medication-taking behavior puts them at risk of therapeutic failure, so that they can be targeted for interventions designed to improve their adherence behavior.

Figure 1 Serum concentration of adalimumab over time for patients whose serum levels (A) did not and (B) did drop below the therapeutic threshold.

Notes: Shown are serum levels of adalimumab for each of the seven patients. The timing of each dose for a patient, not specifically indicated in the figure, occurs immediately before each trough point. The bold dashed lines represent the primary therapeutic threshold (4.9 mg/L). The finer dashed lines are alternative cutoffs of 5.4 and 3.6 mg/L.

For example, patients whose MEMS data indicate their serum drug levels have fallen below the therapeutic range could be prompted in real time to take their next dose.

The pharmacokinetic modeling method represents a conceptual change in the definition of adherence. Instead of a unidimensional measure, in which adherence is defined as a point on a linear scale, adherence is now defined in terms of distributions of dosing intervals. We define pharmacokinetic adherence as any pattern of dosing that maintains a serum concentration within the therapeutic range. More broadly, medication adherence has traditionally been defined as whether a patient complies with a physician's instructions. Instead, we define it as whether or not a pattern of patient behavior results in an unfavorable outcome, ie, whether or not the serum drug level remains within the therapeutic range.

In the present study, all of the seven patients sometimes took adalimumab at shorter dosing intervals than the prescribed 14 days. This would tend to lead to higher serum levels than maintained by the 14-day dosing interval. The levels achieved, however, never exceeded 11 mg/L, and no dose-related adverse effects have been observed at much higher levels of adalimumab.^{14,15} Hence, the upper limit of the therapeutic range, which may be highly relevant to some drugs, does not appear to be important for adalimumab.

While MEMS provides extraordinary detailed information about the timing of doses, there is an inherent limitation – the act of opening/closing the container does not necessarily mean that the patient actually consumed the medication. Other potential limitations of this study include the fact that the dosing intervals were from patients with psoriasis and may not be representative of medication-taking behavior in patients with other inflammatory diseases. In addition, pharmacokinetic parameters for adalimumab were from patients with rheumatoid arthritis, and may differ from values in patients with other inflammatory diseases, due to differences in drug metabolism. However, the FDA-approved maintenance dosing regimen is the same across all indications (40 mg every other week), which suggests that the variations in the pharmacokinetic parameters across different diseases are not expected to be greater than those among patients with the same disease. The pharmacokinetic models we employed did not take into account individual patient variables, such as weight and body mass index, which may affect serum drug levels, as this information was not available. These variables can, in principle, be incorporated in future pharmacokinetic modeling studies. Similarly, other factors that might affect serum drug levels, such as co-medications,

were not considered in this analysis but could in principle be addressed by pharmacokinetic modeling. The pharmacokinetic modeling approach does not take into account the loss of clinical response due to development of antibodies to adalimumab, which occurs in some patients on long-term treatment.¹⁸ Antibody testing would be required to detect this. Finally, the validity of the method depends on knowledge of the relationship between serum drug levels and clinical outcomes. We took a serum concentration of adalimumab of 4.9 mg/L as a cutoff value – the trough level below which mucosal healing does not occur in patients with Crohn's disease or ulcerative colitis.¹³ This is similar to the value of 5.05 mg/L, the median concentration reported in Crohn's disease patients who did not achieve clinical remission.¹¹ Other values reported in patients with rheumatoid arthritis are 5.4 mg/dL (the median serum concentration in non-responders)¹² and 3.6 mg/dL (the serum concentration leading to a 50% decrease in CRP)⁹. The application of the latter two cutoffs would not change the assignment of patients except in the case of patient number 5, whose serum levels never fell below 3.7 (Figure 1B). For the remaining patients with serum levels below the therapeutic threshold, use of the cutoff value of 3.6 mg/L would simply decrease the proportion of days with serum concentrations below the threshold. Further research may be required to examine the relationship between serum threshold levels and clinical response.

In conclusion, this pharmacokinetic modeling study showed that patients with varying patterns of adalimumab dosing could be classified according to whether or not their serum drug concentrations remained within the therapeutic range. A dichotomy between pharmacokinetic adherence and non-adherence can be defined by an objective rather than arbitrary threshold. The concept of pharmacokinetic adherence can be applied to any drug for which pharmacokinetic parameters and a therapeutic range are known. This approach could be used to identify individual at-risk patients who could be targeted for intervention. Furthermore, studies of adherence – eg, estimation of a population adherence rate, studies of factors associated with adherence, and trials of interventions to improve adherence – could be based on an objective definition of adherence rather than an arbitrary threshold.

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Author contributions

AM and MES conceived and designed the study, MES performed the analyses, all authors interpreted the data and critically revised the manuscript, and all authors approved the final version.

Disclosure

The authors have no conflicts of interest to disclose in this work.

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