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Pharmacokinetics of oral and intravenous melatonin in healthy volunteers

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Abstract

Background: The aim was to investigate the pharmacokinetics of oral and iv melatonin in healthy volunteers.

Methods: The study was performed as a cohort crossover study. The volunteers received either 10 mg oral melatonin or 10 mg intravenous melatonin on two separate study days. Blood samples were collected at different time points following oral administration and short iv infusion, respectively. Plasma melatonin concentrations were determined by RIA technique. Pharmacokinetic analyses were performed by "the method of residuals" and compartmental analysis. The pharmacokinetic variables: kₐ, t₁/₂ absorption, tₘₐₓ, Cₘₐₓ, t₁/₂ elimination, AUC₀-∞ and bioavailability were determined for oral melatonin. Cₘₐₓ, t₁/₂ elimination, Vₐ and AUC₀-∞ were determined for intravenous melatonin.

Results: Twelve male volunteers completed the study. Baseline melatonin plasma levels did not differ significantly between the study days (P = 0.067). Mean (SD) t₁/₂ absorption of oral melatonin was 6.0 (3.1) min. Mean tₘₐₓ was 40.8 (17.8) min with a median (IQR) Cₘₐₓ of 3550.5 (2500.5–8057.5) pg ml⁻¹. Mean t₁/₂ elimination was 53.7 (7.0) min. Median absolute bioavailability was 2.5 (1.7–4.7) %. Median Cₘₐₓ after short iv infusion of melatonin was 389,875.0 (174,775.0–440,362.5) pg ml⁻¹. Mean t₁/₂ elimination was 39.4 (3.6) min, mean Vₐ 1.2 (0.6) l kg⁻¹ and mean CL 0.0218 (0.0102) l min⁻¹ kg⁻¹.

Conclusions: This cohort crossover study estimated pharmacokinetics of oral and iv melatonin, respectively in healthy volunteers. Bioavailability of oral melatonin was only 3 %.

Trial registration: Eudra-CT number: 2013-000205-23 (initial registration 27.03.2013). Clinicaltrials.gov Identifier: NCT01923974 (initial registration 08.08.2013).

Keywords: Bioavailability, Intravenous, Melatonin, Oral, Pharmacokinetic

Background

Exogenous melatonin is being increasingly employed as treatment for various medical and surgical diseases [1, 2]. Furthermore, a recent study, administering intravenous (iv) melatonin has documented reduced cardiac morbidity and markers of myocardial ischemia following elective abdominal aortic aneurism repair [3]. Despite its widespread clinical use, the pharmacokinetic properties of exogenous melatonin still need to be established further [4]. A limited number of experimental studies in healthy volunteers have performed direct comparisons of the pharmacokinetics of oral and iv melatonin [5, 6]. The studies differed in number of investigated subjects, dosages, methods and pharmacokinetic analyses [5, 6]. Accordingly, the pharmacokinetic variables varied extensively between the studies [5, 6]. In order to achieve an optimized clinical efficacy of melatonin, further investigation of the pharmacokinetics is clearly needed.

The aim of the study was to investigate the pharmacokinetics of oral and iv melatonin in a cohort of healthy volunteers.

Methods

Approvals were obtained from the Capital Region’s Committee on Health Research Ethics (Protocol number: H-4-2013-013), the Danish Health and Medicines
concentrations was performed by radioimmunoassay (RIA)-technique (Melatonin Direct RIA, DIAsource, Louvaine-La-Neuve, Belgium). Precision of the RIA kit: intra-assay coefficient of variation (CV) = 9.8–13.4 %, inter-assay CV = 8.0–13.3 %. The limit of detection was 2.3 pg ml⁻¹. Linearity of the kit ranged between 8.5–529.0 pg ml⁻¹. If plasma concentrations exceeded detection range of the kit, plasma samples were diluted according to manufacturer’s guidelines. All plasma samples were analysed in duplicate, and the mean value was reported.

Statistical and pharmacokinetic analyses
Normality of data was assessed by visual inspection of residual plots and histograms. Parametric or non-parametric tests were applied according to the distribution of data. Correspondingly, data are presented as mean (SD) or median (IQR), unless stated otherwise. A P-value < 0.05 is considered statistically significant. Data were analysed using IBM SPSS Statistics for Windows version 22.0 (IBM Corp., Armonk, NY, USA) and Graph Pad Prism version 6.0 (Graph Pad Software Inc., La Jolla, CA, USA).

The baseline melatonin plasma concentrations of each study day were compared using a paired sample T-test. Baseline levels were not subtracted from post-treatment (oral or iv) levels.

Pharmacokinetic analyses of oral and iv melatonin were performed separately.

Oral melatonin
Time to maximal concentrations (tmax) and maximal plasma concentrations (Cmax) were assessed directly at the relevant time points. The pharmacokinetic variables: absorption constant (ka), absorption half-life (t1/2 absorption), elimination rate constant (ke) and elimination half-life (t1/2 elimination) were estimated by “the method of residuals” [8]. Areas-under-the-curve (AUC) of plasma concentrations were calculated by applying the trapezoidal rule [9]. AUC0–∞ was estimated as AUC0–420 min + (C420 min / ke). Bioavailability was calculated as (AUC0–∞ oral / AUC0–∞ iv) x 100.

Intravenous melatonin
Cmax was assessed directly at the time point, 0 min after short iv infusion. Pharmacokinetic variables were calculated by compartmental analysis [10]. The pharmacokinetic variables: t1/2 elimination, volume of distribution (Vd) and clearance (CL) were estimated from individual linear regression lines of log-transformed (natural logarithm) plasma concentrations. Following standard equations were applied: t1/2 elimination = ln (2) / ke, Vd = dose / Ce min, CL = ke x Vd. “Goodness of fit” of the individual linear regression lines was assessed by the coefficient of determination, R². AUC0–∞ iv was estimated, as described above.
Results
Twelve male volunteers were included and completed the study. Mean age and body mass index (BMI) were 27.1 (5.2) years and 23.2 (2.7) kg m\(^{-2}\), respectively. Baseline melatonin plasma concentrations did not differ significantly between the study days (before oral melatonin = 27.3 (13.5) pg ml\(^{-1}\); before intravenous melatonin = 18.3 (12.3) pg ml\(^{-1}\) \(P = 0.067\).

The pharmacokinetic variables of oral and iv melatonin are presented in Tables 1 and 2.

Oral melatonin
Oral melatonin demonstrated first-order absorption and elimination kinetics. Mean \(k_o\) was 0.2 (0.1) min\(^{-1}\), and mean \(t_{1/2}^a\) of oral melatonin was 6.0 (3.1) min (Fig. 1). Mean \(t_{max}\) was 40.8 (17.8) min with a median (IQR) \(C_{max}\) of 3550.5 (2500.5–8057.5) pg ml\(^{-1}\). Mean \(t_{1/2}^e\) was 53.7 (7.0) min, \(AUC_{0–\infty}^o\) oral 281,538.3 (232,696.1–546,285.4) pg ml\(^{-1}\) min and median absolute bioavailability was 2.5 (1.7–4.7) %.

Intravenous melatonin
The pharmacokinetic profiles of iv melatonin demonstrated first-order elimination kinetics (Fig. 2). Median \(C_{max}\) after iv bolus injection of 10 mg melatonin was 389,875.0 (174,775.0–440,362.5) pg ml\(^{-1}\). Mean \(t_{1/2}^e\) was 39.4 (3.6) min, mean \(V_d\) 1.2 (0.6) l kg\(^{-1}\) and mean CL 0.0218 (0.0102) l min\(^{-1}\) kg\(^{-1}\). Median \(R^2\) was 0.96 (0.93–0.97). Median \(AUC_{0–\infty}^i\) was 14,179,767.6 (7,063,347.4–18,964,804.0) pg ml\(^{-1}\) min.

Discussion
This cohort crossover study demonstrated a \(t_{max}\) of 41 min following oral administration. \(C_{max}\) and \(AUC\) varied extensively between volunteers in both administration routes. Elimination half-lives were 54 min and 39 min, respectively. Bioavailability of oral melatonin was only 3 %, but demonstrated substantial inter-individual differences.

Oral melatonin
Oral melatonin was absorbed by first-order kinetics, which has previously been demonstrated in doses up to 80 mg [8]. The short \(t_{1/2}^a\) of 6 min, corroborate studies, applying similar oral drug formulations [8]. Accordingly, our \(t_{max}\) value of 41 min is in agreement with other studies, documenting values ranging from 30 to 60 min [6, 11]. Oral administration of exogenous melatonin, approximately 45 min before intended onset of clinical effects therefore seems reasonable, assuming that clinical efficacy coincides with \(t_{max}\) values [12]. Oral administration was associated with extremely variable \(C_{max}\) and \(AUC_{0–\infty}^o\) oral values, which has been described previously [5]. The inter-individual variations are apparently caused by differences in absorption, distribution, metabolism or excretion of the drug, but the exact causes and clinical implications remain unestablished so far [5]. Previous studies demonstrate \(t_{1/2}^e\) values ranging from 46 to 65 min in oral doses from 0.5 to 6 mg [5, 6, 11], which correlates with our findings of 54 min. Our data demonstrated a very low absolute bioavailability of 3 %, albeit with a substantial inter-individual variability. Previous experimental studies have documented higher values ranging between 9 and 33 %, although with comparable inter-individual variability [5, 6, 12]. It is well established in both animal- and human studies that the low bioavailability results from an extensive hepatic first pass metabolism [5]. Similarly, it is also clear that these findings may mandate future dose regulations between different administration routes. However, a general lack of experimental- and clinical studies correlating melatonin plasma concentration levels and clinical effects still remains, and further knowledge is needed, preferably by in-depth pharmacokinetic-pharmacodynamic modelling.

Intravenous melatonin
Previous studies investigating iv administration of melatonin have also demonstrated first-order eliminations kinetics [10], as observed in our study. As with oral melatonin, iv administrations displayed extensive variations in \(C_{max}\) and \(AUC_{0–\infty}^i\) values, which is in accordance with previous studies [6]. Other studies also documented \(t_{1/2}^e\) values ranging between 28 and 60 min in iv doses from 0.005 mg to 2 mg [6, 10, 13], which corresponds to the 39 min, demonstrated in the present study. Several studies confirm that elimination rates of iv melatonin (and oral melatonin) are not related to the administered dose. Similarly, previous studies document CL values of 0.013 l min\(^{-1}\) kg\(^{-1}\) (weight-corrected) [13] and 0.027 l min\(^{-1}\) kg\(^{-1}\) [10], which correspond well to our findings of 0.022 l min\(^{-1}\) kg\(^{-1}\). Cavallo and colleagues also documented a \(V_d\) of 1.8 l kg\(^{-1}\) [10], which is comparable with a value of 1.2 l kg\(^{-1}\), demonstrated in our study.

<table>
<thead>
<tr>
<th>(t_{1/2}^a) absorption min</th>
<th>(t_{max}) min</th>
<th>(C_{max}) pg ml(^{-1})</th>
<th>(t_{1/2}^e) elimination min</th>
<th>(AUC_{0–\infty}^o) oral pg ml(^{-1}) min</th>
<th>(f) %</th>
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<tbody>
<tr>
<td>6.0 (3.1)</td>
<td>40.8 (17.8)</td>
<td>3550.5 (2500.5–8057.5)</td>
<td>53.7 (7.0)</td>
<td>281,538.3 (232,696.1–546,285.4)</td>
<td>2.5 (1.7–4.7)</td>
</tr>
</tbody>
</table>

Absorption half-life, time to maximal concentration and elimination half-life data are presented as mean (SD). Maximal concentration, area-under-the-curve and bioavailability data are presented as median (IQR)
Strengths

Our study is the first to perform direct comparisons of pharmacokinetics of oral and iv melatonin in doses routinely administered perioperatively (approximately 10 mg) [2]. The study was performed as a crossover study to reduce the effect of the inter-individual variability on pharmacokinetic data. Our experimental setup included multiple blood samples for a detailed description of both absorption and elimination phases in both administration routes. Our study also included standard pharmacokinetic methods, such as “the method of residuals” and compartmental analysis [8, 10]. In addition, we chose to include the coefficient of determination ($R^2$) to document the “goodness of fit” of the individual linear regression lines in the compartmental analysis. Our data demonstrated a $R^2$ value of 0.96, indicating a high degree of “fit” of the first-order pharmacokinetic model, and, hence, a considerable accuracy of the derived pharmacokinetic variables.

Limitations

First, this study only included healthy male volunteers in an experimental setup. Hence, a potential gender difference in pharmacokinetic variables may exist. Furthermore, previous experimental studies indicate that the pharmacokinetics of melatonin is affected by age [10] and external factors, such as caffeine intake [14] cigarette smoking [15] and the use of oral contraceptives [16]. Also, a low number of clinical studies have demonstrated altered pharmacokinetic variables of melatonin [17–19] in e.g. critically ill patients [17, 18]. Interestingly, most other patient groups, e.g. surgical patients, still remain to be investigated. Comorbidity and drug interactions may change the pharmacokinetics of melatonin, potentially altering clinical efficacy of the drug [20].

Second, oral and iv study sessions were separated by 3 to 9 months for each volunteer. These time periods may theoretically have affected the comparability of individual pharmacokinetic variables, despite the crossover design. It, however, seems unlikely, as all volunteers were healthy young males in stable physical conditions.

Third, the very low bioavailability of oral melatonin documented in our study may indicate a deficient absorption of the drug in our setup. The volunteers were allowed 5 cl of tap water to facilitate intake of oral melatonin. Hence, it can be discussed, if the restricted liquid volume, despite saliva and gastric/intestinal fluid secretions was sufficient to dissolve and present the ingested melatonin to the small intestine, where absorption is mainly achieved. We, however, chose this amount of water to standardize the experimental conditions and to imitate a clinical premedication scenario [2, 7]. Also, we administered an easily absorbable gelatine capsule in order to optimize dissolution of the drug. Finally, comparable $t_{max}$ values between the volunteers were demonstrated, suggesting that an impeded absorption is rather unlikely.

Conclusions

This crossover cohort study investigated the pharmacokinetics of oral and intravenous melatonin in healthy male volunteers. Oral melatonin was rapidly absorbed, and $T_{max}$ was achieved after 41 min. $C_{max}$ and $AUC$...
varied extensively between volunteers. Elimination half-lives following oral and intravenous melatonin administration was 54 min and 39 min, respectively. The bioavailability of oral melatonin was only 3%, but a considerable variability between the volunteers was noted.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
UPHA participated in the design of the study, collected data, performed data-analysis, and drafted manuscript. MUW, MMR, JR and IG participated in the design of the study, and helped to draft the manuscript. NHG and HF collected data, and helped to draft the manuscript. All authors have read and approved the final manuscript.

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References