Pharmacokinetic Evaluation of Administration of Losartan with Aspirin in Healthy Volunteers

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Abstract

Losartan and aspirin are often used concomitantly in patients with heart failure, ischemic heart disease and hypertension.

Objectives: To investigate whether aspirin co-administration affects losartan bioavailability.

Methods: 1) Twenty-four healthy volunteers from both sexes were recruited. Volunteers received a single 50 mg losartan with or without a 81 mg aspirin tablet. Blood samples were obtained at 0.25, 0.5, 0.75, 1, 1.25, 1.5, 1.75, 2, 2.33, 2.67, 3, 3.5, 4, 4.5, 5, 6, 8, 10, 12 hour post-dosing. The concentrations of losartan were analyzed by LC-MS-MS. Clearance (Cl) and T1/2 were used to evaluate a possible drug-drug interaction. Cmax and AUC0-t were used to evaluate whether co-administration interferes on the bioavailability process.

Results: From the losartan plasma concentrations vs. time curves the following pharmacokinetic parameters were obtained: ASC0-t hours, AUCinf, Cmax, Cl, Vd, T1/2, Ke and T1/2. No significant differences were observed in T1/2 (p-value = 0.431), Cl (p-value = 0.554), AUC0-t, hours (p-value = 0.590), Cmax (p-value = 0.987) and Vd (p-value = 0.647).

Conclusions: Since there is no significant difference in losartan bioavailability and elimination when co-administered with aspirin, we conclude that there is no pharmacokinetic interaction between both drugs. The finding is important since it reassures the safe use of combining AAS to losartan treatment.

Keywords: Mass spectrometry – interaction; LC-MS

Introduction

Losartan is a non-peptide angiotensin II receptor antagonist used as an antihypertensive agent, that block the vasoconstrictor and aldosterone-secreting effects of angiotensin II by type AT1 receptor blockage [1,2]. Losartan may reduce cardiovascular events more significantly than β-blockers in patients with hypertension and left ventricular hypertrophy [3]. Losartan decreased platelet aggregation by a thromboxane A2-dependent mechanism [4]. The losartan pharmacokinetic parameters after single oral administration of 50 mg to healthy volunteers were between 182 - 252 ng/mL, 0.50 – 2.66 h, 323 - 480 ng*h/mL, 1.88 - 2.84 h for Cmax, tmax, AUC0-t, t1/2, respectively [1,2].

Aspirin (acetylsalicylic acid) is an irreversible inhibitor of the cyclooxygenase (COX)-1 and -2 isoenzymes [5] that is used to reduce arterial thrombosis and primary prevention of myocardial infarction and stroke in low daily dose (81 mg) [6-10]. These effects are explained by decreased production of thromboxane A2 (potent stimulant of platelet aggregation and vasoconstriction), and by adequate production of prostaglandin (vasodilator and platelet inhibitor) with COX-2 activity intact [5]. Low doses of aspirin (40–50 mg/day) maintain complete blockade of platelet COX-1 [11].

Thus, association of losartan and aspirin is the treatment of hypertension of patients who have risk of developing thromboembolic disease is often observed [12-14]. Aspirin exerts no significant effect on blood pressure in essential hypertensives taking losartan [14-16], however it may attenuate the action of ACE inhibitors by reducing prostaglandin synthesis [15]. Losartan antihypertensive therapy combined with aspirin was more effective than an atenolol-based treatment with aspirin in reducing the primary composite end point of cardiovascular morbidity and mortality [12]. Interestingly, statistical analysis showed interaction between losartan and aspirin [12]. The conclusion was that further studies were needed to clarify a pharmacologic interaction or a selection by aspirin use of patients more likely to respond to losartan treatment [12].

This study is the first pharmacokinetic evaluation between losartan in combination with aspirin. The study aims to compare the pharmacokinetic (absorption and metabolism/elimination process) of single-dose losartan (50 mg) in the presence or absence of aspirin (81 mg) in healthy volunteers to determine whether co-administration affected the pharmacokinetic profiles of the losartan.

Methods

Pharmacokinetic study

The study began with 24 volunteers and finished with 21 volunteers.
Three volunteers dropped out of the study for personal reasons. The male group was composed of 12 volunteers (32.5 years ± 8.9 years, mean ± s.d.; range 23–52 years), height between 167 and 185 cm (174 ± 7 cm), weighing between 66.6 and 82.8 kg (76.8 ± 5.1 kg). The female group was also composed of 12 volunteers (27.3 years ± 9.2 years; range: 18–52 years), height between 155 and 169 cm (163 ± 5 cm), weighing between 51.0 and 80.0 kg (63.5 ± 9.2 kg).

All subjects gave written informed consent and the State University of Campinas (UNICAMP) ethics committee approved the clinical protocol. All volunteers were healthy as assessed by physical examination, electrocardiogram, and the following laboratory tests: blood glucose, urea, uric acid, creatinine, alkaline phosphatase, aspartate and alanine aminotransferases, gamma-glutamyl transferase, total bilirubin, albumin and total protein, trygliceride, total cholesterol, hemoglobin, hematocrit, erythrocyte sedimentation rate, total and differential white cell and platelet counts, and routine urinalysis. All female volunteers were negative on a pregnancy test. All subjects were negative for human deficiency, hepatitis B (except for serological scar) and hepatitis C virus.

The study was conducted in an open, randomized, 2-period crossover balanced design with a 1-week washout period between doses. During each period, the volunteers were hospitalized at 6:00 p.m. having a normal evening meal, and after an overnight fast starting at 10:00 p.m., they received at 7:00 a.m. a single 50 mg potassic losartan or 50 mg potassic losartan + 81 mg aspirin immediate release tablet. Water (200 mL) was given immediately after drug administration. Blood samples (7 mL) from a suitable antecubital vein were collected at 0.25, 0.5, 0.75, 1, 1.25, 1.5, 1.75, 2, 2.33, 2.67, 3, 3.5, 4, 4.5, 5, 6, 8, 10, 12 hour post-dosing. The blood samples were centrifuged at approximately 2000 x g for 1 min at 4°C, and the decanted plasma stored at -20°C until analysis. All volunteers then fasted for 2 h following the drug administration, after which a standard breakfast was consumed. A standard lunch and an evening meal were provided 4 h and 10 h after dosing, respectively. No other food was permitted during the in-house period. Liquid consumption was permitted ad libitum after lunch but xanthine-containing drinks including tea, coffee, and cola were prohibited. All subjects were requested to stay in the clinical unit for a 12-h period after drug administration.

Systolic and diastolic arterial pressure (measured non-invasively with a sphygmomanometer), heart rate and temperature were recorded just before and at 4, 8 and 12 sampling collection.

### Formulations

The following formulations were employed: losartan (Aradois®) immediate release 50 mg tablets (Biolab Sanus Farmacêutica Ltda., Brazil; lot N° 2120205, expiration date 12/2014) and aspirin (Eclasil®) 81 mg immediate release tablets (Biolab Sanus Farmacêutica Ltda., Brazil; lot N° 2120071, expiration date 12/2014).

### Pharmacokinetics and statistical analysis

The first-order terminal elimination rate constant (ke) was estimated by linear regression from the points describing the elimination phase on a log-linear plot. Half-life (t 1/2) was derived from this rate constant (t 1/2 = ln (2)/ke) and the time taken to achieve this concentration (T max) were obtained directly from the curves. The areas under the losartan plasma concentration vs. time curves from 0-t is the last detectable concentration (AUC0-t) and vs. time curves from 0-8h (AUC0-8) were calculated by applying the linear-log trapezoid rule. Extrapolation of these areas to infinity (AUC0-inf) was done by adding the value C last/ke to the calculated AUC0-t (where C last = the last detectable concentration). The clearance (CL) was calculated by Dose/AUC0-t. The volume of distribution (Vd) was calculated by CI/Ke. The T1/2, CL, ASC0-8 h, Cmax and Vd data for the two formulations were analyzed by paired T-Test, as proposed by the US Food and Drug Administration [17-19]. The software used included WinNonlin Professional Network Edition (Pharsight v. 5.3), Microsoft Excel (v. 7.0), GraphPad Prism (v. 3.02) and GraphPad InStat (v. 3.06).

### Drug analysis

The extraction was performed by vortex-mixing 200 µL of each plasma sample, placed in glass tubes followed by the L.S. (50 µL of 2000 ng/mL) and the samples vortex-mixed for 5s. Formic acid (88%) was added (20 µL) to all tubes and the samples were vortex-mixed for 5s. Ethyl acetate was then added (3 mL) to all tubes and performed the extraction by vortex-mixing for 50 s. The samples were centrifuged at 2000 x g for 2 min. The upper organic phase was transferred to another set of clean glass tubes and evaporated until dry under N2 at 50°C. The dry residues were dissolved with 300 µL of acetonitrile/water (50/50; v/v) by vortex-mixing for 10 s. The solution was transferred to 96-well plates using automatic pipettes with disposable plastic tips.

### Calibration standards and quality control

Stock solutions of losartan and internal standard (valsartan) were prepared in acetonitrile-water (50:50 v/v) at concentrations of 1 mg/mL, respectively. Calibration curves of losartan were prepared by spiking blank plasma at concentrations of 2, 5, 20, 200, 500, 1000, 1500 and 2000 ng/mL and the analysis was carried out in duplicate for each concentration. The quality control samples were prepared in blank plasma at concentrations of 6, 80, 800 and 1600 ng/mL (QCA, QCBI, QCBC, and QC, respectively). The spiked plasma samples (standards and quality controls) were extracted from each analytical batch along with the unknown samples.

### Chromatographic conditions

An aliquot (10 µL) of each plasma extract was injected into an Alltech Prevail C18 5 µm analytical column (150 mm x 4.6 mm i.d.) operating at 50°C. The compounds were eluted by pumping the mobile phase (acetonitrile and water [70/30; v/v] containing 10 mM formic acid) at a flow rate of 1.2 mL/min diluted approximately 1:12. Under these conditions, typical standard retention times were 2.0 min for losartan and 2.7 min for valsartan, and back-pressure values of approximately 50 Bar were observed. The temperature of the autosampler was kept at 8°C and the run-time was 3.75 min.

### Mass-spectrometric conditions

The mass spectrometer (Micromass model Quattro Ultima) was equipped with an electrospray ion source running in positive mode (ES+), was set up in Multiple Reaction Monitoring (MRM) for the transitions 423.30 > 207.15, and 436.50 > 291.00 for losartan and valsartan, respectively. The dwell time, cone voltage and the collision energy were 0.35 sec, 15 V and 22 eV for losartan, 0.35 sec, 20 V and 15 eV for valsartan, respectively. The mass spectrometer (Micromass model Quattro Ultima) was equipped with an electrospray ion source running in positive mode (ES+). The dwell time, cone voltage and the collision energy were 0.35 sec, 15 V and 22 eV for losartan. Data acquisition and analysis were performed using the software Mass Lynx (v. 3.5).

The analysis was conducted and the method was fully validated in accordance with the study protocol and ANVISA (National Health Surveillance Agency) Guidelines for Analytical Methods Validation, including parameters such as sensitivity, specificity, linearity, accuracy, precision and reproducibility.

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The stability of the analytes and internal standard were evaluated in the biological matrix (human plasma) under distinct timing and temperature conditions in two concentrations: 36 days long term at -20°C, 6:55 hours short term at room temperature, 3 freeze and thaw cycles and 70 hours post processing at 8°C. The stability of the compounds in master solutions was also evaluated. The assessment showed that compounds were stable during the time and conditions of analysis.

Results

The formulation was well tolerated at the administered doses and no significant adverse reactions were observed or reported. A total of three adverse events were reported during the study, two of them were considered probably related to the administration (vomit [losartan phase] and headache [losartan phase]). The other adverse event was backache [losartan phase]. The biochemical parameters presented no clinically relevant alterations.

No endogenous peak was observed in the mass chromatogram of blank plasma. Figure 1 shows a chromatogram for the standard LOQ sample in which the retention times for losartan and the internal standard was 2.00 ± 0.3 min and 2.70 ± 0.3 min, respectively. The calibration curves (for losartan) showed good linearity ($r^2 > 0.999396$) within the range 2.00 to 2000 ng/mL for analyte. The limit of quantification (LOQ), defined as the lowest concentration at which both the precision and accuracy were <20%, was 12.0 ng/mL. The within- and between-run precision and accuracy for the LOQ and QCs are summarized in Table 1.

Mean pharmacokinetic parameters of losartan obtained from 21 volunteers after administration of losartan (50 mg) in the presence or absence of aspirin (81 mg) are presented in Figure 2 and Table 2. The paired T test for pharmacokinetic parameters ($T_{1/2}$, CL, AUC$_{0-8}$ horas, C$_{max}$ and Vd) of losartan after administration of losartan (50 mg) in the presence or absence of aspirin (81 mg) are presented in Table 3.
Oral administration of losartan and aspirin was considered safe. After oral administration of losartan (50 mg) or losartan (50 mg) + aspirin (81 mg), the observed pharmacokinetic parameters of losartan of both oral administrations were similar to those reported in the literature for losartan [1,2].

As shown in Figure 3, the systolic and diastolic blood pressures and heart rates measurements revealed no significant alteration after oral administration of losartan (50 mg) or losartan (50 mg) + aspirin (81 mg), even after 12 hours the administration (p = 0.1786). Indeed no significant change in blood pressure was reported when losartan in addition of either 81 or 325 mg aspirin were administrated [14-16].

For pharmacokinetic interaction between the drugs, it is necessary a significant change in the absorption process or in the process of metabolism / elimination. The area under the curve (AUC) until 08 hours after the administration reflects more accurately the amount of drug absorbed, since after this time the dosage form is in the

Table 2: Mean pharmacokinetic parameters of losartan obtained from 21 volunteers after administration of 50 mg losartan in presence or absence of aspirin (81 mg).

<table>
<thead>
<tr>
<th>Variable</th>
<th>Unit</th>
<th>N</th>
<th>Mean</th>
<th>SD</th>
<th>Min</th>
<th>Median</th>
<th>Max</th>
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<tbody>
<tr>
<td>AUC % extrap (%</td>
<td></td>
<td>21</td>
<td>2.46</td>
<td>0.98</td>
<td>1.07</td>
<td>2.06</td>
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<td>39.77</td>
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<td>AUC&lt;sub&gt;0-8h&lt;/sub&gt; ([ng*hr]/mL)</td>
<td>21</td>
<td>361.55</td>
<td>105.08</td>
<td>223.79</td>
<td>378.30</td>
<td>630.95</td>
<td>29.06</td>
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<tr>
<td>AUC&lt;sub&gt;inf&lt;/sub&gt; ([ng*hr]/mL)</td>
<td>21</td>
<td>382.76</td>
<td>112.02</td>
<td>239.09</td>
<td>403.35</td>
<td>665.63</td>
<td>29.27</td>
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<tr>
<td>ASC&lt;sub&gt;LAST&lt;/sub&gt; ([ng*hr]/mL)</td>
<td>21</td>
<td>373.67</td>
<td>110.66</td>
<td>231.19</td>
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<td>655.65</td>
<td>29.62</td>
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<td>C&lt;sub&gt;LAST&lt;/sub&gt; (ng/mL)</td>
<td>21</td>
<td>2.94</td>
<td>0.64</td>
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<td>3.00</td>
<td>4.10</td>
<td>21.89</td>
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<td>Cmax (ng/mL)</td>
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<td>21</td>
<td>211.05</td>
<td>100.61</td>
<td>106.00</td>
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<td>Ke (1/hr)</td>
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<td>0.35</td>
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<td>0.17</td>
<td>0.34</td>
<td>0.47</td>
<td>21.04</td>
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<td>T1/2 (hr)</td>
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<td>2.10</td>
<td>0.58</td>
<td>1.47</td>
<td>2.01</td>
<td>4.08</td>
<td>27.83</td>
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<tr>
<td>T&lt;sub&gt;LAST&lt;/sub&gt; (hr)</td>
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<td>8.00</td>
<td>10.00</td>
<td>12.00</td>
<td>14.61</td>
<td></td>
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<tr>
<td>Tmax (hr)</td>
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<td>21</td>
<td>1.03</td>
<td>0.71</td>
<td>0.50</td>
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<td>2.67</td>
<td>68.81</td>
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<td>Vd (L)</td>
<td></td>
<td>21</td>
<td>417.54</td>
<td>122.19</td>
<td>241.92</td>
<td>401.76</td>
<td>678.22</td>
<td>29.27</td>
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<tr>
<td>Cl (L/h)</td>
<td></td>
<td>21</td>
<td>141.15</td>
<td>39.57</td>
<td>75.12</td>
<td>123.96</td>
<td>209.13</td>
<td>28.03</td>
</tr>
</tbody>
</table>

Table 2: Mean pharmacokinetic parameters of losartan obtained from 21 volunteers after administration of 50 mg losartan in presence or absence of aspirin (81 mg).
large intestine where drug absorption does not occur. The process of metabolism-elimination is responsible for clearance of the drug, which determines the half-life of the drug [23].

More studies were needed to clarify a pharmacologic interaction or a selection by aspirin use of patients more likely to respond to losartan treatment [12]. As noted in the Table 3, there is no significant difference in the area under the curve (AUC) until 8 hours of losartan, as there is no significant difference in the parameter for the clearance (clearance itself, and the half-life). The statistical test used was the paired t test considered by the FDA [18,19] as a test sensitive for identification of pharmacokinetic interactions (in others words, carries the risk of detecting pharmacokinetic interactions that are not clinically relevant).

**Conclusion**

Since there is no significant difference in losartan bioavailability and elimination when co-administered with aspirin, we conclude

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**Table 3:** Paired T test for $T_{1/2}$, CL, AUC$_{0-8}$, Cmax and Vd after administration of 50 mg losartan in presence or absence of aspirin (81 mg).

<table>
<thead>
<tr>
<th></th>
<th>P value</th>
<th>Mean difference</th>
<th>95% confidence interval of the difference</th>
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<td>$T_{1/2}$</td>
<td>0.4315</td>
<td>-0.08667</td>
<td>-0.3118 to 0.1385</td>
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<tr>
<td>$C_L$</td>
<td>0.5547</td>
<td>3.574</td>
<td>-8.835 to 15.982</td>
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<tr>
<td>AUC$_{0-8}$</td>
<td>0.5905</td>
<td>8.082</td>
<td>-22.746 to 38.910</td>
</tr>
<tr>
<td>C$_{max}$</td>
<td>0.9878</td>
<td>-0.4238</td>
<td>-57.370 to 56.522</td>
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<tr>
<td>V$_d$</td>
<td>0.6472</td>
<td>12.457</td>
<td>-68.381 to 43.467</td>
</tr>
</tbody>
</table>

---

**Figure 3:** Mean sitting systolic and diastolic blood pressures and sitting heart rates of losartan (50 mg) in the presence or absence of aspirin (81 mg) in 21 healthy volunteer.
that there is no pharmacokinetic interaction between both drugs. The finding is important since it reassures the safe use of combining AAS to losartan treatment.

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References


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