INTRODUCTION

As obesity reaches current pandemic proportions, the number of bariatric surgeries also increases worldwide. This surgical treatment has been consolidated as an effective therapy for obesity and for the long-term maintenance of weight loss. Nevertheless, surgery for obesity has been associated with some early and late complications that account for significant morbidity. The major villain in this group of complications is infection; therefore, prevention is paramount in the management of these patients.

For infection prophylaxis to be successful, the spectrum, pharmacokinetics and toxicity of the selected antimicrobial agents need to be taken into consideration, as well as other factors such as the appropriate duration of therapy and the maximum concentration of the drug in the tissues at the time of the incision. However, the currently available guidelines on antibiotic prophylaxis are based on healthy, non-obese patients.

The literature in general, and that related to obese surgical patients in particular, is sparse with respect to the use of antibiotics and their distribution in the body of obese patients. Little is known on the pharmacokinetics of antibiotics in patients whose body mass index (BMI) is >40 kg/m$^2$. Even so, the manner in which obese patients absorb, distribute, metabolize and excrete drugs in general is known to be different from that of non-obese individuals.

Indeed, the relationship between body size and physiological and pharmacokinetic variables in the obese population implies that some physiological changes that are characteristic of morbid obesity affect the kinetics of drugs. These changes involve an increase in cardiac output, total blood volume and renal clearance, as well as the occurrence of fat deposits in the liver and changes in plasma proteins.
Taking into consideration that the incidence of surgical site infection in patients operated on for morbid obesity is high and the current recommendations for prophylactic antibiotics are flawed, with surgical site infection tending to result in significant morbidity, it is assumed that the current prophylactic regimens probably fail to provide adequate tissue levels of the drugs in the morbidly obese. Therefore, this study was conducted to analyze the rates of surgical site infection (SSI) with three different antibiotic prophylactic regimens.

METHODS

A prospective, cross-sectional study including a total of 896 Roux-en-Y gastric bypasses performed to treat obesity was conducted between January 2009 and January 2013 at the Hospital of the Federal University of Pernambuco, Northeastern Brazil (Table 1). The institute’s internal review board approved the study protocol prior to its initiation and all patients signed an informed consent form.

All the procedures were performed by laparotomy. The study analyzed the rate of SSI according to the prophylactic regimen used and also evaluated associations with other factors such as age, sex, preoperative weight, BMI and comorbidities.

At admission, patients were examined to rule out any possible sites of infection and to identify any community-acquired infection, with the procedure being cancelled if any were found. Two hours prior to surgery, patients were asked to take a shower and wash their hair, after which the surgical site was washed using a chlorhexidine solution. Trichotomy was performed after anesthesia was induced. After a skin incision was made, subcutaneous tissue was carefully retracted, limiting injury to as few adipocytes as possible and minimizing the risk of developing a seroma. The aponeurosis was closed using continuous absorbable sutures. The subcutaneous tissue was sutured to diminish the dead space. No drains were used in the subcutaneous tissue. In all cases, the skin was closed by suturing.

The study compared three groups of patients according to the type of antibiotic prophylaxis used for bariatric surgery, with the antibiotics being administered intravenously at the induction of anesthesia in all cases. In Group I, 194 patients were treated with two doses of ampicillin (2.0 grams) / sulbactam (1.0 gram); in the 303 patients in Group II, treatment consisted of a single 1-gram dose of ertapenem; and the 399 patients in Group III received a single 2-gram dose of cefazolin at induction of anesthesia followed by a continuous infusion of 1 gram of cefazolin throughout the surgical procedure. Patients were followed up for a minimum of 30 days to diagnose and control any possible infections.[12, 14] The presence of pus was the criterion used to classify the wound as infected. Whenever infection was detected at the surgical site, the wound was opened. Dressings were changed daily. Seromas were managed by manual expression, without need to reopen the wound.

The data were presented as tables of frequency distribution. Means and medians were calculated and the chi-square test was used to evaluate differences at a significance level of 95% (P<0.05).

RESULTS

The most common surgical complications recorded were related to the surgical site No patients had to be re-operated and no deaths occurred as a result of these complications. There were no statistically significant differences between the three groups with respect to surgical site infection (Table 2). The Frequency of postoperative complications according to the type of antibiotic prophylaxis used were no statistically significant differences.

### TABLE 1. Patient-related variables according to the type of antibiotic prophylaxis used for bariatric surgery

<table>
<thead>
<tr>
<th>Variable</th>
<th>Ampicillin/sulbactam (n = 194)</th>
<th>Ertapenem (n = 303)</th>
<th>Continuous Cefazolin (n = 399)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Females</td>
<td>116 (59.8%)</td>
<td>181 (59.7%)</td>
<td>248 (62.2%)</td>
</tr>
<tr>
<td>Males</td>
<td>78 (40.2%)</td>
<td>122 (41.3%)</td>
<td>151 (37.8%)</td>
</tr>
<tr>
<td>Mean age (range) (years)</td>
<td>35.1 (16-61)</td>
<td>34.7 (20-64)</td>
<td>35.1 (16-70)</td>
</tr>
<tr>
<td>Mean weight (range) (kg)</td>
<td>124.8 (86-215)</td>
<td>136.2 (87-232)</td>
<td>136.9 (85-202)</td>
</tr>
<tr>
<td>Mean BMI (range) (kg/m²)</td>
<td>43 (35-79)</td>
<td>45 (33-72)</td>
<td>45 (33-65)</td>
</tr>
<tr>
<td>Comorbidities</td>
<td>139 (71.6%)</td>
<td>205 (67.6%)</td>
<td>278 (69.7%)</td>
</tr>
</tbody>
</table>

Group I versus Group II: P=0.371; Group I versus Group III: P=0.143; Group II versus Group III: P=0.890.
significant differences between the groups (Table 3). No adverse reactions occurred during the study that could be attributed to the antimicrobial agents used as prophylaxis.

DISCUSSION

Recommendations regarding the prophylactic treatment of infection in obese patients are currently identical to those adopted for non-obese patients. The guidelines issued by the Centers for Disease Control and Prevention (CDC) in 1999 recommend the prophylactic use of antibiotics, administered intravenously to ensure peak plasma concentrations at the time of incision, maintaining therapeutic blood and tissue levels throughout surgery and up to a few hours after skin closure.

Drugs are absorbed, distributed, metabolized and excreted differently in obese compared to non-obese patients. Few data are available in the literature with respect to pharmacotherapy in obese patients and the available data are limited to only a few drugs. As with anesthetics, chemotherapy drugs and certain other drugs, antimicrobials may be significantly affected by obesity and by changes determined by surgical procedures.

The volume and the speed of distribution of drugs are determined by many factors including body mass, blood flow to the tissues, tissue protein binding and the kinetics of drug excretion. The relative importance of each of these factors varies with the physical and chemical characteristics of the drug, including its lipid solubility. Most antimicrobial substances are polar, or hydrophilic, meaning that they are easily distributed in water but not in adipose tissue.

Surgical site infections develop in 5.6%-20% of patients undergoing gastroplasty when cefazolin is administered prophylactically. The incidence of surgical site infections in patients operated on for obesity is high and there is a need to establish recommendations for antibiotic prophylaxis in this population. For this reason, since 1997 this research group has been involved in studying safe and effective prophylaxis.

The water content of adipose tissue is approximately 30% of that found in other tissues. Consequently, the volume of distribution of hydrophilic drugs in this type of tissue may be around 30% of that found in other tissues. The distribution of hydrophilic antimicrobials in the water content of adipose tissue explains the need to increase the dose proportionally to compensate for excess body weight, using the dose correction factor for the patient’s weight.

In a study conducted by Edmiston et al., patients submitted to Roux-en-Y gastric bypasses for morbid obesity were given 2 grams of cefazolin preoperatively followed by a second dose three hours later. Thirty-eight patients were assigned to one of three groups according to their BMI: A) 40-49 (n=17), B) 50-59 (n=11) and C) ≥60 kg/m² (n=10). Multiple timed serum and tissue samples were collected and cefazolin levels were assessed by microbiological assay. Significantly lower concentrations of cefazolin were found in the adipose tissue (P=0.04), initial (P=0.03) and omental flap closure (P=0.05) in groups B and C compared to group A. Therapeutic tissue levels were achieved in only 48.1%, 28.6% and 10.2% of the specimens in groups A, B and C, respectively.

In a study carried out by Forse et al., morbidly obese patients submitted to gastroplasty were randomly selected to receive 1 gram of cefazolin either into the fatty tissue of the gluteal region, intramuscularly in the gluteal region or by intravenous injection, while a fourth group of morbidly obese patients were given 2 grams of cefazolin intravenously. At incision and closure, both blood and tissue levels of cefazolin were significantly (P<0.001) lower in all the morbidly obese patients who received 1 gram of cefazolin compared to the blood and tissue levels of the drug found in patients of normal weight. Cefazolin levels below the minimum inhibitory

<table>
<thead>
<tr>
<th>Complications</th>
<th>Ampicillin/sulbactam</th>
<th>Ertapenem</th>
<th>Continuous Cefazolin</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Seroma</td>
<td>76</td>
<td>15</td>
<td>151</td>
<td>342</td>
</tr>
<tr>
<td>Atelectasis</td>
<td>09</td>
<td>19</td>
<td>18</td>
<td>46</td>
</tr>
<tr>
<td>Respiratory infection</td>
<td>09</td>
<td>03</td>
<td>5</td>
<td>13</td>
</tr>
<tr>
<td>Urinary tract infection</td>
<td>03</td>
<td>06</td>
<td>5</td>
<td>14</td>
</tr>
<tr>
<td>Gastric fistula</td>
<td>03</td>
<td>03</td>
<td>3</td>
<td>7</td>
</tr>
<tr>
<td>Thromboembolism</td>
<td>02</td>
<td>03</td>
<td>2</td>
<td>10</td>
</tr>
<tr>
<td>Intra-abdominal infection</td>
<td>03</td>
<td>03</td>
<td>3</td>
<td>9</td>
</tr>
<tr>
<td>Postoperative death</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>

TABLE 3. Frequency of postoperative complications according to the type of antibiotic prophylaxis used for bariatric surgery.
concentration of >2 micrograms/mL for gram-positive cocci and >4 micrograms/mL for gram-negative rods were found. When the morbidly obese patients were given 2 grams of cefazolin, the incidence of surgical site infection fell from 16.5% to 5.6%; therefore, the recommended dose of cefazolin for the morbidly obese patient was increased from 1 to 2 grams(15).

Ferraz et al. evaluated two groups of patients undergoing Roux-en-Y gastric bypasses in a study in which patients receiving two 3-gram doses of ampicillin-sulbactam as antimicrobial prophylaxis were compared with patients receiving a single 1-gram dose of ceftriaxone. No statistically significant differences were found between these two groups with respect to the incidence of surgical site infection(11).

In a study conducted by van Kralingen et al., younger age rather than body weight was shown to be associated with a significantly higher clearance of cefazolin. However, since the unbound plasma concentrations of cefazolin remained above 1 mg L(-1) in all patients weighing ≤260 kg up to 4 hours after intravenous administration of a 2-gram dose, repeating the dose within 4 hours of administration or giving the patients a dose of another class of antibiotic should only be considered in the case of a higher minimum inhibitory concentration for 90% of the isolates tested(18).

Current analysis of pharmacokinetic dosing suggests that the strategies adopted may fail to provide adequate perioperative prophylaxis in gastric bypass patients. The results of the present study show rates of surgical site infection that range from 4.16% when prophylaxis with ampicillin-sulbactam is used to 1.98% when ertapenem is used and 1.55% with continuous cefazolin. The use of continuous cefazolin as prophylaxis during the surgical treatment of morbid obesity has shown very promising results. However, further studies are required to evaluate the effect of this therapy on hospital microflora and bacterial resistance. Therefore, based on these findings, some prophylactic regimens should be reconsidered and even substituted to ensure the prevention of surgical site infection in bariatric patients.

Authors’ contributions
Ferraz AAB: literature review, data analysis, preparation of the manuscript; Siqueira LT: survey data, data analysis; Campos JM: survey data, data analysis; Araújo Jr GC: literature review, data analysis, preparation of the manuscript; Martins Filho ED: literature review, survey data, data analysis, preparation of the manuscript; Ferraz EM: survey data, data analysis.
REFERENCES


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