Knowledge and Perceptions of Final Year Medical Students in Iraqi Universities about Generic Medicines

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Abstract

Objective: To explore and evaluate final year medical students’ perceptions of and knowledge about generic medicines in Iraqi universities.

Methods: A cross-sectional survey was conducted between 1st June 2010 and 31st July 2010 in six Iraqi public universities offering courses in medicine.

Results: Responses were received from 546 medical students (response rate 69.6%). More than 60% of study participants thought that generic medicines are inferior, less effective and produce more side effects compared to brand name medicines. These findings highlight that final year medical students need a better understanding of the principles and concepts of bioavailability and bioequivalence if they are to contribute appropriately to generic medicine use.

Conclusions: This study clearly shows that medical students in Iraq lack of understanding on issues related to generic medicine use. In order to encourage the use of generic medicines in Iraq, medical students need to be receiving a better education on the issues relating to generic medicines and generic prescribing. This could be achieved by including the relevant topics in the current medical education curriculum.

Keywords: Generic medicines; Generic prescribing; Bioequivalence; Perceptions

Introduction

The reasons especially among authorities in Europe for the increasing focus on medicines is their expenditure growth in recent years, greater that other components, leading to pharmaceutical expenditure now the largest or equal largest component. In addition, the need to continue to provide equitable and comprehensive healthcare in the face of pressures including demographics, rising patient expectations and the continued launch of new premium priced drugs with pressures from consultants, patients and others to prescribe these [1-3].

An increasing strategy employed by authorities surrounds increasing the utilisation of generics as well as introducing strategies to obtain low prices to take full advantage of over $100bn/ year of products likely to lose their patent by 2012 - especially with such wide variation in current prices of generics [3-6].

The debate surrounding generic prescriptions has centred on issues relating to bioequivalence, quality and safety, [2,7-13]. Some of these issues relate to the fact that in some European countries there are branded generics. This can cause patient confusion leading potentially to medication errors. In Iraq, Concern has been expressed by physicians practicing elsewhere about the efficacy of generic Medication. This debate has centered on issues related to bioequivalence and potential confusion that might arise when changes of medicine brands occur in some patient populations [14]. Furthermore, previous studies have shown that changing existing prescribing behaviors is difficult. [15,16]. So, it is important to train medical students with getting used to INN prescribing as happens in the UK for instance and reinforced in the community [1,17]; influencing existing prescribers can appreciably enhance prescribing efficiency. Consequently, would suggest a dual strategy. However, may be easier to start with students as the first group. The aims of this study are to explore and evaluate final-year medical students’ perception and knowledge of generic medicines and generic prescribing in Iraqi universities.

Methodology

Development of the survey questionnaire

As Hassali and associates used the questionnaire in their study and is validated also, it is reasonable to use pre-validated questionnaire in survey research. Therefore, on this pretext we decided to use the validated questionnaire which we revalidated before the execution in the current study [18].

A total of 23 survey items, classified into four groups, were developed. The first part consisted of three demographic questions: age, gender, and the university at which the participants were currently enrolled. In the second part, items were pooled under the heading of ‘knowledge of bioequivalence of generic medicines’. This section included a question which asked the student to select the correct bioequivalence regulatory limits when comparing generic medicines with innovator brands according to the requirements of the US Food and Drug administration (FDA). The other four questions were framed in a five-point Likert scale format (5 = ’strongly agree’, 4 = ’agree’, 3= ’neutral’, 2= ‘disagree’ and 1 = ’strongly disagree’). This type of scale was chosen because its construction is relatively simple and the interpretation of results is straightforward. [19].

The third part of the survey contained eight items which evaluated the medical students’ understanding of brand name medicines versus generic medicines.

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These items were also framed in a five-point Likert rating scale. The fourth part of the questionnaire consisted of seven items designed to evaluate the medical students’ perceptions of education they were currently receiving. Similarly to part three, these questions were also answered using a five-point Likert scale.

In order not to overlook any of the students’ opinions on the issue of generic and brand name medicines, a blank space was provided on the last page of the survey for the respondents to write down their views.

Face and content validation of the paper survey

Three lecturers of pharmacology from Basra college of Medicine and Pharmacy with experience in the field of medicines were asked to evaluate the relevance, clarity and conciseness of the items included in the questionnaire. The observations and comments given by the lecturers were taken into account [20]. In order to test the validity and the reliability [21,22] of the survey form, the revised questionnaire was pilot-tested by administering it to a convenient sample of 20 medical students who were not participating in the main study. The overall Cronbach’s alpha value was 0.72.

Study design and sample size

A cross-sectional survey was conducted between 1st June 2010 and 31st July 2010. Convenience sample method was used, six universities were chosen from the middle and south of Iraq. The sampling frame was all final-year medical students in six universities who were enrolled during the period of the study. The number of enrolled students during the study period was obtained from the Dean’s Office in each university as shown in Table 1.

Survey administration and timeframe

This descriptive and analysis study was conducted over a period of two months, between the 1st Jun 2010 and the 31st of July 2010. In order to inform the students about the objectives of the survey, an explanatory letter attached to the survey questionnaire was distributed to all of the participants. The students received the survey questionnaire from the respective lecturers in each university. Permission to approach the participants. The students received the survey questionnaire from the respective lecturers in each university. Permission to approach the students and to conduct the study was obtained from the Ministry of Higher Education and Scientific Research in Iraq.

Data collection

Responses from the students were collected by the medical lecturers in each university and were entered (key - in) into the Statistical Package for Social Sciences (SPSS) for Windows, version 16, by the researcher in order to perform statistical analysis.

Data analysis

Data analysis was performed using SPSS® for Windows, version 16 [23] The mean (± SD) was calculated for the continuous variables, and frequencies were measured for the categorical variables. Associations between groups were examined using chi-square or Fisher exact tests whenever necessary. For categorical variables, Fisher’s exact test is preferred over the chi-square test if 20-25% or more cells in the table have expected frequencies of less than five, or if any of the expected frequencies are less than one [22,24,25]. Data which emerged from the domains using the Likert scale as a measurement were analysed statistically as non-parametric data [26-28]. However, the response categories in the Likert scale have a ranking order, but the intervals between the values cannot be presumed to be equal [29,30].

For this survey data, a default Monte Carlo simulation in the SPSS software was used to estimate Fisher’s exact p-values as the data set was large and normal exact computations require a significant amount of time and memory from the computer. A two-sided 99% confidence level Monte Carlo estimate of Fisher’s exact p-values was computed, with the p-value of 0.05 or less taken to demonstrate statistical significance.

Results

The total number of medical students in the six universities in question in Iraq during the time of the survey was 785. By the end of the survey period, 546 students had responded to the survey (response rate = 69.6%). The responses of each university are shown in Table 1.

The average age of the respondents was 23.78 ± 0.72 years. Of the respondents, 248 (45.4%) were 24 years old, and 283 (51.8%) of the respondents who responded were female (Table 3).

The first item on the questionnaire asked the students to select the correct bioequivalence limits allowed by the FDA when comparing an innovator drug with a generic one. The following explanation was provided with the question:

Table 1: Final-year medical student enrolment at six Iraq universities, 2010.

<table>
<thead>
<tr>
<th>University</th>
<th>Number of enrolled final year students</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>286</td>
</tr>
<tr>
<td>2</td>
<td>110</td>
</tr>
<tr>
<td>3</td>
<td>56</td>
</tr>
<tr>
<td>4</td>
<td>258</td>
</tr>
<tr>
<td>5</td>
<td>20</td>
</tr>
<tr>
<td>6</td>
<td>75</td>
</tr>
<tr>
<td>Total</td>
<td>785</td>
</tr>
</tbody>
</table>

*20 students were excluded for pilot study

Table 2: Response rates according to universities.

<table>
<thead>
<tr>
<th>Response</th>
<th>Frequency</th>
<th>Percent (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>80%-120%</td>
<td>54</td>
<td>9.9</td>
</tr>
<tr>
<td>80%-125%</td>
<td>49</td>
<td>9.0</td>
</tr>
<tr>
<td>90%-100%</td>
<td>76</td>
<td>13.9</td>
</tr>
<tr>
<td>95%-100%</td>
<td>99</td>
<td>18.1</td>
</tr>
<tr>
<td>95%-105%</td>
<td>64</td>
<td>11.7</td>
</tr>
<tr>
<td>i don't know</td>
<td>204</td>
<td>37.4</td>
</tr>
<tr>
<td>Total</td>
<td>546</td>
<td>100.0</td>
</tr>
</tbody>
</table>

Table 3: Demographic characteristics according to university.

<table>
<thead>
<tr>
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<td>546</td>
<td>100.0</td>
</tr>
</tbody>
</table>

Table 4: Responses to question 4.
In pharmacology, the term ‘bioavailability’ refers to the rate (how fast) and the extent (how much) to which an active ingredient is absorbed and becomes available at the site of the drug action. The FDA organisation considers a generic product to be bioequivalent if its bioavailability is within a specified range compared with the currently marketed branded product.

After this statement, the following question was asked:

The regulatory limits applied are that the 90% confidence intervals for the ratios (generic product: brand name product) of the areas under the plasma drug concentration versus time curves and the maximum plasma drug concentrations must fall between...

Six answer options were given, the correct answer being 80-125%.

The responses are shown in Table 4. Only 49 (9.0%) of the students answered the question correctly, while 293 (53.6%) answered incorrectly and 204 (37.0%) did not attempt to provide a numerical answer.

The responses to other questions relating to the students' knowledge of bioequivalence are shown in Table 5.

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The responses to other questions relating to the students' knowledge of bioequivalence are shown in Table 5.
More than 70% of the respondents did not believe that generic medicines are therapeutically equivalent to the corresponding brand name medicines, or even to each other. There was a statistically significant difference between the responses to these two questions according to universities (p<0.001 for both questions). More than 70% of the respondents did not possess sufficient information about bioequivalence for generic medicine and how bioequivalence tests are conducted for generic medicines. There were no statistically significant differences according to gender and the different universities for these two questions.

The responses to questions about understanding of brand name versus generic name are shown in Table 6. More than 60% of the respondents thought that a generic medicine was not bioequivalent to the corresponding branded medicine. There was a statistically significant difference in the responses to this question from different universities. More than half of the respondents did not know that the generic medicine must be presented in the same dose and dosage form as the brand name medicine. There was a statistically significant gender difference in the responses to the question regarding the form of the dosage (p=0.006). With regard to quality and efficacy, around two-thirds of the respondents agreed that generic medicines are of a lower quality (69%), are less effective (66%), produce more side effects (61.5%) and adhere to lower safety standards (68.8%) than branded medicines. There was a statistically significant difference in the responses to the question about quality between the different universities (p<0.001). According to the question regarding the price of generic medicines, 84.4% of the respondents believed that generic medicines are less expensive than brand name medicines. A statistically significant difference (p<0.001) was found between the responses of the different universities.

With regards to the participants’ perceptions of the current education in prescriptions, Table 7 shows the responses to these questions. Around 80% of the respondents needed more information on issues pertaining to the safety and efficacy of generic medicines. Statistically significant differences (p = 0.013) were noted in the responses to this question from the various universities. Of the respondents, 353(64.7%) felt confident enough to prescribe medicines according to their generic name rather than a brand name. Statistically significant differences (p<0.001) were noted in the responses to this question from the different universities. Of the respondents, 63.2% found it easier to recall a medicine’s therapeutic class using generic names rather than brand names. Statistically significant differences (p = 0.015) were found in the responses to this question from the different universities and between genders (p = 0.050). Of the respondents, 68.3% believed that pharmacists would give them advice in the future with regard to generic medicines. Statistically significant differences were found between the responses to this question from the different universities (p<0.001). Almost 68.3% of the respondents agreed that their future prescribing habits would probably be influenced by their senior colleagues and medical consultants. Statistically significant gender differences were found in the responses to this question (p=0.020), as a higher proportion of female students strongly agreed or agreed with this item. More than half of the respondents (56.8%) believed that advertisements by drug companies would influence their future prescribing patterns. Statistically significant differences were found in the response to this question from the various universities (p<0.001). Around 60% of the respondents believed that their university did not adequately cover the topic of cost-effective prescribing on their medical curriculum.

Discussion

Part 1 - Demographics and response rate

To the best of our knowledge, this is the first study in Iraq which has evaluated the knowledge and perceptions of final-year medical students in universities in Iraq towards generic medicine. In the present study, an overall response rate of 69.6% was recorded. One reason for a lack of responses to this survey may be the lack of information regarding generic medicines among the students. With regard to the respondents’ demographic characteristics, 51.8% of the respondents were female, while 48.2% were male. The average age of the students was 23.78 ± 0.72 years old.

Part 2 - Understanding of the bioequivalence of generic medicines

According to the FDA, the determination of the bioequivalence of a generic medicine involves an assessment of pharmacokinetic parameters such as the area under the curve (AUC) and peak concentration (Cmax), as well as the calculation of a 90% confidence interval for the ratio of the average of these parameters for the two products that are compared, usually a test product (generic product) against a reference product (the innovator or the branded product). The calculated confidence interval should fall within a conventionally established bioequivalence range of 80% to 125% for the ratio of the product averages. This reflects that the difficulty of the topics of pharmacokinetics and bio-pharmaceutics make it difficult for students to understand them. In the first question, which asked the students about their knowledge of the bioequivalence limits for the approval of generic medicine by the FDA, only 9.0% (n = 49) of the students gave the right answer and 37.4% (n=204) of the respondents had no idea. This reflects the deficiency in the students’ knowledge of pharmacokinetic parameters, which are important in the areas of bioequivalence and bioequivalence testing with generic medicine. The poor response to this question in this study confirmed the response to the same question in the study which was conducted in Australia by Hassali and his group [18]. These two studies in Australia and Iraq clarify that topics relating to bioequivalence are not covered and that these topics are important for medical students during their education, as their education will help them to be more confident in prescribing generic medicine in the future.

Part 3 - Understanding of brand name versus generic medicines

More than 70% of the respondents did not believe that generic medicines were therapeutically equivalent to the corresponding brand name medicines, or even to each other. This is due to bad experiences among patients with regard to the generic medicines used in Iraq over the last 20 years. During this time, most medicines which have been imported into the country have not undergone a quality control check, due to the shortage of medicine as a result of an international embargo which has been in place since 1990'(31). This result was in line with the results of other studies conducted in different countries [2,7-13]. While The findings of the current study are not in line with those of the previous study which was carried out in Australia by Hassali et al. [18], and in which more than 80% of the respondents believed that a generic medicine was bioequivalent to the corresponding brand name medicine. Statically analysis show significant differences among universities, the differences in the responses might occur due to deficiency in the universities curricula in teaching this topic to their students. According to questions which explored the medical students’ knowledge of dosage forms and doses when comparing generic medicines with branded
medicines according to the FDA’s requirements, more than half of the respondents did not know the definition of generic medicine according to this organisation. Under the FDA’s regulations, a generic medicine must contain identical amounts of the same active ingredients in the same dosage form as a branded medicine [32]. Around half of the respondents were not aware of these two facts. This response pattern might be influenced by lack of education about generic medicines. Statistically analysis show significant differences according to gender with the fact that the generic medicine must be in same dosage forms, a higher proportion of male students either strongly disagreed or disagreed with this question compared to their female counterparts. This response reflects a lack of information regarding the requirements of generic medicines according to the FDA. The findings of the current study are in line with those of a previous study carried out in Australia by Hassali [18].

According to the questions that evaluated the medical students’ knowledge of the quality and effect of generic medicines when compared to brand name medicines, 377 of the respondents (69%) believed that the generic medicines in Iraq were of a lower quality. Of the respondents, 336 (61.5%) thought that generic medicines would cause more side effects than brand name medicines. Statistically analysis showed significant differences among universities and the differences in the responses might occur due to differences in the universities curricula in teaching this topic to their students. While 433 of the respondents (66.7%) believed that they were less effective than their brand name counterparts. These results were in line with the results of the study conducted in Australia by Hassali [18]. Which illustrated the deficit in medical students’ knowledge of generic medicines. More than 80% of the respondents were fully aware that generic medicines are less expensive than brand name medicines. Statistically analysis showed significant differences among universities, the differences in the responses might occur due to differences in the universities curricula in teaching this topic to their students. Of the respondents, 376 (68.8%) thought that brand name medicines were required to meet higher standards than generic medicines.

Part 4 - Perceptions of the current education in prescribing

Most of the respondents in the current study (74.9%) said that they would have liked more information on the safety and efficacy of generic medicines during their years of medical education. The highest proportion of respondents from the University No. 6, while the lowest came from the University of respondents No. 5. Of the respondents, 353 (64.7%) were more confident in prescribing using generic names rather than brand names because the curriculum deals with generic names more than brand names, the highest proportion of respondents from the university No.3, while the lowest came from the university No.5. This may be reflective of different styles of teaching among universities. Of the respondents, 345(63.2%) found it easier to recall a medicine’s therapeutic class using generic names rather than brand name, The highest proportion of respondents from the university No.6, while the lowest came from the university No.2. The proportion of males is higher than the proportion of females in relation to this question. Nearly two-third of the respondents thought that pharmacists were one of the most important groups of healthcare professionals in terms of providing medical students with advice on generic medicines, while in a similar study in Australia [18]. The respondents were more confident in pharmacists, with 86.8% thinking that pharmacists comprised one of the most important teams of healthcare professionals in terms of providing the respondents with advice. This reflects the different views held with regard to pharmacists in both countries; in Iraq, many private pharmacies are run by non-medical staff, and this decreases patients’ confidence in pharmacists’ advice. More than half of the students in Iraq agreed that advertising by drug companies would influence their future prescribing patterns, while in Australia, one quarter of the students believed that advertising from companies would influence their prescribing patterns [18]. These results was in line with the study conducted by Godman et al. [33] where more than half of the physicians stated that Pharma Company activity would influence future prescribing habits as Pharma Companies spend a great deal of money on marketing activities as well as physician education [33]. More than half of the students in Iraq and Australia believed that their universities did not adequately cover the topic of cost-effective prescribing in their medical curriculum. A statistically significant difference was noted for all these six responses among universities; The differences in the responses might occur due to differences in the universities curricula in teaching these topics to their students. In the present study, almost two-thirds of the medical students strongly agreed or agreed that their future prescribing habits would probably be influenced by their senior colleagues and medical consultants. Statistically analysis showed significant differences according to gender, where high response from Female students than male students was noted. In psychology it is generally seen that female have a tendency to follow their seniors and in that context, in the current study, females are influenced by their senior colleagues opinions and decisions. This result is in line with the results of the study conducted in Australia by Hassali et al. [18], in which 92% of the respondents strongly agreed or agreed that their future prescribing habits would probably be influenced by their senior colleagues and medical consultants. Similarly, in a previous study conducted among interns in Australia, 80% of the medical interns believed that their consultants would influence their prescribing practices [34]. This clarifies the deficiency in topics that encompass prescription-related issues in our universities, as their actual prescribing training will only start in the hospital.

Limitations of the study

The number of students who participated in this study was relatively small compared to the total number of students who are enrolled in medical colleges in Iraq. Therefore, these results may not necessarily be generalisable to all medical students.

Conclusion

This study clearly shows that medical students in Iraq lack an extensive understanding of the concept of bioequivalence and the safety and quality of generic medicines. In order to encourage the use of generic medicines in Iraq, medical students need to be receiving a better education on the issues relating to generic medicines and generic prescribing. This could be achieved by including the relevant topics in the current medical education curriculum.

Reference


