



Knowledge and Attitude About Research Ethics Among Iranian Dental Students

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ABSTRACT

Objective: To evaluate the knowledge and attitude of the students of Kerman dental school (Iran) about ethics in dental research. **Material and Methods:** This cross-section study was conducted on 307 dental students selected through the census sampling method. Data were collected by a researcher-made questionnaire consisting of 12 items about knowledge and 17 items about attitude toward research ethics. Data analyzed in SPSS software using t-test and linear regression test. P-values of less than 0.05 were considered statistically significant. **Results:** Of the respondents, 33.9% were male and 66.1% were female, and 44% had good knowledge and 20.8% had a positive attitude about research ethics. A significant correlation was found between knowledge and attitude. A significant correlation was also observed between knowledge and participation in research workshops. Knowledge and attitude showed no significant correlation with gender or year of admission. **Conclusion:** Participants had appropriate knowledge and attitude about research ethics education concerning experimental works and retrospective studies on biologic samples. Holding research workshops with an introduction to ethical codes of research is recommended.

Keywords: Bioethics; Ethics, Research; Health Knowledge, Attitudes, Practice; Students, Dental.

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Introduction

The past decades have seen rapid progress in dental research with significant contributions to oral health and public health in general [1]. In many cases, health researchers have to use human participants in their studies, which brings up ethical requirements for protecting the rights, dignity, and well-being of participants [2].

There are numerous ethical and legal requirements for clinical research, which must be given the highest priority to ensure that participating patients are completely protected. This is especially important when working with vulnerable groups such as children and patients who cannot complete the consent form. Despite the global acceptance of the ethical guidelines stated in the Declaration of Helsinki [3], it is still possible to encounter violations and dishonesty in adhering to these requirements [4]. The act of acquiring informed consent is an essential prerequisite for any biomedical research involving diagnosis and treatment because, without it, the dentist will be vulnerable to litigation [5].

A study by Deolia et al. [6] on research ethics in India showed that dental professionals had a good attitude toward the issue, but there was room for improvement in their knowledge and behavior. This study also reported a statistically significant association between attitude toward the issue and age and gender [6]. In a study carried out by Mallela et al. [7] on research ethics in northern India, more than 90% of the participants were aware of ethics committees' presence but had little knowledge about international ethical guidelines. Also, 20% of the participants believed that research ethics committees cause delays in research [7].

According to Kakar et al. [8], informed consent is even more critical when providing medical care to children and foreigners and using medical images in medical and dental research. In the research conducted by Taiwo and Kass [9] in Nigeria on the people involved in oral health research, 93.8% of the respondents said they were not asked whether they want to volunteer, and all of them generally had poor knowledge about the key items of informed consent. El-Dessouky et al. [10] also reported the low awareness of participants in dental research. In a study by Ghaderi et al. [11], university instructors and professors had moderate knowledge and post-graduate students had poor knowledge about research's ethical guidelines.

Considering the involvement of biomedical research with human studies and the general scarcity of information in this regard, especially in dental field in Iran, and in continuing our interest in medical research [12,13], the present study aimed to assess the knowledge and attitude of Kerman Medical University dental students about research ethics.

Material and Methods

Study Design and Sample

This descriptive cross-sectional study was performed through a census sampling method. All students of Kerman dental school (capital city of Kerman province in South, Eastern of Iran) were included. First, a list of college students in the academic year was prepared, and briefing sessions were held at the theoretical classrooms of students of each entrance classroom by an assistant (a senior dentistry student) with sufficient knowledge about the questionnaire and the aim of the study. In these sessions, research objectives were explained and verbal consent was obtained. It was also explained that participation is completely optional and responses with be anonymous.

Data Collection

Questionnaires were administered and collected at the same session. The approximate time for completing the questionnaire was 15 minutes. The questionnaire consisted of three parts. The first part was dedicated to collecting information about the student's demographic characteristics, including gender, year of admission, participation in research methodology workshops, participation in ethics workshops, published papers, and research proposals. The second part included 12 questions measuring the students' knowledge about ethical requirements of dental research (about the Declaration of Helsinki, health requirements of participants, research purpose, intervention complications, informed consent, informing participants of results, use of human data or materials such as blood, saliva and extracted teeth, approval of the ethics committee). For this part of the questionnaire, correct answers were awarded a score of 1 and wrong answers were given a score of 0. Therefore, the total score of this part ranged from 0 to 12. The third part of the questionnaire comprised of 17 questionnaires measuring the students' attitude toward ethics in dental research (ethical considerations regarding laboratory samples, obtaining consent to perform radiography for research, obtaining informed consent for the use of saliva and blood, obtaining consent from vulnerable patients, such as the elderly or intellectually disabled). This part of the questionnaire was designed based on the 5-point Likert scale with answers ranging from completely disagreed to completely agree. Hence, the total score of this part ranged from 17 to 85. For validity assessment verification, the questionnaire's initial draft was reviewed by a group of experts and modifications were made according to their comments. After a second review, experts verified the validity of the modified questionnaire. The validity coefficient of the questionnaire as a whole was estimated at 0.98.

For reliability assessment, the questionnaire was administered to 20 students from different years of admission and the process was repeated two weeks later with the same 20 subjects. Based on this assessment, the reliability coefficient was calculated to 0.91.

Data Analysis

Data were coded and entered into SPSS software, version 21 (IBM Corp., Armonk, NY, USA), and analyzed by linear regression and t-test at p<0.05 significance level.

Ethical Clearance

This study was approved by the competent ethics committee with the ethics code IR.KMU.REC.1395.755.

Results

Of 307 students who participated in the research, 104 (33.9%) were male and 199 (64.8%) were female. One hundred and sixty-seven people (54.4%) did not participate in research methodology workshops. Two hundred and forty people (78.2%) had not taken the medical ethics course, and 36.2% (n=111) were interested in participating in medical ethics workshops (Table 1).

Table 1. Distribution of respondents accord	ing to demographic	variable	- 1
Variables	Ν	%	
Gender			
Male	104	33.9	
Female	199	64.8	
No answer	4	1.33	
Participated in research methodology workshops			

Table 1. Distribution of respondents according to demographic variables.

Yes	140	45.6			
No	167	54.4			
Participated in medical ethics course					
Yes	57	18.5			
No	240	78.2			
Had research proposals					
Yes	66	21.5			
No	241	78.5			
Published papers					
Yes	53	17.3			
No	254	82.8			
Participated in scientific congresses					
Yes	123	40.1			
No	184	59.9			
Participated in medical ethics workshops					
Yes	66	21.5			
No	241	78.5			
Interested in taking the research ethics course					
Yes	111	36.2			
No	196	63.9			

In the knowledge dimension, the item correctly answered most frequently was "physician is required to fully inform the patient about the possible complications of the research and how they must be addressed", which was answered correctly by 248 people (80.8%). On the other hand, only 67 people (12.1%) gave a correct answer to the item "declaration of Helsinki is about the ethical requirements of human research". Among the respondents, 62.2% gave a correct answer regarding the necessity of getting approval from ethics committees in all research projects and 70.7% correctly answered whether it is required to protect the confidentiality and privacy of research participants.

In the attitude dimension, 71.5% of the respondents disagreed or completely disagreed with the statement that "Dental research on people with an intellectual disability is allowed". After reading the statement, "It is required to obtain informed consent from patients before using blood samples in research (during and after taking samples)", 63.6% of the student responded that they disagreed or completely disagreed with this statement. Among the respondents, 16.3% believed that it is not necessary to obtain informed consent to conduct research on the elderly, 44.9% agreed or completely agreed with the statement that there is no ethical consideration regarding the use of prepared dental blocks for research into dental materials, and 46.4% agreed or completely agreed with the statement that there is no ethical consideration regarding the use of prepared dental blocks for research into dental materials, and 46.4% agreed or completely agreed with the statement that there is no ethical consideration regarding the use of prepared dental blocks for research into dental materials, and 46.4% agreed or completely agreed with the statement that there is no ethical consideration regarding the use of prepared dental blocks for research into dental materials, and 46.4% agreed or completely agreed with the statement that there is no ethical consideration regarding the use of extracted teeth in research.

The mean and standard deviation of the knowledge score was 7.43 ± 2.99 from 12. For attitude, the mean score was 39.47 ± 10.05 from 85. Figure 1 shows the distribution of knowledge and attitude scores of the students. As can be seen, 40.4% of the students had good knowledge and 20.8% had a positive attitude about the issue.

A significant direct relationship was observed between the knowledge and attitude of the students (p=0.000). In this study, female students were more knowledgeable and had a more positive attitude about the ethics of dental research. In the case of knowledge, the difference between genders was statistically significant, although marginally (p=0.077). In the case of attitude, the difference was not statistically significant (p=0.41). Participation in research congresses was significantly related to knowledge (p=0.010) but not attitude (p=0.706). There was a significant relationship between participation in ethics workshops and knowledge or

attitude (p=0.529 and p=0.291). Also, no statistically significant relationship was found between the students' knowledge and attitude and their admission year (Table 2).



Figure 1. Distribution of knowledge and attitude scores of the students.

Variables			Mean	Standard Deviation	p-value	
Participated in research methodology workshops	Knowladaa	Yes	7.93	2.71	0.010*	
r articipated in research methodology workshops	Miowiedge	No	7.02	3.16	0.010**	
	Attitudo	Yes	39.73	11.05	0.700	
	Attitude	No	39.26	9.22	0.706	
Doution at his would have	Knowladaa	Yes	7.62	2.83	0.500	
i ai ticipated in etnics workshops	Milowieuge	No	7.37	3.06	0.529	
	Attitudo	Yes	38.25	11.75	0.001	
	Attitude	No	39.88	9.41	0.291	
II - J h	Vl.d	Yes	7.06	3.03	0.050	
Had research proposals	Knowledge	No	7.53	2.90	0.272	
	A 1	Yes	37.63	11.78	0.150	
	Attitude	No	40.02	9.49	0.150	
	17 1 1	Yes	6.96	2.48	0.000	
Published papers	Knowledge	No	7.52	3.04	0.228	
	A 1	Yes	40.64	9.67	0.000*	
	Attitude	No	33.85	9.87	0.000*	
Denti instalin si stifica su marca	Vl.d	Yes	7.98	8 2.67	0.040*	
Farticipated in scientific congresses	Knowledge	No	7.16	3.17	0.042*	
	Attitudo	Yes	40.01	11.33	0.505	
	Attitude	No	39.19	9.17	0.525	
Participated in medical othics workshops	Knowledge	Yes	7.33	2.58	0.754	
i ai ticipated in medical etnics workshops	Milowieuge	No	7.46	3.04	0.754	
	A'. 1	Yes	40.18	9.23	0.051*	
	Attitude	No	36.72	12.46	0.051**	
Interested in taking the research othics course	Knowledge	Yes	7.52	2.72	0.690	
Interested in taking the research ethics course	Milowieuge	No	7.37	3.15	0.080	
	Attitudo	Yes	38.75	10.77	0.387	
	Attitude	No	39.87	9.64		
Condon	Knowledge	Male	7.03	3.30	0.077	
Genuer	Miowiedge	Female	7.69	2.81		
	Attitude	Male	38.81	11.04	0.041	
	Attitude	Female	39.89	9.56	0.041	

	Table 2. Relat	ionship betweer	n the students'	knowledge and	attitude and	demographic	variables.
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Discussion

In the present research, (12.1%) knew that Helsinki's declaration is about ethical consideration of human research. In the study conducted by Mallela et al. [7], 34.8% of the faculty members in northern India said they knew about this declaration's topic. Hariharan et al. [14] also reported that very few of their respondent were aware of this declaration.

It has been shown that researchers must be aware of the ethical and legal requirements of human research in their own country and at the international level [15]. The Council for International Organizations of Medical Sciences (CIOMS) guidelines consists of valuable content regarding the challenges of contemporary clinical research, complex issues such as HIV/AIDS research, the availability of treatments after the research, research on women, protection of privacy and confidentiality, compensation for undesirable outcomes and complications, and guidelines for obtaining consent [16].

In this research, 21.6% of the students had participated in research ethics workshops. In the study of El-Dessouky et al. [10], 36.8% of respondents had participated in research ethics workshops or courses. Reddy et al. [1] reported that 67% of respondents had been educated about research ethics. Compared to the study of El-Dessouky et al. [10], a larger fraction of students surveyed in this study had participated in research ethics workshops and courses, which reflects the higher importance given to the topic at the surveyed university.

In this study, 36.1% of the students believed that "ethics committees often cause delays in research". This is consistent with the findings reported by El-Dessouky et al. [10], where 44% of respondents believed that ethics committees delay the research, but is higher than the 26% reported in the study of Reddy et al. [1]. In the research carried out by Gopinath et al. [15], 31% of university instructors believed that ethical committees delay the research. Given that all research projects need the approval of ethics committees, this finding suggests that the process of obtaining this approval should be accelerated.

In the present study, 22.4% of the students agreed fully or partially that there is no need to inform patients about the materials used in human research, given their limited knowledge about the subject. In the study of El-Dessouky et al. [10], 10% of respondents believed that patients need not be aware of the risk of research. Ethical guidelines, however, state that participants should be made aware of the nature of the research and its possible consequences and make a free choice without any concern about possible effects on their own treatment process [17].

In the present study, 63.3% and 55.7% of the respondents partially or completely disagreed with the necessity of obtaining consent for taking blood or gingival crevicular fluid samples for use in research. El-Dessouky et al. [10] reported that about one-third of their respondents believed that consent is not necessary in such cases. Reddy et al. found that 94 of the 115 respondents partially or strongly agreed with this statement [1].

In our study, 16.3% and 13.3% of the students partially or completely agreed with the necessity of obtaining written consent from the elderly and intellectually disabled. In the study of El-Dessouky et al. [10], almost 40% of people believed that written consent should be obtained from vulnerable people. In this context, the term vulnerable groups refer to people who are relatively or completely unable to protect themselves, including psychological patients, intellectually disabled, prisoners, pregnant women, embryos, people with terminal diseases, students, elderly, and anesthetized people [17]. Helsinki's declaration states that "medical research with a vulnerable group is only justified if the research is responsive to the health needs or priorities of this group" [18].

More than 95% of respondents in the study of Gopinath et al. [15] believed that informed consent should be obtained in writing from all patients to ensure they understand both benefits and risks of research. In the present study, 62.2% of the students partially or completely agreed that "It is necessary to obtain consent before performing radiography for research". In fact, the respondents agreed with this statement more than they did with any other statement, most likely because they were aware of the risks of radiation exposure.

In this survey, 40.4% of the respondents had a good level of knowledge about dental research ethics. However, knowledge about retrospective studies on pathology and biologic samples was tragically scarce. This finding suggests that universities and workshops should put more effort into closing the existing gap in the knowledge of students and researchers about the ethics of retrospective studies that use paraffin blocks or stored biologic samples. On the question that whether retrospective studies need to gain informed consent, the consensus in most countries is that these studies do not require written consent, provided that the issue is disclosed to and verified by the competent ethics committee [19].

In the present study, 16% of the students had poor knowledge and 38.8% had moderate knowledge about ethics in dental research. In the study of Gopinath et al. [15], 38% of instructors had poor knowledge about this topic. Mallela et al. [7] reported that faculty members had a fair knowledge about research ethics. In a study carried out by Ghaderi et al. [11], the students of Kurdistan University showed higher than average knowledge about this subject.

In this study, there was a significant relationship between students' knowledge about ethics in dental research and their participation in scientific congress. The reason for this may be that many lectures given in scientific congresses are based on the principles of ethics in research and have been considered by the participating student.

There was a significant correlation between students' knowledge about ethics in dental research and their participation in research methodology workshops. The reason for this may be that ethical issues have been well discussed in research methodology workshops and have therefore raised awareness.

No significant relationship was observed between participation in ethics workshops and knowledge about ethics in dental research. This is perhaps because these workshops often involve more general principles of research than moral principles in particular.

There was a significant relationship between participation in ethics workshops and attitude about ethics in dental research. In some studies, the reason for this may be that the researcher does not have the necessary knowledge but has a positive attitude towards the observance of ethical principles.

In this study, 52.1% of the respondents partially or completely disagreed with the statement that "Verbal consent is enough for all human research". Obtaining informed consent before clinical trials is one of the basic principles of ethics in research. Kakar et al. [8] argue that informed consent is essential for providing medical care to children and foreigners and for using medical images in medical and dental research. A study carried out in Nigeria showed that 93.8% of the people who had participated in oral health research were not asked whether they want to volunteer, and all of them had generally poor knowledge about the key aspects of informed consent [9]. Informed consent is the foundation of ethics in medical and health care research. This issue is especially important for research on the elderly, where their poor health, cognitive impairments, and passiveness may complicate the situation [20].

It is not arguable that every adult has the right to decide whether or not to perform medical treatment, even if the refusal results in his/her death [21]. The systematic review carried out by Mukherjee et al. [20] showed that informed consent is a neglected subject in dental care and research the elderly.

A significant direct relationship was observed between the students' attitude and knowledge about ethics in dental research, in the sense that more knowledgeable people had more positive attitudes. Thus, it seems in knowledgeable participants have been able to use their awareness to improve their attitude.

In this study, 20.8% of the participants had a positive attitude toward research ethics. In contrast, Kandeel et al. [22] reported that more than 80% of their respondents had a positive attitude toward ethics committees. It should, however, be noted that Kandeel's study was conducted on faculty members rather than students [22]. The study of Gopinath et al. [15] also reported that the attitude of participants about the ethics of research was not desirable.

The present research found a statistically significant relationship between attitude toward research ethics and having published a paper. This is perhaps because the process of writing and publishing a paper involves obtaining approval from competent ethics committees, which requires the researchers to educate themselves about the subject, thereby affecting their attitude.

There was no significant difference between male and female students in terms of attitude toward research ethics. Since ethics is an issue in developing, this lack of difference in attitude between genders is not unexpected.

However, female students had higher mean knowledge scores, although the difference was not significant (p>0.05). This is perhaps because female students have paid more attention to ethics-related discussions in workshops and classes.

In this research, 36.2% of the students wanted to participate in research ethics workshops and there was no significant relationship between this inclination and attitude toward research ethics. In contrast, Gopinath's study reported that most of their respondents were inclined to participate in research ethics workshops [15]. This difference can be attributed to the fact that the participants in our study were university students.

Conclusion

Participants had appropriate knowledge and attitude about research ethics. Overall, there is much room for improvement in research ethics education, including those concerning experimental and laboratory works and retrospective studies on human samples.

Authors' Contributions



Financial Support

None.

Conflict of Interest

The authors declare no conflicts of interest.

Data Availability



The data used to support the findings of this study can be made available upon request to the corresponding author.

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