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RESEARCH NOTE

Assessing knowledge of genetics in undergraduate students in Quito, Ecuador [version 1; peer review: 1 approved, 1 approved with reservations]

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Abstract

Knowledge of genetics is crucial for understanding genetic and genomic tests and for interpreting personal genomic information. Despite this relevance, no data are available about the level of knowledge of genetics in an Ecuadorian population. This investigation sought to survey such knowledge in undergraduate students affiliated with private and public institutions in Quito, the capital city of Ecuador. A total of 350 individuals responded to a validated questionnaire measuring knowledge of genetics. Scores ranged from 45% to 87% (mean: 66.8%), and students achieved slightly better results when asked about genetics and diseases (mean score: 68.3%) than when asked about genetic facts (mean score: 64.9%). Additionally, no significant differences in performance were found among students from private and public institutions. Surprisingly, the lower score obtained (45%) was from a question about how chromosomes are passed to the next generation. The highly educated status of the surveyed population could explain the overall adequate results; nonetheless, the possibility that the correct responses were given by chance cannot be ignored. Therefore, the actual knowledge of genetics among the participants might be less than that revealed by the percentages of correct answers. Consequently, to achieve the goal of ensuring informed decision-making concerning genetic and genomic tests, it seems evident that the national education programs of Ecuador require improvement in teaching of genetic concepts.

Keywords

Ecuador, knowledge of genetics, genetic literacy, undergraduate students, survey

Open Peer Review

Approval Status

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version 1
14 Mar 2019	view	view

1. **Vasiliki Mollaki** , Hellenic National Bioethics Commission, Athens, Greece

2. **Rebecca Carver** , Norwegian Institute of Public Health, Oslo, Norway

Any reports and responses or comments on the article can be found at the end of the article.

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Introduction

Genetic and genomic testing have transformed our understanding of our health, personal well-being and recreational consumerism. Advances in powerful and cheap genetic analyses have allowed new opportunities to generate information about important conditions, such as cancer, diabetes, and cardiovascular diseases (Burton, 2015; Perkins *et al.*, 2018; Rafiq *et al.*, 2015; Roberts & Middleton, 2018; Wu *et al.*, 2019). In recent years, access to pharmacogenomics, nutrigenomics, disease risk, ancestry and ethnicity tests, as well as access to sport genetic analyses, has become widespread in low- and middle-income countries. Such genetic and genomic practices are carried out by health care institutions and, moreover, direct-to-consumer (DTC) genetic tests are easily available on the internet (Covolo *et al.*, 2015; Phillips, 2016). In Ecuador, a case study by the Red Cross found that rape, intimate partner violence and femicide rates are high. Ecuadorian laws offer mothers the right to ask for a free paternity test; a positive result automatically obliges fathers to provide support for their children. Additionally, genetic tests are routine in Ecuador for police investigating rape cases. For these reasons, increasing knowledge about how DNA can be a link between parents and children or between a sexual offender and a crime seems to be a powerful tool for women's empowerment. Several studies have demonstrated that the understanding and interpretation of personal genomic information is biased by one's own knowledge and appreciation of basic genetic facts, namely, their level of genetic literacy (Hooker *et al.*, 2014; Lea *et al.*, 2011; Lontok *et al.*, 2015; Rafiq *et al.*, 2015). Evidently, an adequate amount of basic knowledge about genetics is essential to understand and interpret the results of genetic and medical analyses. Therefore, various studies have focused on assessing the impact of knowledge of genetics on perception of genetic facts and understanding of disease onset (Haga *et al.*, 2013; Hollands *et al.*, 2016; Lea *et al.*, 2011). Unfortunately, despite the obvious necessity to determine knowledge of genetics, to our knowledge there is no available information regarding this matter in our country. Moreover, recent research has shown differences in quality between public and private higher education institutions in Colombia (Cayon *et al.*, 2017). Therefore, it seems important to assess such differences in Ecuador. The data gathered from these kinds of studies could contribute to the development of programs to reinforce the teaching of genetics to a wider population, which will undoubtedly have a positive impact on national educational programs. Therefore, as a baseline report, we decided to determine the basic knowledge of genetics in undergraduate students in Quito, the capital city of Ecuador. This study provides a glimpse of student perspectives toward genetics and the relation of genetics to disease in a relatively highly educated population based in a developing country. Furthermore, this investigation represents one of the first steps required for building the appropriate strategies to comprehensively assess knowledge of genetics and to ultimately increase the level of genetic literacy in the region.

Methods

Setting, recruitment and questionnaire

Participants' knowledge of genetics was measured as a baseline report on the attitude among undergraduate students toward genetic concepts, who were intentionally chosen because they did

not follow programs involving biologically related courses ($n=350$ by convenience sampling method). The main objective of this research was to assess the competence of students to respond to a validated survey evaluating a minimum, adequate amount of knowledge about genetics (Fitzgerald-Butt *et al.*, 2016). Surveys were carried out from August to October 2018. Individuals were recruited from 3 public and 4 private institutions located in Quito, the capital city of Ecuador. The identity of the institutions was handled in an anonymous form. The participants were approached at random inside the campuses and asked to fill out a questionnaire consisting of 18 statements, provided in Dataset 1 (Larrea, 2019), which measured both the actual knowledge of the associations of genetic conditions with diseases and the actual knowledge of genetic facts. For each question, the results are presented as the percentage of correct answers.

Statistical analysis

Pearson's chi-square test was used to determine the likelihood that the results (answers) supporting the null hypothesis are not due to chance. Additionally, Student's *t*-test was used to assess whether the two groups, composed of publicly and privately educated students, presented any significant differences regarding their measure of knowledge about genetics (assuming equal variances). *P* values are reported using a Type I error level of 0.05, 0.01 and 0.001. All data analyses were carried out with MATLAB® version 9.9.9341360 (R2016a). A MATLAB script to repeat the analysis is available in Dataset 2 (Larrea, 2019).

Ethics approval

This survey was performed under the format of "common social topics". Because of the low-risk nature of the study, approval from a committee was not sought. The participants were informed about the objective of the questionnaire; the survey was voluntary and anonymous, and information that could put the person at risk was not collected. All surveyed students provided prior verbal consent. Written consent was not sought from the participants due to the low-risk nature of the study.

Results

In this research, we present the data gathered as a reference study outlining the knowledge of genetics in undergraduate students. Overall, 350 participants were enrolled in this research (average age: 21.8 years old, SD: ± 2.8); individuals came from diverse backgrounds that did not involve life sciences or medicine. The results varied from 45% to 87% (mean: 66.8%, median: 65%) (Table 1). The responses to each question can be found in Dataset 3 (Larrea, 2019). The percentage scores were higher for the subsection regarding the relationship between genetics and the presence of illness (mean: 68.3%). The lower scores within this section were observed when individuals were asked about the inheritance of diseases (mean: 56%, $p=0.019$) and when questioned about the health status of a person carrying an altered gene (mean: 55%, $p=0.069$). The percentage scores were lower for the subsection regarding genetic facts (mean: 64.9%). In particular, the students seemed to have difficulty answering correctly when asked about the quantity of chromosomes present in humans (mean: 58%, $p=0.004$) and about the number of copies of each chromosome passed down

Table 1. Knowledge of disease related-concepts and genetic facts of undergraduate students with percentages of correct answers.

Disease-related concepts	Total population (n=350)		Private institutions (n=170)		Public institutions (n=180)	
	% correct	p-Value ^a	% correct	p-Value ^a	% correct	p-Value ^a
1. Some diseases are caused by genes, environment and lifestyle. (T)	87	<0.001	89	<0.001	85	<0.001
2. A gen is a disease. (F)	61	<0.001	63	<0.001	65	<0.001
3. Healthy parents can have a child with an inherited disease. (T)	74	<0.001	76	<0.001	71	<0.001
4. A person with altered (mutated) gene may be completely healthy. (T)	55	0.069	57	0.011	53	0.443
5. All serious diseases are inherited. (F)	56	0.019	54	<0.001	59	0.014
6. The child of a person with an inherited disease will always have the same disease. (F)	58	0.002	58	<0.001	58	0.032
7. Altered (mutated) genes can cause disease. (T)	84	<0.001	88	<0.001	81	<0.001
8. A genetic test can tell you if you have a higher chance to develop a specific disease (T)	80	<0.001	82	<0.001	78	<0.001
Average percentage for this section	68.3		70		68.4	
Genetic facts						
1. You can see a gene with the naked eye. (F)	59	<0.001	58	<0.001	61	0.004
2. Genes are instructions for making proteins, which help the body grow and work properly. (T)	57	0.008	61	0.004	54	0.357
3. A gene is a piece of DNA. (T)	77	<0.001	74	<0.001	80	<0.001
4. Genes are inside cells. (T)	69	<0.001	71	<0.001	71	<0.001
5. A chromosome contains many genes. (T)	78	<0.001	77	<0.001	79	<0.001
6. Genes determine traits such as height, eye color and facial appearance. (T)	84	<0.001	82	<0.001	86	<0.001
7. A person has thousands of genes. (T)	73	<0.001	74	<0.001	73	<0.001
8. Identical twins have different sets of genes. (F)	49	0.915	47	0.307	53	0.443
9. Humans have 20 pairs of chromosomes. (F)	58	0.004	53	0.027	61	0.004
10. Parents pass both copies of each chromosome to their child. (F)	45	0.054	49	0.610	41	0.014
Average percentage for this section	64.9		63.8		68.6	
Overall average percentage	66.8		66.6		67	

^ap-values for determining answers provided by chance were calculated using Pearson's Chi squared test. T, true; F, false.

to the next generation (mean: 45%, $p=0.054$). In addition to the lower scores, the hypothesis that the questions were answered correctly without any previous knowledge (provided by chance) could not be significantly rejected. Generally, no differences in the overall knowledge of genetics could be found among students enrolled in private and public institutions ($p=0.9405$). Likewise, no differences between these two groups were observed regarding disease-related questions ($p=0.7844$) and genetic facts ($p=0.7318$).

Discussion

In this report, we portray the percent of correct answers to an 18-item questionnaire measuring a minimum, adequate amount of knowledge about genetics. Overall, this Andes-located population of undergraduate students demonstrated some basic

knowledge toward genetic concepts and their relation to diseases. Nonetheless, student knowledge on facts about genetic proved to be less strong. This tendency was observed in both privately and publicly educated individuals with no significant difference. These results are lower in comparison to the published reports on general populations that have made extensive use of similar survey instruments to determine knowledge about genetics. For instance, Haga & colleagues (2013) found higher scores in a general population based in the US. However, similar scores to those reported here were found by Jallinoja & Aro (1999) in a study performed on a general population in Finland. Furthermore, a group composed of adolescents and young adults suffering from congenital heart disease scored similar results (Fitzgerald-Butt *et al.*, 2016). Notably, the present results are some what higher than those obtained from a Dutch population

suffering from asthma, diabetes mellitus type II and cardiovascular disease (Calsbeek *et al.*, 2007). It is evident that demographic differences may account for the variances in the results. Nevertheless, these results may also imply notable differences between Ecuadorian, US and European science and health education programs (Lontok *et al.*, 2015). To our surprise, the lowest scores obtained were for the two questions involved in how chromosomes are passed down to the next generation. This means that students may not understand the power of genetics to address important issues for the Ecuadorian population, such as determining paternity, solving crimes or understanding our ethnic genetic background. To the best of our knowledge, this study is the first to report a measurement of knowledge of genetics in an Ecuadorian population.

Additionally, the presented results indicate that, despite the relatively adequate scores, the probability that participants were providing correct responses by chance could not be significantly discarded for the overall test and for the two subsections (Table 1). This fact implies that the actual knowledge might be truly weaker than the one asserted by the percentages of correct answers. However, individuals affiliated with private and public universities responded with similar accuracy. The observed average scores might reflect the high level of education of the respondents. It is worth mentioning that the interviewed people did not follow any biologically/medically related courses, which indicates that a non-specialized population may have adequate knowledge about the essential genetic concepts and their involvement in disease. Nevertheless, the participants' knowledge may not be as strong as it appears. As mentioned earlier, the scores do not differ substantially from the earlier studies making use of similar surveys. Nonetheless, the scores were lower than those obtained from a study performed in the US (Haga *et al.*, 2013) where genetic education is constantly improving (Lontok *et al.*, 2015). Based on these observations, a revision of the genetic content covered in educational programs and the implementation of science popularization initiatives seem imperative.

Some limitations of this study should be mentioned. This investigation did not attempt to address the differences in knowledge of genetics among groups classified by characteristics such as sex, ethnic group, age, family history of inherited diseases or level of education. Instead, this study was intended to be focused solely on a general undergraduate population not studying biology or medicine. Furthermore, more universities in different cities should be sampled to have a national perspective on students' insights about genetics. Overall, these results provide a glimpse of the students' standpoint toward genetics and its involvement in disease. Nevertheless, more effort is decisively

needed to design and execute plans that will ensure an optimized method to measure knowledge of genetics in a larger and more diverse population. The data generated using these approaches will be proven essential when designing educational programs involving genetics and health. The application of such programs will be fundamental for the general population to avoid misunderstandings about genetics and to avoid the incorrect utilization of scientific terms.

Follow-up studies will try to explore the knowledge about genetics and the attitudes toward related subjects, including genetic testing, stem cells, regenerative medicine and genetically modified organisms (GMOs). The expected results will provide improved insight into the population's knowledge and will serve as a foundation for developing better strategies to increase the level of genetic literacy in our community.

Data availability

Extended data

Open Science Framework: Assessing genetic knowledge in Ecuador. <https://dx.doi.org/10.17605/OSF.IO/ZUVMN> (Larrea, 2019)

This project contains the following extended data:

- Dataset 1.pdf (the questionnaire in the original Spanish language and its translation into English)
- Dataset 2.pdf (the MATLAB script to reproduce the analysis)

Underlying data

Open Science Framework: Assessing genetic knowledge in Ecuador. <https://dx.doi.org/10.17605/OSF.IO/ZUVMN> (Larrea, 2019)

This project contains the following underlying data:

- Dataset 3.csv (a spreadsheet containing all responses to the evaluation)

Grant information

The author(s) declared that no grants were involved in supporting this work.

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Current Peer Review Status:

Version 1

Reviewer Report 21 May 2019

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Rebecca Carver

Department of Communication, Norwegian Institute of Public Health, Oslo, Norway

This study provides valuable insight into the level of knowledge of genetics in an Ecuadorian population.

Introduction

Be careful with terminology regarding “perspectives” towards genetics, as this is not the same as knowledge about genetics. The same goes for “attitudes”.

Study design

The first sentence in the methods section does not make sense. How is this a “baseline report on the attitude among undergraduate students toward genetic concepts?” What do the authors mean? Knowledge is not the same as attitudes.

The authors of the chosen instrument (Fitzgerald-Butt et al 2016¹) have stated in their paper that the instrument can be used for older teenage and young adult patients and parents in the pediatric setting, and that is most informative for individuals with below average genetic knowledge. How is this instrument applicable to the current study's Ecuadorian student population, which is a very different sample population than the one for which the instrument was developed? What were the reasons for choosing this particular instrument? Can the authors justify their choice of instrument?

Results

What percentage of correct answers is “adequate” in the chosen instrument? Is there a reference value in the original instrument that can be used to compare the results of this survey? (E.g. what minimum percentage of answers should the respondents answer correctly in order to have an adequate amount of knowledge?).

Discussion

The results show that students achieved slightly better results when asked about genetics and diseases than when asked about genetic facts. One possible explanation for this could be that the

questions about disease relate more to people's lives than genetic facts. The authors state that they were surprised by the lower score obtained on a question about how chromosomes are passed to the next generation, but students may regard this type of "textbook knowledge" as less relevant to their everyday lives, and thus be less inclined to remember it. There is a lot of literature on how the genetics content in textbooks is lagging behind modern scientific developments – e.g. see previous literature by Gericke et al. and Dougherty et al. – and this may be an explanation for the results in this study. The authors might like to comment on this.

Conclusions

The authors conclude by saying that despite the relatively adequate score overall (66.8% correct answers), the possibility that the correct responses were given by chance cannot be ignored – and that the actual knowledge of genetics among the participants might be less than that revealed by their answers. Consequently, the authors assert; "it seems evident that the national education programs of Ecuador require improvement in teaching of genetic concepts". However, if the responses were given by chance, is it not also possible for the actual knowledge of genetics among the participants to be *higher* than that revealed here? Could the authors comment on this?

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Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Partly

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Yes

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Partly

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Science communication, Public understanding of Genetics, Science Education, Public Health and Survey design

I confirm that I have read this submission and believe that I have an appropriate level of

expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Reviewer Report 07 May 2019

<https://doi.org/10.5256/f1000research.20127.r47621>

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Vasiliki Mollaki

Hellenic National Bioethics Commission, Athens, Greece

The study aimed to assess the knowledge of genetics in a selected population of students from both public and private institutions.

The work is described in sufficient detail (regarding the samples size, study design and statistical analysis). The authors acknowledge the study limitations.

Although similar studies have been performed in other countries, this is the first study assessing genetic knowledge in Ecuador. The result of the study could be proved useful for policy makers and educational programs.

Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Yes

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Yes

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Human genetics, molecular genetics, bioethics

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

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