Characteristics of health information channels usage by

**Deaf and hard of hearing people in Israel** / Liat Winter

Abstract

As technology and communications have developed, the methods used for seeking

information in many fields have also changed. Health is one of the fields that has

experienced change because of technological developments, and one expression of

these changes is the increase in health care issues.

The issue of health inequality has been recognized for 150 years, but only in recent

decades have there been efforts to reduce health inequality. Health inequality appears

in disadvantaged populations, including ethnic and linguistic minorities, and also

among people with physical and sensory disabilities. The Deaf and hearing-impaired

are among the populations whose health is adversely affected by health inequality.

This inequality is evident, among other things, in their ability to obtain, understand

and use health information. The Deaf and hearing-impaired (especially those who use

Israeli sign language) may suffer impaired health due to the difficulty they have

accessing health information, as a result of accessibility and communication

problems.

Goal

The aim of this study was to examine of how Deaf and hearing-impaired people seek

health information. Do they derive benefit from the health information they find?

What are their preferred information channels? What are the leading health

information topics that interest the research population? This study did not deal with

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the overall information behavior of the Deaf, but rather focused on one important

component of their information behavior, namely, the selection and evaluation of

information channels.

**Research Methods** 

The study was conducted among 79 Deaf and hearing-impaired people (of whom 49

identified themselves Deaf, and 30 as hearing-impaired). It was a quantitative study

based on a questionnaire that was available in two versions. The first, an online form

created in Google Forms, was shared through Facebook groups of Deaf and hearing-

impaired people, by e-mail, and among friends. The second, print version was

distributed to people who do not regularly use Facebook, had difficulty filling out the

online questionnaire, or rarely use a computer.

The questionnaire examined the effect of independent variables (gender, marital

status, age, income level and hearing-related variables such as hearing level, type of

communication, reading literacy, and level of education) on the dependent variables

concerning information channels, both traditional (newspapers, books, advertising

brochures, television, family, friends, health professionals) and online (social

networks, health websites, forums, videos). To compare the responses of the two

research populations (Deaf and hearing-impaired) on these subjects, the questionnaire

was analyzed statistically using measures of frequency, standard deviation, and t-tests

for independent variables for the questions relating to computer knowledge and use,

and health questions.

**Findings** 

The findings showed that the leading health information subjects sought by

participants were general topics that also interest the general population (Information

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on drugs, making appointments, diseases, physicians, hospitals, healthy lifestyle),

followed by topics specific to the research population (e.g. ENT specialists, Cochlear

implant, hearing test and speech speech-language therapists). Study participants

sought health information for their parents, children, themselves and, to a lesser

extent, friends. Most of the Deaf and hearing-impaired participants communicate

using speech, because most were born to hearing families, use speech to communicate

with their families and environment, and were educated in individualized,

mainstreaming programs.

Participants search for health information using online information channels, informal

and formal channels, as well as traditional channels (radio, telephone, facsimile,

television, print newspapers, libraries, print books, print leaflets). The preference for

using online channels (social networks, health websites, forums, videos) can mostly

be explained by the problem of accessibility presented by traditional channels,

because people with severe hearing loss cannot use hearing aids, and these channels

do not include accessibility features such as subtitles or interpretation in Israeli sign

language. An additional reason relates to the low level of literacy, and poor ability to

understand material written for and targeting a more literate population.

Within the study population, participants with severe hearing loss had difficulty using

traditional channels that do not include accessibility features such as subtitles or

interpretation in Israeli sign language. The Deaf participants preferred to seeking

information on the Internet over going to the doctor. By its nature, the internet is an

information source that does not require hearing, where they can be express

themselves freely; understanding speech is not a prerequisite, and the information is

accessible from anywhere, at any time. In the real-world, these factors create

difficulties for some members of the research population, but internet blurs their

difficulty. However, hearing-impaired people who can use the auxiliary aids for

communication do use traditional channels also.

Moreover, it was found that the level of knowledge and use of computers and the

internet among the Deaf and hearing-impaired was good; participants know how to

evaluate the quality of a website, and adopt every technological innovation. They use

e-mail, social networks and read the news. There were no differences in level of

computer and internet use between the Deaf and hearing-impaired.

In other findings regarding a majority of the research population: more women than

men sought health information; their average age was 42-53, born in Israel, married,

up to 3 children, educated in individualized mainstreaming programs, were Deaf or

hearing-impaired from birth or age of two years, have severe hearing loss, have

difficulty speaking on the phone, believe that listeners understand their speech, prefer

to speak Hebrew, self-evaluated their literacy for reading and writing Hebrew as very

good but very low in English. Research participants who defined themselves as Deaf

were characterized by a better command of Israeli sign language than the hearing-

impaired. The Deaf had more Deaf/hearing-impaired relatives, and more contact with

other Deaf/hearing-impaired people.

**Conclusions and Solutions** 

It is necessary to increase public awareness, and expand the use of subtitles and sign

language in traditional media. (Among older Deaf people, there may be a problem

with literacy, and it would be desirable to have professionals in the community adapt

publications and information to be linguistically and culturally appropriate.) Creating

a free database of reliable health resources built specifically for the Deaf and hearing-

impaired population should be considered.

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Encouraging Deaf speakers of Israeli sign language and hearing-impaired people to

pursue careers in public health, health research and health care should be considered,

thereby creating collaboration between the public health community, and the Deaf and

hearing-impaired community, which ought to create more effective ways of

disseminating health information, and for developing appropriate and accessible

programs.

Despite the similarities between Deaf and hearing-impaired populations, some

differences were found between the two. The differences included:

Physician-patient relationship. Absent good communications between doctors and

patients, uncertainty leads patients to seek other information channels, including the

internet. Compared to the Deaf, the hearing-impaired have better communications

with doctors; therefore, they did not seek additional information to fill-in gaps in

understanding that remained after an appointment.

Seeking health information. On a question regarding their internet search preferences,

Deaf people preferred searching the internet because hearing is unnecessary, and they

use it as a substitute for going to a doctor. The Deaf also estimated that they received

more information from family and friends, while the hearing-impaired estimated that

they received more information from information centers, organizations,

audiologists/speech-language therapists.

Possible solutions may include training medical teams in ways of communicating

with the Deaf and hearing-impaired. These solutions must be provided by the public

health authorities, who must work with representatives of the Deaf and hearing-

impaired population to adapt solutions for the Deaf and hearing-impaired population.

Among the proposed solutions: providing a basic course in Israeli sign language for

medical teams; assistance by a professional interpreter trained in medical

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interpretation, as necessary and at the patient's request; and using written notes. Care

should be taken to ensure clear speech, with good lighting and no background noise,

not hiding lips, removing masks, facing the patient to facilitate lip reading, using

vocabulary without complicated medical terms, and pictures if necessary; requesting

the assistance of a family member; and remembering that the Deaf or hearing-

impaired person is the patient and speaking directly to him rather than the assistant

(family, friend or interpreter).

Another solution involves conducting surveys that include useful medical data and

facilitate future research. This data would allow creating programs to prevent chronic

diseases, and improve the health status of the Deaf and hearing-impaired population,

according to appropriate priorities, and evaluate the effectiveness of these programs.

It is also necessary to create a training program for sign language interpreters who

specialize in health, know medical terminology and translate it effectively for Deaf

patients. They must also be aware of confidentiality requirements. It is important to

note that it is necessary to define the research populations uniformly and use

terminology for who is Deaf and who is hearing-impaired consistently, in order to

ensure more accurate and consistent studies.

With all these steps, we can hope that Deaf and hearing-impaired people will be better

able to seek and benefit from health information, thereby improving.

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