Digital technology use among disadvantaged Australians: implications for equitable consumer participation in digitally-mediated communication and information exchange with health services

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Abstract

Objective. To present research findings on access to, and use of, digital information and communication technologies (ICTs) by Australians from lower income and disadvantaged backgrounds to determine implications for equitable consumer access to digitally-mediated health services and information.

Methods. Focus groups were held in 2008-09 with 80 residents from lower income and disadvantaged backgrounds in South Australia, predominantly of working- and family-formation age (25 to 55 years). Qualitative analysis was conducted on a-priori and emergent themes to describe dominant categories.

Results. Access to, and use of, computers, the Internet and mobile phones varied considerably in extent, frequency and quality within and across groups due to differences in abilities, resources and life experience. Barriers and facilitators included English literacy (including for native speakers), technological literacy, education, income, housing situation, social connection, health status, employment status, and trust. Many people gained ICT skills by trial and error or help from friends, and only a few from formal programs, resulting in varied skills.

Conclusion. The considerable variation in ICT access and use within lower income and disadvantaged groups must be acknowledged and accommodated by health initiatives and services when delivering digitally-mediated consumer-provider interaction, online health information, or online self-management of health conditions. If services require consumers to participate in a digitally-mediated communication exchange, then we suggest they might support skills and technology acquisition, or provide non-ICT alternatives, in order to avoid exacerbating health inequities.

1. What is known about the topic? Government and health provider use of digitally-mediated information and communication is rapidly increasing. However, national data show that ICT access is distributed unevenly across Australia’s population. Furthermore, this distribution mirrors the health gradient. There is little qualitative data on the extent to which, and ways in which, ICTs are used within lower income and disadvantaged groups - those with greater health need.

2. What does this paper add? This paper augments the scant literature to describe ICT access and use in a range of lower income and disadvantaged groups. It indicates barriers and facilitators, and highlights the need for formal supports to level up the whole population to have the skills, confidence and resources to use and benefit from ICT-mediated communication.

3. What are the implications for practitioners? As health services and governments increase the level of digitally-mediated information and communication connection with consumers/patients, it is important to understand and find ways to address differential consumer access to and use of ICTs, so that equity of access to services and information is promoted. This is particularly important as lower income and disadvantaged groups are likely to have both poorer health and lower ICT use.

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Introduction

The ability to access information and communication technologies (ICTs), particularly via the Internet and mobile phones, is increasingly vital to full participation in economic, social, and political life.1-4 This is particularly so as institutions move towards greater ICT-mediated provision of services, support and
information. There is also increasing focus on encouraging digitally-mediated consumer-to-provider and consumer-to-consumer connection in the health sector (within ‘e-health’), and more recently through ‘m-health’ (via mobile devices, including mobile phones).

The spread of ICTs hides the uneven distribution of digital access across Australia’s population, for example ~28% of households have no home Internet access, although 82% of Australians aged 14 years or older use a mobile phone and 31% of children aged 5–14 years own one. Socioeconomic factors influence these distributions. For example, only around 7% of households earning $120 000 or more are without home Internet, whereas 42% of households earning less than $40 000 are without home Internet. Similar inequities are evident by geographic area.

However, connection data do not explain the extent to which ICTs are actually used and how this may, in turn, influence health. Australia’s National E-Health Strategy intends to ‘transform the way healthcare professionals practice and consumers interact with the health system’, to ‘reduce costs and demands on the health system’. The Strategy aims to ‘empower consumers’ by encouraging ‘electronic access to the information needed to better manage and control their personal health outcomes’, and to focus on ‘those segments of the population that interact frequently with the health system’. However, the Strategy does not acknowledge that those more likely to interact with the health system are those lower down the social gradient with less means to use ICTs.

In Australia, there is little qualitative research exploring digital access and use (or non-use) among lower socioeconomic status (SES) groups, or considering the implications for equitable consumer access to digitally-mediated health services. Much research on e-health is technically or provider-focussed, with an excessive focus on ‘new and exciting opportunities to empower individuals’ (e.g. through electronic health records) without explaining how benefits will result. Indeed, more critical approaches find potential for increased discrimination and inequality as digital inequalities are intertwined with other inequalities, such as poverty and remoteness, and parallel the uneven distribution of chronic disease. Furthermore, some assume that all consumers are abandoning traditional media channels and that the Internet enables ‘unfettered access’ to information for all. However, the small amount of research on non-users finds those with lower education and income being least likely to have access, for example, to Internet-based medical information and support.

Methods

Our study is a primary analysis of data from 80 individuals in nine focus groups. Between August and November 2008, six groups were held across Adelaide with 55 participants exploring ICT use, and in November 2009, three groups were held with 30 participants about Internet access via mobile phones. Focus groups are valuable for research with lower SES or ‘vulnerable’ populations because they give ‘voice’ to the participants, allowing them to define what is important to understand their experiences. Local service workers provided recruitment assistance in lower SES areas, as per Glover et al. and Australian Bureau of Statistics. Participants were aged 25 to 55 years and recruited from: a women’s support group; two men’s support groups; an employment support group; a community housing group; an Aboriginal students group; and an African recent-refugee group. Research approval was obtained from the Flinders University Social & Behavioural Research Ethics Committee.

A semi-structured discussion schedule was developed based on a literature review and previous surveys and allowed participants to raise additional issues. All but six participants also agreed to complete a short survey with demographic and technological information. Questions relevant to this paper were on ICTs being used or not; description of use; ways of learning; perceived facilitators and barriers; and preferred contact methods with service providers. Assistance was offered to participants to read project information and forms, and the refugee group used translated documents and an interpreter. Discussions were recorded and transcribed. Participants were thanked with lunch and a $30 voucher.

Transcripts were analysed according to established methods to provide a descriptive account. Two of the authors conducted all groups, which allowed continual data immersion; they then used a constant comparative, iterative method to analyse two transcripts, individually allocated text to a-priori and emergent codes, and compared and discussed interpretations to derive a coding framework. This framework was applied to the other transcripts and new emergent codes were discussed and added. As analysis proceeded with subsequent groups, new questions were incorporated into the schedule to further explore emerging ideas. The final framework consisted of dominant categories.

Results

Socioeconomic and demographic profile of participants

The survey confirmed that participants represented lower SES and disadvantaged groups: three-quarters had household incomes equal to or less than average; two-thirds were living with some perceived financial strain; two-fifths were educated to year 11 or less; most were in lower level occupations; 1 in 10 were unemployed; almost all resided in lower SES areas; three-quarters lived in rental housing; over one-quarter were Aboriginal; 1 in 10 were sole parents; and two-fifths spoke a language other than English at home.

ICT access and use

Access to, and use of ICTs – computers, Internet, mobile phones – varied considerably in extent, frequency and quality, as did perceived effectiveness: ‘I do have a computer at home but it sits in the bedroom collecting dust. I’ve got the Internet on but I never use it’; ‘My computer I am on 90% of the time. I use it for anything from Internet banking; we purchase things worldwide from various websites, check the weather, my TV guide’; ‘Don’t have anything to do with the Internet. I don’t have a lot of technology stuff’. People often found mobile phones cheaper to buy and maintain, and easier to use, than a computer, although a few shared mobiles (e.g. with a partner), and one-third had no
landline home phone. Although some had no Internet access, others mentioned a range of access opportunities, including home, work, college, or community locations: ‘I’ve had dial-up for years now and I’m just used to it. If I need something faster I go here [common room in community housing] or to [homeless women’s centre].’ Nevertheless, those with home or community access did not necessarily get time to use this, have skills to use it well, or choose to use it all the time. Many could not afford Internet access at home or on their mobile phone, yet public access varied in usefulness and availability: ‘I’ve got another 18 months to go of [home] broadband, then I think I’ll start using the library. Costs, yes’; ‘The library is good because it’s got free Internet, but some people go there every day [and] they’ve booked it up [so] that you have trouble getting time’; ‘At [disability support centre] I’d feel like... people would be watching me’. Furthermore, women with young children found public computers hard to use without a crèche, and one single dad needed home access to search for work once his daughter was asleep at night.

There were also across-group differences in ICT use. For example, younger Aboriginal participants described difficulties with intergenerational communication, as grandparents would not answer calls from mobile phones, which was the younger people’s preferred communication method. The refugee group was the only group to discuss high costs using phone cards to call overseas; they also criticised Australian government websites (e.g. Immigration) for not clearly providing non-English language options, and used the Internet to access news about Australia from non-Australian sources. Some older participants appeared to have greater difficulty with technological literacy and manual dexterity, but some participants in their 20s also had barriers to use (e.g. affordability), with one young woman still using her local payphone.

How people learned to use technologies
Most technology use was self-taught or via friends, so that those with few social connections were at a distinct disadvantage. A smaller number gained skills through school or workplaces. Others reported reluctance to learn, were fearful of technology, or felt appropriate training (i.e. respectful, non-judgmental, one-on-one needs-based training at familiar places) was unavailable.

Basic skills, such as mobile texting and Internet searching, were difficult or off-putting, including for some younger people. Therefore, lack of confidence, skills or resources often restricted use and limited the benefit people derived: ‘I have a mobile but I only know how to answer it. Going into and sending messages, I wouldn’t have a clue’.

Preferred service contact
In light of barriers to use, participants varied in their willingness to have digitally-mediated contact with services. Some liked electronic contact for straightforward transactions but wanted personal contact for complex queries: ‘You can just jump online for Centrelink and find out how much your next pay is’; ‘I don’t have my [driving] licence so it’s better for me to [do banking] over the phone’; ‘I applied for TAFE just a couple of weeks ago through the Internet’. However, for a high proportion of participants, only traditional contact was effective: ‘I prefer to ring up because they explain it to me better than what is written on the Net’; ‘Sometimes you’ve got a question, which in a pile of writing you’ve got to search for. But if you ask a question with someone face-to-face - there’s your answer’. The move to electronic contact was interpreted by some as cost-shifting to the consumer rather than improving services, and those without the resources or desire to change also felt that direction to increased digital contact (e.g. ‘Visit our website’) caused stigmatisation and stress: ‘It implies you should be connected – you’re no-one if you aren’t’. However, Internet or mobile contact from service providers (e.g. text reminders for appointments) was welcomed by customers who gave authorisation.

Discussion
Although digital technology use is increasing rapidly across Australia, this is not occurring equally across the population. This has implications for equitable access to health services and information. Among our research participants, some did not use ICTs, and those that did varied in their quality, type and frequency of use. Our findings suggest that initiatives which increase digital contact with or between consumers, on the assumption of improved service efficiency and effectiveness, should assess whether the whole consumer population has the resources and skills to benefit from such contact, and then provide resources and training to fill identified gaps. That is, using Dahlgren and Whitehead’s
term, to ‘level-up’ less privileged groups across the social gradient.

Despite the almost universal ownership of mobile phones, many in our study perceived they had insufficient income to make mobile calls, limiting mobiles as a reliable communication option. For example, until 1800 numbers become free from mobile phones, services could consider how to support mobile-free calling such that a 1800 Smoking Quitline, for example, is affordable for the mobile-only population. Otherwise, such strategies will disproportionately affect lower SES groups, who are more likely to smoke. As a positive example, South Australia’s Royal District Nursing Service established and paid for videophones and Broadband connections to introduce virtual nurse visiting in clients’ homes, which saved resources compared with personal visits.

Although there is a national focus on digital up-skilling for particular sub-groups (e.g. Aboriginal, older, culturally and linguistically diverse, remote Australians), there seems less focus on the mainstream, lower income, working age population whose levels of literacy, trust and confidence may also be low. We therefore suggest services consider the communication needs of their whole population. Researchers could further explore consumer needs in mainstream and sub-groups to identify the best ways to fill ‘communication gaps’.

Our findings also suggest that services should be aware that pressuring unprepared or unwilling consumers to use ICTs can further undermine health by creating stigma, distrust and feelings of losing control. A reasonable conclusion in this context is that ‘encouraging consumers to use electronic access’ could not only undermine health in lower SES groups, but that encouragement (or coercion) into digital communication may cause them to opt out of services, putting their health in further jeopardy.

Conclusion

Our study findings augment the scant literature to provide insight into the implications of increased ICT-mediated health services for disadvantaged groups. Although qualitative findings from non-random selected groups are not generalisable, this limitation was partly addressed by our participants having broadly representative characteristics of the more disadvantaged in the Australian population. Our study suggests that ICT-mediated health initiatives should be considered in more complex and diverse ways to ensure that health inequities are not exacerbated by digitally-mediated communication methods, excluding those who lack skills, resources or capabilities to use ICTs. Unless lower SES groups receive appropriate support (e.g. skills training, funded equipment), ICT-mediated communication may become a new barrier to health service access. Consumers should be included in communication planning, rather than taking a ‘one-size-fits-all’ approach, and further research on the topic is indicated. Without such compensatory measures, Australia’s E-Health Strategy may end up increasing inequities in health service access.

Competing interests

The authors disclose that they have no financial or personal relationships with organisations that could inappropriately influence their work, nor do they have any financial or other support which may pose a competing interest. The source providing funding agrees to the publication of this material, but had no influence over the way in which the data in this paper were analysed or interpreted. Dr Newman discloses that she is an unremunerated member of the Australian Communications Consumer Action Network (ACCAN).

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