



BMJ Open is committed to open peer review. As part of this commitment we make the peer review history of every article we publish publicly available.

When an article is published we post the peer reviewers' comments and the authors' responses online. We also post the versions of the paper that were used during peer review. These are the versions that the peer review comments apply to.

The versions of the paper that follow are the versions that were submitted during the peer review process. They are not the versions of record or the final published versions. They should not be cited or distributed as the published version of this manuscript.

BMJ Open is an open access journal and the full, final, typeset and author-corrected version of record of the manuscript is available on our site with no access controls, subscription charges or pay-per-view fees (<http://bmjopen.bmj.com>).

If you have any questions on BMJ Open's open peer review process please email [info.bmjopen@bmj.com](mailto:info.bmjopen@bmj.com)

# BMJ Open

## Breast cancer screening based on personalized risk assessment: Experience and satisfaction of primary care providers with receiving risk communication documents for patients enrolled in the PERSPECTIVE I&I project

Journal:	<i>BMJ Open</i>
Manuscript ID	bmjopen-2024-093936
Article Type:	Original research
Date Submitted by the Author:	19-Sep-2024
Complete List of Authors:	<p>Omeranovic, Arian; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Lapointe, Julie; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Fortier, Philippe; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Bergeron, Anne-Sophie; Université du Québec à Rimouski - Campus de Lévis, Département des sciences infirmières; Research Centre of the Chaudière-Appalaches Integrated Health and Social Services Centre</p> <p>Dorval, M; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center; Research Centre of the Chaudière-Appalaches Integrated Health and Social Services Centre</p> <p>Chiquette, Jocelyne; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Boubaker, Asma; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Eloy, Laurence; Ministère de la Santé et des Services sociaux du Québec, Programme québécois de cancérologie</p> <p>Turgeon, Annie; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Lambert-Côté, Laurence; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Joly, Yann; McGill University, Center of Genomics and Policy; McGill University Faculty of Medicine, Human Genetics Department and Bioethics Unit</p> <p>Brooks, Jennifer; University of Toronto Dalla Lana School of Public Health</p> <p>Walker, Meghan J. ; University of Toronto Dalla Lana School of Public Health; Ontario Health</p> <p>Stockley, Tracy; University Health Network, Laboratory Medicine Program</p> <p>Pashayan, Nora; University of Cambridge, Department of Public Health and Primary Care</p> <p>Antoniou, Antonis; University of Cambridge, Department of Public Health and Primary Care</p> <p>Easton, Douglas; University of Cambridge, Department of Public Health and Primary Care</p> <p>Chiarelli, Anna; University of Toronto Dalla Lana School of Public Health; Ontario Health</p> <p>Knoppers, Bartha; McGill University, Center of Genomics and Policy</p>

	Simard, Jacques; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center; Laval University Faculty of Medicine, Department of Molecular Medicine Nabi, Hermann; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center; Laval University Faculty of Medicine, Department of Social and Preventive Medicine
Keywords:	Epidemiology < ONCOLOGY, Breast tumours < ONCOLOGY, PUBLIC HEALTH, Cancer genetics < GENETICS
Note: The following files were submitted by the author for peer review, but cannot be converted to PDF. You must view these files (e.g. movies) online.	
Supplementary file 3.zip	

SCHOLARONE™  
Manuscripts



I, the Submitting Author has the right to grant and does grant on behalf of all authors of the Work (as defined in the below author licence), an exclusive licence and/or a non-exclusive licence for contributions from authors who are: i) UK Crown employees; ii) where BMJ has agreed a CC-BY licence shall apply, and/or iii) in accordance with the terms applicable for US Federal Government officers or employees acting as part of their official duties; on a worldwide, perpetual, irrevocable, royalty-free basis to BMJ Publishing Group Ltd ("BMJ") its licensees and where the relevant Journal is co-owned by BMJ to the co-owners of the Journal, to publish the Work in this journal and any other BMJ products and to exploit all rights, as set out in our [licence](#).

The Submitting Author accepts and understands that any supply made under these terms is made by BMJ to the Submitting Author unless you are acting as an employee on behalf of your employer or a postgraduate student of an affiliated institution which is paying any applicable article publishing charge ("APC") for Open Access articles. Where the Submitting Author wishes to make the Work available on an Open Access basis (and intends to pay the relevant APC), the terms of reuse of such Open Access shall be governed by a Creative Commons licence – details of these licences and which [Creative Commons](#) licence will apply to this Work are set out in our licence referred to above.

Other than as permitted in any relevant BMJ Author's Self Archiving Policies, I confirm this Work has not been accepted for publication elsewhere, is not being considered for publication elsewhere and does not duplicate material already published. I confirm all authors consent to publication of this Work and authorise the granting of this licence.

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

**Breast cancer screening based on personalized risk assessment: Experience and satisfaction of primary care providers with receiving risk communication documents for patients enrolled in the PERSPECTIVE I&I project**

**Word count: Abstract= 249; document = 3180**

**Total number of figures and tables = 4**

For peer review only

## Abstract

**Objective:** We aim to document primary care providers' (PCPs) experience and satisfaction with receiving risk communication documents on their patient's breast cancer (BC) risk assessment and proposed screening action plan.

**Methods and Analysis:** A survey was sent to all 763 PCPs designated by participants in PERSPECTIVE I&I research project, about one to four months after the receipt of risk communication documents. Descriptive analyses were used to report on participants' responses. Responses to two open-ended questions were subjected to content analysis.

**Results:** A total of 168 PCPs answered the survey, from which 72.6% reported being women and 74.4% having more than 15 years of practice. Relatively few (38.1%) were familiar with the risk-based BC screening approach prior to receiving their patient risk category. A vast majority (85.7%) agreed with the screening approach and would recommend it to their patients if implemented at the population level. PCPs reported understanding the information given to them (92.3%) and agreed with the proposed BC screening action plan (88.7%). Some PCPs suggested to simplify the materials, to stay mindful of the fact that the approach could increase their workload, and to invest efforts in the planning of professional training.

**Conclusion:** PCPs displayed positive attitudes about a risk-based BC screening approach and were satisfied with the information provided. They also highlighted that if the approach were to be implemented at the population level, it would be important to address issues such as professional training, impact on workload. Future qualitative studies may help further characterize PCPs' perspectives.

**Keywords:** Risk-based breast cancer screening, risk assessment, primary care providers, polygenic risk score.

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

**Strengths and limitations of this study:**

- This study is among the first to report on the experience and satisfaction of PCPs receiving patients’ personalized BC risk assessment and proposed action plans in order to adapt screening for their individual patients in a real-life scenario. This makes it possible to collect real-life PCPs feedback compared to feedback based on hypothetical scenarios.
- The main limitation of this study is related to its relatively low response rate, which resulted in a limited sample size. Such a sample cannot be deemed to represent the PCPs population at large.
- Our small sample size also prevented us from conducting more complex multivariable analyses.

Enseignement Supérieur (ABES) .  
Protected by copyright, including for uses related to text and data mining, AI training, and similar technologies.

## Introduction

According to the World Health Organization, more than 2.3 million women were diagnosed with breast cancer (BC) in 2020, leading to more than 600,000 deaths <sup>1</sup>. In Canada, BC is the second most commonly diagnosed cancer and more than 28,600 women were diagnosed with this disease in 2022 <sup>2</sup>. Fortunately, the death rate of from BC has steadily declined since its peak in 1986 <sup>3 4 5</sup>. Such a decline in mortality can be attributed to improved treatments and more efficient screening programs <sup>6 3</sup>.

While the current age-based BC screening programs have been successful in reducing mortality, there are still areas for improvement <sup>7</sup>. This includes reducing BC overdiagnosis <sup>8</sup>, which can have several consequences such as unnecessary medical exams, treatments, and psychological impacts on patients <sup>9</sup>. Moreover, age-based screening recommendations ignore several BC risk factors, such as genetic susceptibility, lifestyle habits, or reproductive history <sup>10</sup>. Evidence suggests that a more personalized risk-based approach could be a cost-effective way to improve BC screening programs <sup>11 12 13 14 15</sup>. This personalized approach involves targeting women at the highest risk for developing BC. <sup>15</sup> First, such risk stratification is expected to allow for reducing BC mortality through early detection of tumors in high-risk patients, thereby significantly increasing the chances of effective therapeutic management, cure, and long-term survival <sup>15</sup>. Moreover, by focusing screening efforts on specific populations, this personalized approach would lead to a more rational and cost-effective allocation of limited healthcare resources, representing a significant benefit in terms of cost optimization and spending efficiency within the healthcare system <sup>13 15</sup>.

Although a personalized risk-based approach appears promising, its implementation does represent a challenge <sup>14 16</sup>. Part of this challenge concerns the coordination of health services through adequate preparation of, and efficient communication with, primary care providers (PCPs) <sup>14</sup>. Several studies indicated that PCPs seem to have positive attitudes towards the implementation of risk-based assessment for BC <sup>17 18 19 20</sup>. The use of genomic technologies for multifactorial risk assessment in other types of cancer also seems to be generally well received by various PCPs <sup>21 22 23</sup>. However, they tend to report a lack of training in conducting BC risk assessment <sup>24</sup>. Other barriers to implementing a risk-based screening approach were identified, such as an increased workload, a lack of financial and human resources, and a lack of coordination between public and private PCPs <sup>25</sup>. Primary healthcare professionals, such as nurse practitioners and family



1  
2 physicians, are in a prime position to facilitate the implementation of BC screening approaches tailored to  
3  
4 each patient's individual risk level <sup>20</sup>. Their role is essential on several instances <sup>20 26 27 28</sup>. First, they are  
5  
6 expected to clearly explain to their patients the advantages and disadvantages of different screening  
7  
8 methods based on personal risk assessments. Second, to able to effectively interpret and communicate each  
9  
10 patient's calculated risk level using specific prediction tools. Finally, to advise their patients on the most  
11  
12 appropriate screening and prevention strategies based on their individual risk profile. In definitive, PCPs in  
13  
14 this context are expected to have the crucial responsibility of educating, raising awareness, and guiding their  
15  
16 patients towards the screening options best suited to their personal risk of BC <sup>20</sup>.  
17  
18

19  
20 Most previous studies collected PCPs' opinions and attitudes on hypothetical implementation  
21  
22 scenarios <sup>17 18 21 23 29</sup>. To our knowledge, very few evaluated the experience and satisfaction of PCPs on the  
23  
24 actual receipt of a patient's personalized BC risk category in real-life practice. This important feedback is  
25  
26 needed to guide future implementation efforts <sup>30</sup>.  
27  
28

29  
30 Our study aimed therefore to document the experience and satisfaction of PCPs relative to the  
31  
32 receipt of information on their patients' personalized BC risk category and proposed screening action plan in  
33  
34 their real-life practice in the context of the PERSPECTIVE Integration and Implementation study <sup>16 31</sup>.  
35  
36

37  
38 **Materials and Methods**

39  
40 **Setting**

41  
42 The present study is part of a major Canadian research project entitled PERSPECTIVE: I&I (Personalized Risk  
43  
44 Assessment for Prevention and Early Detection of Breast Cancer: Integration and Implementation), which  
45  
46 aims to improve breast cancer risk assessment and determine optimal approaches for implementing risk-  
47  
48 based screening and prevention within the Canadian health system <sup>32 16</sup>. This project included a pre-  
49  
50 implementation research activity recruiting more than 3,750 women from Quebec and Ontario, Canada's  
51  
52 two most populous provinces <sup>31</sup>. Participating women underwent a comprehensive BC risk assessment using  
53  
54 the Breast and Ovarian Analysis of Disease Incidence and Carrier Estimation Algorithm (BOADICEA) model  
55  
56 implemented in the CanRisk prediction tool. This tool estimates participants' 10-year BC risk using the  
57  
58 polygenic risk score (PRS) and multiple risk factors, namely age at menarche, age at menopause, number of  
59  
60

children, age at first live birth, use of oral contraception, use of hormone replacement therapy, body mass index, height and alcohol use<sup>33 34 35,36</sup>. By identifying and combining common, low-penetrance genetic variants, PRS is considered as useful tool for estimating the genetic risk of developing disease at both individual and population levels<sup>37 36 38 39</sup>. The information from the risk assessment was then used to inform patients about their risk category and of possible screening action plan. In Quebec province only, the risk communication documents were sent to both the women and their designated family physician or primary care nurse practitioner. This includes a risk letter that reported on women's 10-year estimated risk, stratified into three risk categories using age-dependent risk thresholds. The remaining lifetime risks (from age 30 to 80 years) for these three categories—referred to as "average," "higher than average," and "high"—are based on percentages of less than 15%, 15%–24%, and more than 25%, respectively<sup>16</sup>. It also included the proposed screening action plan based on that risk category. Finally, it also includes a 2-page information booklet on the study, risk assessment, the importance of discussing their risk level with their patient, and a follow-up decision tree detailing the proposed action plan based on risk category<sup>16,32</sup>. All the documents are available in the Supplementary files.

### Design and participants

All family physicians and primary care nurse practitioners designated by each of the 1,642 women participating in PERSPECTIVE I&I in the province of Quebec were sent an invitation letter and the survey. They were mailed about one to four months after the letter informing women of their risk category and the corresponding screening action plan were sent out. To increase participation, two additional reminders were sent by fax one to six months after the initial mailing. The recruitment phase took place from July 2021 to July 2022. Participants were consented by completing the questionnaire. In addition, we specified the terms of confidentiality and participation in the first paragraph of the survey, while also providing a telephone number and e-mail address for any questions concerning the study. The Ethics Review Boards of the *CHU de Québec-Université Laval* Research Center (Quebec City University Hospital) approved this study (MP-20-2020-4670).

### Survey instrument development

1  
2 The survey was based on previous work <sup>15,22,23,40-42</sup> and developed in French by a multidisciplinary team of  
3  
4 clinicians and scientists with expertise in epidemiology, social science, and medicine. This 9-item  
5  
6 questionnaire was pilot-tested with a dozen of clinicians and scientists not involved in the study. In the survey  
7  
8 introduction, PCPs were reminded that, as part of the PERSPECTIVE I&I research project, they have received  
9  
10 a letter reporting risk category for at least one of their patient's BC risk assessments. After this introduction,  
11  
12 PCPs were invited to provide their experience and satisfaction through seven close-ended multiple-choice  
13  
14 questions related to the following aspects:  
15

- 16 • Familiarity with the risk-based BC screening approach, clarity of the letter used to inform women on  
17  
18 their risk category, usefulness of the information booklet in understanding the result letter, attitudes  
19  
20 and readiness regarding the proposed screening action plan, and perceived needs for more training  
21  
22 (1 question with 8 statements);  
23  
24 • Use and appreciation of the PERSPECTIVE I&I project website, which provides further information  
25  
26 about the risk-based BC screening approach (1 question with 4 statements);  
27  
28 • Attitudes towards implementing a risk-based approach at the population level and its perceived  
29  
30 benefits (2 questions);  
31  
32 • Socio-demographic information such as profession, gender, and years of practice (3 questions).  
33  
34  
35  
36  
37

38 The survey also had two open-ended questions on possible ways to improve the risk-based BC screening  
39  
40 approach and the material provided and on additional resources that would be needed to support their  
41  
42 practice (2 questions).  
43

44 **Statistical analysis**

45  
46 Descriptive analyses were used to report participants' responses to the seven closed-ended questions. We  
47  
48 used Fisher's exact test with the SAS software, Version 9.4 (Copyright © 2016 by SAS Institute Inc., Cary, NC,  
49  
50 USA) for our bivariate analyses. Specifically, we tested whether participants' attitudes towards the risk-based  
51  
52 BC screening approach differed across years of practice and across gender. Years of practice were classified  
53  
54 as follows: less than 5 years, 5-10 years, 11-15 years, 16-20 years, 21 years and over.  
55  
56

57  
58 A content analysis was performed on the two open-ended questions. Responses were coded by AO  
59  
60 and JL to group them into larger themes using an Excel spreadsheet. ASB also independently coded the data.

AO, JL, and ASB then deliberated over their respective coding to come up with intercoder agreement to assure the reliability of the identified themes<sup>43</sup>. An inductive approach was favored for the coding and analysis of our qualitative data where codes were selected without prior theoretical framework. It should be noted that the answers to our open-ended questions were generally short. To remain faithful to the perspectives of our participants, the themes identified are also presented in general terms.

## Results

Out of the 763 PCPs contacted, 168 (22%) participated in our study. Most of them (i.e., 72.4%) had only one patient participating in the PERSPECTIVE I&I project, while 27.6% had two patients or more. Among participants, 72.6% were female and 74.4% had more than 15 years of practice (Table 1).

### Quantitative results

Only 38.1% of our participants knew about screening based on personalized BC risk assessment. Despite this, 86.9% of participants believed it is appropriate to carry out BC risk assessment prior to screening. The vast majority also found the proposed action plan appropriate (85.7%) and were ready to follow it (88.7%). Moreover, 92.3% reported understanding the information provided in the risk letter and 89.3% of them agreed that the information booklet enabled them to understand the description of their patient's risk category. Finally, the perception of participants was mixed about the need for more training; 44.1% of them agreed that they need more training while 34.5% neither agreed nor disagreed with this statement (Figure 1).

When asked about their appreciation of the study's online resource, 158 (94%) participants reported that they did not visit the website mentioned in the risk communication documents. As for the 10 (6%) participants who did visit it, all agreed that the website answered their questions, that the information was clearly presented and easily accessible, and that they would recommend the website to their colleagues if they wish to learn more about risk-stratified BC screening approach.

When we asked participants how likely they would encourage their patients to participate in a risk-based BC screening program, 87.5% of them responded that they were likely or very likely to encourage their patients to take part in such a program (Figure 2). There was no evidence from our bivariate analysis that

1 years of practice or gender influence the likelihood of encouraging patients to participate in a risk-based BC  
2 screening program (data not shown).  
3  
4

5  
6 When presented with various statements about the benefits of personalized risk-stratified approach  
7 for BC screening, 82.1% of the participants agreed that it could screen high risk women and 69% of them  
8 responded that it could both reduce unnecessary mammograms in the future and screen women of less than  
9 50 years of age. A little more than 40% agreed that it could reduce the number of false positive mammograms  
10 and that it could lead to cost savings for society. Only a third (34.5%) of our participants believed that the  
11 approach could reduce the number of BC deaths (Figure 3).  
12  
13

14  
15  
16  
17  
18  
19  
20 **Qualitative results**

21 A total of 42 participants provided an answer to the open-ended question on whether there are aspects to  
22 be modified in the risk-based BC screening approach and in the material provided. A few participants deemed  
23 the approach ineffective, superfluous, or irrelevant. The main concern was related to the potential increase  
24 in PCPs' workload. Addressing the follow-up of high-risk patients was also judged important as well as putting  
25 efforts in the promotion of the risk-based BC screening approach in the population. Concerning the  
26 documents received, some would have appreciated a shorter, simplified, version of the documents while  
27 others felt that it would be necessary to clarify which information should be provided to patients. Lastly,  
28 some participants stated that there were no aspects to change to the risk-based approach or to the risk  
29 communication documents received.  
30  
31

32 A total of 21 participants responded to the open-ended question on additional resources that would  
33 help their practice within a risk-based BC screening approach. Participants mentioned the need to develop  
34 resources for PCPs such as a mobile application. They were concerned about improving access to information  
35 about genetic and mentioned the importance of developing information tools for patients. The need for  
36 additional training and case discussions was also raised. Finally, participants suggested transferring the  
37 follow-up role to nurses.  
38  
39

40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58 **Discussion**

59  
60 **Summary of results and perspective of the literature**

According to our results, risk-based approach to guide BC screening is receiving a strong support from PCPs with a vast majority considering the approach appropriate and being likely or very likely to recommend the approach if it were to be implemented at a population level. This positive appraisal of the risk-based BC screening approach echoed the results reported in previous <sup>17 18 19 20</sup>.

However, one of the main concerns was related to the potential impact on workload. This concern is also reported in other studies and is deemed as an important barrier to implementation <sup>44 17 45</sup>. Future risk-based BC screening initiatives will need to invest on the development and implementation of an efficient operational integration of this approach <sup>46</sup>. As our participants suggested, this could involve a greater role for nurses in assessing and communicating breast cancer risk category to patients. Several implementation scenarios such as self-management by women themselves are possible for the risk-based BC screening approach and should be considered and pilot-tested <sup>40</sup>.

The need for more professional training was mentioned in both our qualitative and quantitative results. As with workload, the need for training is a recurring aspect in previous studies looking at the implementation of risk-based screening <sup>18 24</sup>. This highlights the necessity of leading concerted multi-level strategies to offer adequate training in personalized risk assessment and stratification that includes genomics and precision medicine approaches <sup>22</sup>. In the context of the PERSPECTIVE I&I project, a website was available and mentioned in the documents for PCPs wishing to have additional information about the approach. It was concerning to know that only 6% of our participants consulted the website despite admitting their need for more training. This lack of use of the website might be explained by two aspects. First, participants found the information in the documents clear and sufficient to understand the BC risk assessment and screening action plan. Second, it is consistent with PCPs expressed concerns with increased workload. In light of these results, risk-based BC screening initiatives should ensure that risk assessment letters describing the risk category be concise, clear and in an easy-to-read format. This also highlighted the importance of involving all relevant stakeholders, particularly PCPs, when designing communication tools. If implementation efforts do provide a website or online resources, the latter should not replace the information booklet provided along with the risk category letter. The website should offer complementary and detailed information for those wishing to learn more about personalized risk assessment and risk-stratified BC screening approach. As mentioned by

1  
2 PCPs, given the limited resources in healthcare systems, the most important objective is to ensure the  
3  
4 simplicity and clarity of the information provided rather than quantity.  
5

6 **Strength and Limitations**  
7

8  
9 To our knowledge, our study is among the first to report on the experience and satisfaction of PCPs receiving  
10  
11 patients' personalized BC risk assessment and proposed action plans in order to adapt screening for their  
12  
13 individual patients. This makes it possible to collect real-life PCPs feedback compared to feedback based on  
14  
15 hypothetical scenarios. The addition of qualitative open-ended questions added context to our quantitative  
16  
17 results, by allowing our participants to give suggestions and answers that our research team had simply not  
18  
19 anticipated when constructing the questionnaire, particularly regarding the fear of an increase in their  
20  
21 workload or the way in which information was presented in the letter and information leaflets. In this sense,  
22  
23 the answers to our open-ended questions are a way of enriching the information gathered in the other  
24  
25 questions by providing new answers or more in-depth explanations of the PCPs' perspective on  
26  
27 implementation of the approach<sup>47</sup>. However, further qualitative studies would be necessary to gain an even  
28  
29 more contextualized and in-depth understanding of the issue. Finally, our results are timely given that several  
30  
31 major research projects are underway to study the implementation of risk-based BC screening approaches<sup>14</sup>  
32  
33  
34  
35 48 16 49.

36  
37  
38 The main limitation of this study is related to its relatively low response rate, which resulted in a  
39  
40 limited sample size. Such a sample cannot be deemed to represent the PCPs population at large. It also  
41  
42 prevented us from conducting more complex multivariable statistical analyses. In addition, our sample is not  
43  
44 representative of PCPs in the province of Quebec. Notably we have an over-representation of PCPs  
45  
46 identifying themselves as women and of PCPs with more than 21 years' experience, potentially affecting the  
47  
48 generalizability of our findings.  
49

50  
51 Nonetheless, our sample size is within the range of previous similar work<sup>23,29,50-52</sup> and our findings  
52  
53 are poised to offer a glimpse into the experience and satisfaction of family physicians and nurse practitioners  
54  
55 upon receiving their patient's BC risk-category and action plan for BC screening. Finally, it is worth reminding  
56  
57 that the survey was launched in the middle of the third wave of the COVID-19 pandemic, during which health  
58  
59 care professionals were overwhelmed and strained by the massive influx of patients.  
60

## Implications for clinical practice and future research

Overall, our results show that PCPs are in favour of the integration of the risk-based BC screening approach when provided with real-life information about risk category and screening action plan. With the reduction of costs associated with genome sequencing and the rapid advancement of technologies<sup>53</sup>, it is becoming increasingly feasible for healthcare systems to allocate resources in calculating patients' genomic risk to include in risk assessment tools in order to offer to patients a risk stratified approach for screening tailored to their risk category. This study contributes to the growing body of scientific evidence evaluating the potential of implementing personalized risk assessment to offer a risk-based BC screening approach. Specifically, our findings highlight the importance of considering PCPs' perspectives when planning to implement this BC screening approach. In addition, future studies with a qualitative design would probably provide a unique opportunity to further explore PCPs views about the approach and put our findings into a wider context.



References

1. World Health Organization. Breast cancer. <https://www.who.int/news-room/fact-sheets/detail/breast-cancer#:~:text=In%202020%2C%20there%20were%202.3%20million%20women%20diagnosed,years%2C%20making%20it%20the%20world's%20most%20prevalent%20cancer.>

2. Brenner DR, Poirier A, Woods RR, et al. Projected estimates of cancer in Canada in 2022. *Canadian Medical Association Journal*. 2022;194(17):E601-E607. doi:10.1503/cmaj.212097

3. Canadian Cancer Society. Breast cancer statistics. <https://cancer.ca/en/cancer-information/cancer-types/breast/statistics>

4. Dibden A, Offman J, Duffy SW, Gabe R. Worldwide Review and Meta-Analysis of Cohort Studies Measuring the Effect of Mammography Screening Programmes on Incidence-Based Breast Cancer Mortality. *Cancers*. 2020;12(4):976.

5. Duffy SW, Vulkan D, Cuckle H, et al. Effect of mammographic screening from age 40 years on breast cancer mortality (UK Age trial): final results of a randomised, controlled trial. *The Lancet Oncology*. 2020;21(9):1165-1172. doi:10.1016/S1470-2045(20)30398-3

6. Maajani K, Jalali A, Alipour S, Khodadost M, Tohidinik HR, Yazdani K. The Global and Regional Survival Rate of Women With Breast Cancer: A Systematic Review and Meta-analysis. *Clin Breast Cancer*. Jun 2019;19(3):165-177. doi:10.1016/j.clbc.2019.01.006

7. Román M, Sala M, Domingo L, Posso M, Louro J, Castells X. Personalized breast cancer screening strategies: A systematic review and quality assessment. *PloS one*. 2019;14(12):e0226352. doi:10.1371/journal.pone.0226352

8. Mandrik O, Zielonke N, Meheus F, et al. Systematic reviews as a 'lens of evidence': Determinants of benefits and harms of breast cancer screening. *International Journal of Cancer*. 2019;145(4):994-1006. doi:<https://doi.org/10.1002/ijc.32211>

9. Marmot MG, Altman DG, Cameron DA, Dewar JA, Thompson SG, Wilcox M. The benefits and harms of breast cancer screening: an independent review. *Br J Cancer*. Jun 11 2013;108(11):2205-40. doi:10.1038/bjc.2013.177

10. Nelson HD, Zakher B, Cantor A, et al. Risk factors for breast cancer for women aged 40 to 49 years: a systematic review and meta-analysis. *Ann Intern Med*. May 1 2012;156(9):635-48. doi:10.7326/0003-4819-156-9-201205010-00006

11. Shieh Y, Eklund M, Madlensky L, et al. Breast Cancer Screening in the Precision Medicine Era: Risk-Based Screening in a Population-Based Trial. *JNCI: Journal of the National Cancer Institute*. 2017;109(5)doi:10.1093/jnci/djw290

12. Pashayan N, Morris S, Gilbert FJ, Pharoah PDP. Cost-effectiveness and Benefit-to-Harm Ratio of Risk-Stratified Screening for Breast Cancer: A Life-Table Model. *JAMA Oncology*. 2018;4(11):1504-1510. doi:10.1001/jamaoncol.2018.1901

13. Sankatsing VDV, van Ravesteyn NT, Heijnsdijk EAM, Broeders MJM, de Koning HJ. Risk stratification in breast cancer screening: Cost-effectiveness and harm-benefit ratios for low-risk and high-risk women. *Int J Cancer*. Dec 1 2020;147(11):3059-3067. doi:10.1002/ijc.33126
14. Pashayan N, Antoniou AC, Ivanus U, et al. Personalized early detection and prevention of breast cancer: ENVISION consensus statement. *Nat Rev Clin Oncol*. Nov 2020;17(11):687-705. doi:10.1038/s41571-020-0388-9
15. van den Broek JJ, Schechter CB, van Ravesteyn NT, et al. Personalizing Breast Cancer Screening Based on Polygenic Risk and Family History. *JNCI: Journal of the National Cancer Institute*. 2021;113(4):434-442. doi:10.1093/jnci/djaa127
16. Brooks JD, Nabi HH, Andrulis IL, et al. Personalized Risk Assessment for Prevention and Early Detection of Breast Cancer: Integration and Implementation (PERSPECTIVE I&I). *J Pers Med*. Jun 4 2021;11(6)doi:10.3390/jpm11060511
17. French DP, Woof VG, Ruane H, Evans DG, Ulph F, Donnelly LS. The feasibility of implementing risk stratification into a national breast cancer screening programme: a focus group study investigating the perspectives of healthcare personnel responsible for delivery. *BMC Womens Health*. May 2 2022;22(1):142. doi:10.1186/s12905-022-01730-0
18. Blouin-Bougie J, Amara N, Simard J. Toward a Population-Based Breast Cancer Risk Stratification Approach? The Needs and Concerns of Healthcare Providers. *J Pers Med*. Jun 10 2021;11(6)doi:10.3390/jpm11060540
19. Kaplan CP, Karliner L, Lee A, Livaudais-Toman J, Tice JA, Ozanne E. Acceptability of an mHealth breast cancer risk-reduction intervention promoting risk assessment, education, and discussion of risk in the primary care setting. *Mhealth*. 2021;7:54. doi:10.21037/mhealth-20-82
20. Lapointe J, Buron A-C, Mbuya-Bienge C, et al. Polygenic risk scores and risk-stratified breast cancer screening: Familiarity and perspectives of health care professionals. *Genetics in Medicine*. 2022/11/01/ 2022;24(11):2380-2388. doi:<https://doi.org/10.1016/j.gim.2022.08.001>
21. O'Shea R, Rankin NM, Kentwell M, et al. How can Australia integrate routine genetic sequencing in oncology: a qualitative study through an implementation science lens. *Genetics in medicine : official journal of the American College of Medical Genetics*. Sep 2020;22(9):1507-1516. doi:10.1038/s41436-020-0838-x
22. Owusu Obeng A, Fei K, Levy KD, et al. Physician-Reported Benefits and Barriers to Clinical Implementation of Genomic Medicine: A Multi-Site IGNITE-Network Survey. *J Pers Med*. Jul 24 2018;8(3)doi:10.3390/jpm8030024
23. Hann KEJ, Fraser L, Side L, et al. Health care professionals' attitudes towards population-based genetic testing and risk-stratification for ovarian cancer: a cross-sectional survey. *BMC Womens Health*. Dec 16 2017;17(1):132. doi:10.1186/s12905-017-0488-6
24. Bellhouse S, Hawkes RE, Howell SJ, Gorman L, French DP. Breast Cancer Risk Assessment and Primary Prevention Advice in Primary Care: A Systematic Review of Provider Attitudes and Routine Behaviours. *Cancers*. Aug 18 2021;13(16)doi:10.3390/cancers13164150

25. Laza-Vásquez C, Hernández-Leal MJ, Carles-Lavila M, et al. Barriers and Facilitators to the Implementation of a Personalized Breast Cancer Screening Program: Views of Spanish Health Professionals. *Int J Environ Res Public Health*. Jan 27 2022;19(3)doi:10.3390/ijerph19031406
26. Mbuya Bienge C, Pashayan N, Brooks JD, et al. Women's Views on Multifactorial Breast Cancer Risk Assessment and Risk-Stratified Screening: A Population-Based Survey from Four Provinces in Canada. *J Pers Med*. Feb 2 2021;11(2)doi:10.3390/jpm11020095
27. Chowdhury S, Henneman L, Dent T, et al. Do Health Professionals Need Additional Competencies for Stratified Cancer Prevention Based on Genetic Risk Profiling? *J Pers Med*. Jun 9 2015;5(2):191-212. doi:10.3390/jpm5020191
28. Lapointe J, Côté J-M, Mbuya-Bienge C, et al. Canadian Healthcare Professionals' Views and Attitudes toward Risk-Stratified Breast Cancer Screening. *Journal of personalized medicine*. 2023;13(7):1027. doi:10.3390/jpm13071027 Accessed 2023/06//.  
<http://europepmc.org/abstract/MED/37511640https://www.mdpi.com/2075-4426/13/7/1027/pdf?version=1687338552https://doi.org/10.3390/jpm13071027https://europepmc.org/article/PMC10381377https://europepmc.org/articles/PMC10381377?pdf=render>
29. Owusu Obeng A, Fei K, Levy KD, et al. Physician-Reported Benefits and Barriers to Clinical Implementation of Genomic Medicine: A Multi-Site IGNITE-Network Survey. *Journal of personalized medicine*. 2018;8(3):24. doi:10.3390/jpm8030024
30. Dent T, Jbilou J, Rafi I, et al. Stratified cancer screening: the practicalities of implementation. *Public Health Genomics*. 2013;16(3):94-9. doi:10.1159/000345941
31. Walker MJ, Blackmore KM, Chang A, et al. Implementing Multifactorial Risk Assessment with Polygenic Risk Scores for Personalized Breast Cancer Screening in the Population Setting: Challenges and Opportunities. *Cancers (Basel)*. May 31 2024;16(11)doi:10.3390/cancers16112116
32. Pashayan N, Antoniou AC, Lee A, et al. Should Age-Dependent Absolute Risk Thresholds Be Used for Risk Stratification in Risk-Stratified Breast Cancer Screening? *Journal of Personalized Medicine*. 2021;11(9):916.
33. Lee A, Mavaddat N, Wilcox AN, et al. BOADICEA: a comprehensive breast cancer risk prediction model incorporating genetic and nongenetic risk factors. *Genet Med*. Aug 2019;21(8):1708-1718. doi:10.1038/s41436-018-0406-9
34. Carver T, Hartley S, Lee A, et al. CanRisk Tool-A Web Interface for the Prediction of Breast and Ovarian Cancer Risk and the Likelihood of Carrying Genetic Pathogenic Variants. *Cancer Epidemiol Biomarkers Prev*. Mar 2021;30(3):469-473. doi:10.1158/1055-9965.epi-20-1319
35. Lakeman IMM, Rodríguez-Girondo M, Lee A, et al. Validation of the BOADICEA model and a 313-variant polygenic risk score for breast cancer risk prediction in a Dutch prospective cohort. *Genetics in medicine : official journal of the American College of Medical Genetics*. Nov 2020;22(11):1803-1811. doi:10.1038/s41436-020-0884-4
36. Mavaddat N, Michailidou K, Dennis J, et al. Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. *Am J Hum Genet*. Jan 3 2019;104(1):21-34. doi:10.1016/j.ajhg.2018.11.002

37. Mavaddat N, Ficorella L, Carver T, et al. Incorporating Alternative Polygenic Risk Scores into the BOADICEA Breast Cancer Risk Prediction Model. *Cancer Epidemiol Biomarkers Prev*. Mar 6 2023;32(3):422-427. doi:10.1158/1055-9965.epi-22-0756
38. Lewis CM, Vassos E. Polygenic risk scores: from research tools to clinical instruments. *Genome Medicine*. 2020/05/18 2020;12(1):44. doi:10.1186/s13073-020-00742-5
39. Mbuya-Bienge C, Pashayan N, Kazemali CD, Lapointe J, Simard J, Nabi H. A Systematic Review and Critical Assessment of Breast Cancer Risk Prediction Tools Incorporating a Polygenic Risk Score for the General Population. *Cancers (Basel)*. Nov 12 2023;15(22)doi:10.3390/cancers15225380
40. Esquivel-Sada D, Levesque E, Hagan J, Knoppers BM, Simard J. Envisioning Implementation of a Personalized Approach in Breast Cancer Screening Programs: Stakeholder Perspectives. *Healthcare Policy*. Nov 2019;15(2):39-54. doi:10.12927/hcpol.2019.26072
41. Koitsalu M, Sprangers MA, Eklund M, et al. Public interest in and acceptability of the prospect of risk-stratified screening for breast and prostate cancer. *Acta Oncol*. 2016;55(1):45-51. doi:10.3109/0284186x.2015.1043024
42. Puzhko S, Gagnon J, Simard J, Knoppers BM, Siedlikowski S, Bartlett G. Health professionals' perspectives on breast cancer risk stratification: understanding evaluation of risk versus screening for disease. *Public Health Reviews*. 2019/02/28 2019;40(1):2. doi:10.1186/s40985-019-0111-5
43. Popping R. Analyzing Open-ended Questions by Means of Text Analysis Procedures. *Bulletin de méthodologie sociologique: BMS*. 10/01 2015;128:23-39. doi:10.1177/0759106315597389
44. Laza-Vásquez C, Codern-Bové N, Cardona-Cardona À, et al. Views of health professionals on risk-based breast cancer screening and its implementation in the Spanish National Health System: A qualitative discussion group study. *PloS one*. 2022;17(2):e0263788. doi:10.1371/journal.pone.0263788
45. Rainey L, van der Waal D, Jervaeus A, et al. Are we ready for the challenge of implementing risk-based breast cancer screening and primary prevention? *Breast*. Jun 2018;39:24-32. doi:10.1016/j.breast.2018.02.029
46. Onega T, Beaber EF, Sprague BL, et al. Breast cancer screening in an era of personalized regimens: A conceptual model and National Cancer Institute initiative for risk-based and preference-based approaches at a population level. *Cancer*. 2014;120(19):2955-2964. doi:<https://doi.org/10.1002/cncr.28771>
47. Rouder J, Saucier O, Kinder R, Jans ME. What to Do With All Those Open-Ended Responses? Data Visualization Techniques for Survey Researchers. *Survey Practice*. 2021;
48. Esserman LJ. The WISDOM Study: breaking the deadlock in the breast cancer screening debate. *NPJ Breast Cancer*. 2017;3:34. doi:10.1038/s41523-017-0035-5
49. Roux A, Cholerton R, Sicsic J, et al. Study protocol comparing the ethical, psychological and socio-economic impact of personalised breast cancer screening to that of standard screening in the "My Personal Breast Screening" (MyPeBS) randomised clinical trial. *BMC Cancer*. 2022/05/06 2022;22(1):507. doi:10.1186/s12885-022-09484-6

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

50. Ayoub A, Lapointe J, Nabi H, Pashayan N. Risk-Stratified Breast Cancer Screening Incorporating a Polygenic Risk Score: A Survey of UK General Practitioners' Knowledge and Attitudes. *Genes (Basel)*. Mar 16 2023;14(3)doi:10.3390/genes14030732

51. Sabatino SA, McCarthy EP, Phillips RS, Burns RB. Breast cancer risk assessment and management in primary care: provider attitudes, practices, and barriers. *Cancer Detect Prev*. 2007;31(5):375-83. doi:10.1016/j.cdp.2007.08.003

52. Corbelli J, Borrero S, Bonnema R, et al. Use of the Gail model and breast cancer preventive therapy among three primary care specialties. *J Womens Health (Larchmt)*. Sep 2014;23(9):746-52. doi:10.1089/jwh.2014.4742

53. National Human Genome Research Institute. The Cost of Sequencing a Human Genome. 2021.

For peer review only

Enseignement Supérieur (ABES) .  
Protected by copyright, including for uses related to text and data mining, AI training, and similar technologies.

## Supplementary materials

**Supplementary file 1:** Tables and Figures

**Supplementary file 2:** Questionnaire (in French)

**Supplementary file 3:** Risk letters and proposed screening action plans based on that risk category (in French)

**Supplementary file 4:** CROSS-Checklist

## Footnotes

**Contributors:** All authors have contributed to the development of this research. HN and JS participated in the conceptualization of the research. AO, JL and PF wrote the first draft of this manuscript. All authors critically reviewed and commented on drafts of this manuscripts and approved the first version submitted. AO, AB, PF and HN revised the manuscript following the reviewers' comments, before the final version submitted.

**Funding:** This work was support by the Canadian Institutes of Health Research (#155865), the "Fondation du cancer du sein du Québec", Genome Canada (#13529), Génome Québec and the Ontario Research Foundation.

**Ethics approval and consent to participate:** Ethics approval was obtained. Ethics Research Committees of the CHU de Québec-Université Laval (MP-20-2020-4670). Participants were consented by completing the questionnaire.

**Competing interests:** The authors declare that they have no competing interests.

**Patient and public involvement:** Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

**Availability of data and materials:** Data are available upon a reasonable request to the corresponding author.

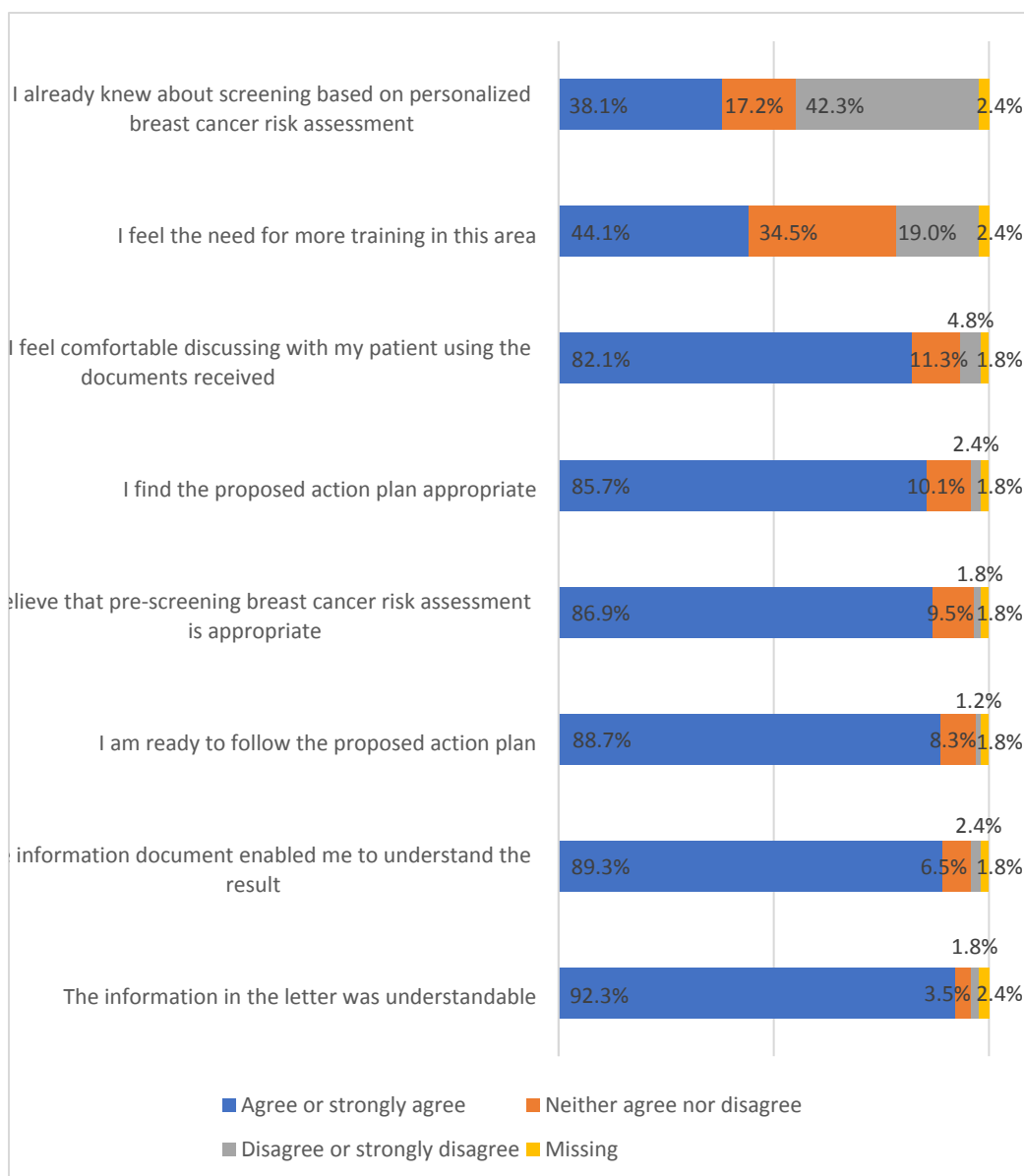
**Acknowledgements:** We would like to extend our sincere thanks to all the co-authors in our study, who generously shared their time, experiences, and insights with us. We would also like to thank the participants, without whom this study would simply not have been possible.



Supplementary file 1: Tables and Figures

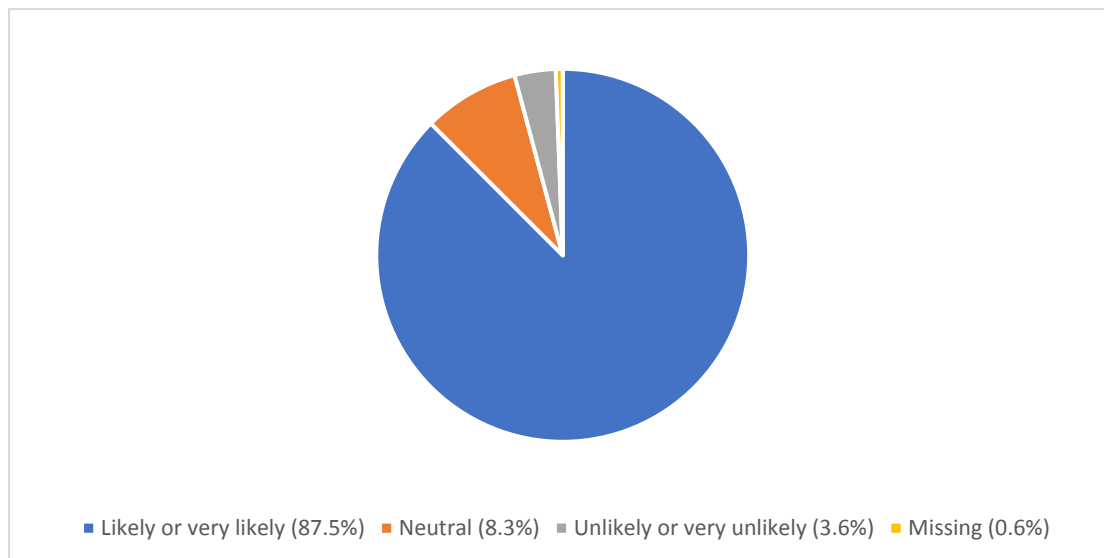
Table 1. Participants’ characteristics (n=168)

Sociodemographic variables	Frequency n (%)
<b>Profession</b>	
Physician	163 (97.0)
Primary Care Nurse Practitioner	5 (3.0)
Prefer not to answer	0 (0.0)
<b>Gender</b>	
Women	122 (72.6)
Men	46 (27.4)
Other	0 (0.0)
<b>Years of practice</b>	
< 5 years	14 (8.3)
5-10 years	20 (11.9)
11-15 years	8 (4.8)
16-20 years	20 (11.9)
> 21 years	105 (62.5)
Prefer not to answer	1 (0.6)

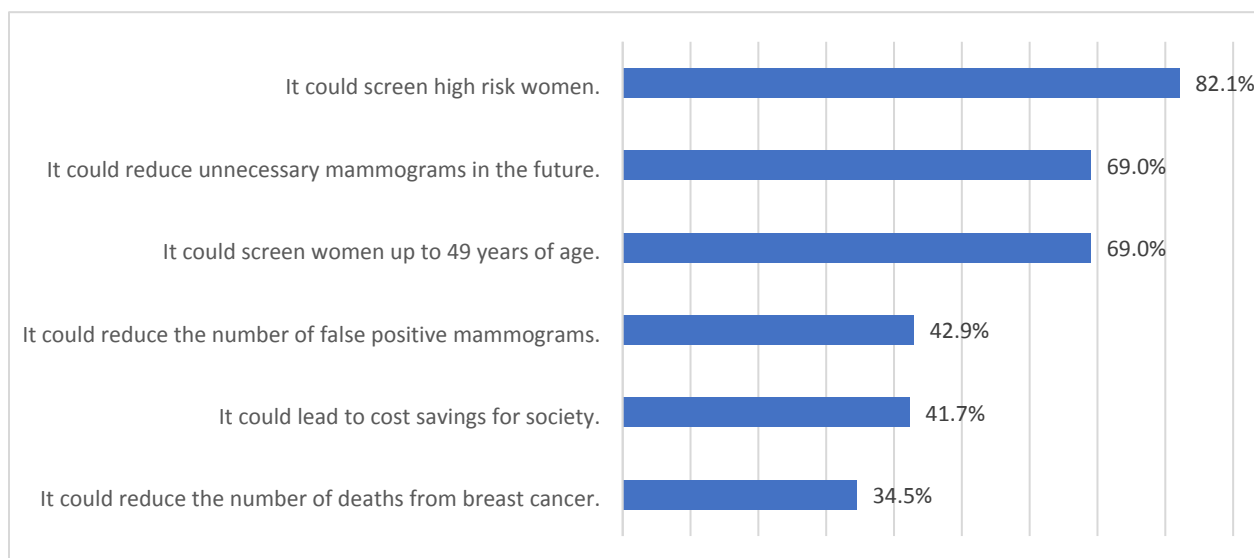


**Figure 1.** Participants' experience and satisfaction with the risk letter and the risk-based breast cancer screening approach.





**Figure 2.** Participants' likeliness to encourage patients to participate in programs that offer personalized risk assessment for breast cancer screening if it were to be offered at population level.



**Figure 3.** Benefits of personalized risk assessment for breast cancer screening according to participants.

# Questionnaire de retour d'expérience concernant l'étude PERSPECTIVE I&I

Vous avez reçu récemment un ou plusieurs résultats de l'évaluation personnalisée du risque de cancer du sein de votre patiente ou vos patientes réalisée dans le cadre du projet de recherche PERSPECTIVE. Afin de nous permettre de tenir compte de vos avis et préférences dans l'amélioration de nos processus, nous vous invitons à répondre à ce court sondage qui prendra environ 10 minutes.

Vos réponses seront rendues anonymes : nous ne collecterons pas votre nom ni aucune information permettant de vous identifier en lien avec vos réponses aux questions.

Soyez à l'aise de formuler vos remarques à l'égard de l'étude.

Vous pouvez en tout temps refuser de répondre à une question.

Si vous avez des questions sur le questionnaire, contactez-nous par téléphone au 418 682-7391 (sans frais 1 888 682-7391) ou par courriel à l'adresse [info@etudeperspective.ca](mailto:info@etudeperspective.ca).

**Q1 : Suite à votre lecture des documents reçus (lettre et document d'information), veuillez SVP indiquer votre niveau d'accord avec les énoncés suivants :**

	Très en accord	En accord	Ni en accord ni en désaccord	En désaccord	Très en désaccord
Les informations dans la lettre étaient compréhensibles	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Le document d'information m'a permis de bien comprendre le résultat	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Je connaissais déjà le dépistage basé sur l'évaluation personnalisée du risque de cancer du sein	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Je juge que l'évaluation du risque de cancer du sein avant dépistage est pertinente	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Je trouve le plan d'action proposé approprié	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Je suis prêt à suivre le plan d'action proposé	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Protected by copyright, including for uses related to text and data mining, AI training, and similar technologies. Enseignement Supérieur (ABES).

1 Je me sens à l'aise de discuter  
2 avec ma patiente à l'aide des  
3 documents reçus

☐☐☐☐☐

4 Je ressens le besoin d'avoir  
5 davantage de formation dans ce  
6 domaine  
7

☐☐☐☐☐

10 Q2 : Avez-vous consulté le site internet mentionné  
11 dans la lettre?

☐ Oui☐ Non

13 Q2a Si oui, pour quelles raisons ? (Cochez tout ce qui  
14 s'applique)

☐ J'avais des interrogations suite à la lecture des documents☐ Je voulais valider ma compréhension suite à la lecture des documents☐ Je voulais en savoir plus sur l'évaluation personnalisée du risque de cancer du sein☐ Autre

21 Veuillez préciser

\_\_\_\_\_

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28

29

30

31

32

33

34

35

36

37

38

39

40

41

42

43

44

45

46

47

48

49

50

51

52

53

54

55

56

57

58

59

60

Q2b Si oui, suite à votre consultation du site Internet, veuillez SVP indiquer votre niveau d'accord avec les énoncés suivants :

Très en accord

En accord

Ni en accord ni en désaccord

En désaccord

Très en désaccord

Le site Internet a répondu à mes interrogations

○

○

○

○

○

Les informations sur le site Internet étaient présentées clairement

○

○

○

○

○

J'ai trouvé les informations que je cherchais sur le site Internet

○

○

○

○

○

Je recommanderais le site Internet à mes collègues pour qu'ils s'informent sur l'évaluation personnalisée du risque de cancer du sein

○

○

○

○

○

Q3 : Si l'évaluation personnalisée du risque pour le dépistage du cancer du sein était proposée au niveau populationnel, quelle est la probabilité que vous encouragiez vos patientes à participer à un tel programme?

○ Très probable

○ Probable

○ Neutre

○ Improbable

○ Très improbable

○ Ne sais pas

Q3a : Si vous avez répondu > ou <, quelles sont les raisons qui expliquent votre choix?

☐ Cette approche demande trop de temps de consultation

☐ Je juge cette approche superflue par rapport au PQDCS

☐ Ce n'est pas de ma responsabilité

☐ Je me questionne sur la validité médicale de cette approche

☐ Je ne suis pas assez à l'aise avec cette approche

☐ Je vais utiliser cette approche lorsque d'autres le feront

☐ Autre

Veuillez préciser

Q4 : Cochez SVP le ou les énoncés avec lesquels vous êtes en accord concernant l'utilisation de l'évaluation du risque personnalisée pour le dépistage du cancer du sein.

☐ Cela pourrait réduire des mammographies inutiles dans le futur

☐ Cela pourrait permettre de dépister des femmes de 49 ans et moins

☐ Cela pourrait permettre de dépister des femmes ayant un risque élevé

☐ Cela pourrait permettre de réduire le nombre de mammographies faussement positives

☐ Cela pourrait permettre de diminuer le nombre de décès liés au cancer du sein

☐ Cela pourrait mener à des économies à la société

Q5 : Êtes-vous :

☐ Médecin

☐ Infirmière praticienne spécialisée en 1er ligne


☐ Préfère ne pas répondre

Protected by copyright, including for uses related to text and data mining, AI training, and similar technologies. Enseignement Supérieur (ABES).

For peer review only - <http://bmjopen.bmj.com/site/about/guidelines.xhtml>

2024-05-08 12:13

projectredcap.org



Q6 : Quel est votre sexe?

- ☐ Femme  
☐ Homme  
☐ Autre  
☐ Préfère ne pas répondre

Q7 : Depuis combien de temps pratiquez-vous votre profession?

- ☐ Moins de 5 ans  
☐ De 5 à 10 ans  
☐ De 11 à 15 ans  
☐ De 16 à 20 ans  
☐ 21 ans et plus  
☐ Préfère ne pas répondre

For peer review only

Protected by copyright, including for uses related to text and data mining, AI training, and similar technologies.  
Enseignement Supérieur (ABES).

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28

29

30

31

32

33

34

35

36

37

38

39

40

41

42

43

44

45

46

47

48

49

50

51

52

53

54

55

56

57

58

59

60

Questions ouvertes optionnelles

Q8 : Selon vous, y aurait-il des aspects à modifier dans l'approche personnalisée d'évaluation du risque et le matériel fourni? Si oui, lesquels?

Q9 : Quelles ressources supplémentaires pourraient vous aider?

## Checklist for Reporting Of Survey Studies (CROSS)

Section/topic	Item	Item description	Reported on page #
<b>Title and abstract</b>			
Title and abstract	1a	State the word “survey” along with a commonly used term in title or abstract to introduce the study’s design.	1-2-3
	1b	Provide an informative summary in the abstract, covering background, objectives, methods, findings/results, interpretation/discussion, and conclusions.	1-2-3
<b>Introduction</b>			
Background	2	Provide a background about the rationale of study, what has been previously done, and why this survey is needed.	4-5
Purpose/aim	3	Identify specific purposes, aims, goals, or objectives of the study.	5
<b>Methods</b>			
Study design	4	Specify the study design in the methods section with a commonly used term (e.g., cross-sectional or longitudinal).	5
	5a	Describe the questionnaire (e.g., number of sections, number of questions, number and names of instruments used).	6-7
Data collection methods	5b	Describe all questionnaire instruments that were used in the survey to measure particular concepts. Report target population, reported validity and reliability information, scoring/classification procedure, and reference links (if any).	6-7
	5c	Provide information on pretesting of the questionnaire, if performed (in the article or in an online supplement). Report the method of pretesting, number of times questionnaire was pre-tested, number and demographics of participants used for pretesting, and the level of similarity of demographics between pre-testing participants and sample population.	6-7
	5d	Questionnaire if possible, should be fully provided (in the article, or as appendices or as an online supplement).	Supp. Files
	6a	Describe the study population (i.e., background, locations, eligibility criteria for participant inclusion in survey, exclusion criteria).	6
Sample characteristics	6b	Describe the sampling techniques used (e.g., single stage or multistage sampling, simple random sampling, stratified sampling, cluster sampling, convenience sampling). Specify the locations of sample participants whenever clustered sampling was applied.	6
	6c	Provide information on sample size, along with details of sample size calculation.	6
	6d	Describe how representative the sample is of the study population (or target population if possible), particularly for population-based surveys.	11
Survey	7a	Provide information on modes of questionnaire administration, including the type and number of contacts, the location where the survey was conducted (e.g., outpatient	6-7



	administration	room or by use of online tools, such as SurveyMonkey).	
		7b Provide information of survey's time frame, such as periods of recruitment, exposure, and follow-up days.	6
		Provide information on the entry process:	N/A
		7c ->For non-web-based surveys, provide approaches to minimize human error in data entry.	
		->For web-based surveys, provide approaches to prevent "multiple participation" of participants.	
	Study preparation	8 Describe any preparation process before conducting the survey (e.g., interviewers' training process, advertising the survey).	6-7
	Ethical considerations	9a Provide information on ethical approval for the survey if obtained, including informed consent, institutional review board [IRB] approval, Helsinki declaration, and good clinical practice [GCP] declaration (as appropriate).	6
		9b Provide information about survey anonymity and confidentiality and describe what mechanisms were used to protect unauthorized access.	6
		10a Describe statistical methods and analytical approach. Report the statistical software that was used for data analysis.	8
		10b Report any modification of variables used in the analysis, along with reference (if available).	N/A
	Statistical analysis	10c Report details about how missing data was handled. Include rate of missing items, missing data mechanism (i.e., missing completely at random [MCAR], missing at random [MAR] or missing not at random [MNAR]) and methods used to deal with missing data (e.g., multiple imputation).	Supp. file 1
		10d State how non-response error was addressed.	N/A
		10e For longitudinal surveys, state how loss to follow-up was addressed.	N/A
		10f Indicate whether any methods such as weighting of items or propensity scores have been used to adjust for non-representativeness of the sample.	N/A
		10g Describe any sensitivity analysis conducted.	N/A
	Results		
	Respondent characteristics	11a Report numbers of individuals at each stage of the study. Consider using a flow diagram, if possible.	8
		11b Provide reasons for non-participation at each stage, if possible.	N/A
		11c Report response rate, present the definition of response rate or the formula used to calculate response rate.	8

Descriptive results	11d	Provide information to define how unique visitors are determined. Report number of unique visitors along with relevant proportions (e.g., view proportion, participation proportion, completion proportion).	8
	12	Provide characteristics of study participants, as well as information on potential confounders and assessed outcomes.	8
	13a	Give unadjusted estimates and, if applicable, confounder-adjusted estimates along with 95% confidence intervals and p-values.	8-9 and Sup. File
Main findings	13b	For multivariable analysis, provide information on the model building process, model fit statistics, and model assumptions (as appropriate).	N/A
	13c	Provide details about any sensitivity analysis performed. If there are considerable amount of missing data, report sensitivity analyses comparing the results of complete cases with that of the imputed dataset (if possible).	N/A
<b>Discussion</b>			
Limitations	14	Discuss the limitations of the study, considering sources of potential biases and imprecisions, such as non-representativeness of sample, study design, important uncontrolled confounders.	11
Interpretations	15	Give a cautious overall interpretation of results, based on potential biases and imprecisions and suggest areas for future research.	9-10-11-12
Generalizability	16	Discuss the external validity of the results.	11
<b>Other sections</b>			
Role of funding source	17	State whether any funding organization has had any roles in the survey's design, implementation, and analysis.	18
Conflict of interest	18	Declare any potential conflict of interest.	18
Acknowledgements	19	Provide names of organizations/persons that are acknowledged along with their contribution to the research.	18

# BMJ Open

## Primary care providers' experience and satisfaction with personalized breast cancer screening risk communication: A descriptive cross-sectional study

Journal:	<i>BMJ Open</i>
Manuscript ID	bmjopen-2024-093936.R1
Article Type:	Original research
Date Submitted by the Author:	03-Feb-2025
Complete List of Authors:	<p>Omeranovic, Arian; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Lapointe, Julie; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Fortier, Philippe; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Bergeron, Anne-Sophie; Université du Québec à Rimouski - Campus de Lévis, Département des sciences infirmières; Research Centre of the Chaudière-Appalaches Integrated Health and Social Services Centre</p> <p>Dorval, M; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center; Research Centre of the Chaudière-Appalaches Integrated Health and Social Services Centre</p> <p>Chiquette, Jocelyne; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Boubaker, Asma; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Eloy, Laurence; Ministère de la Santé et des Services sociaux du Québec, Programme québécois de cancérologie</p> <p>Turgeon, Annie; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Lambert-Côté, Laurence; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center</p> <p>Joly, Yann; McGill University, Center of Genomics and Policy; McGill University Faculty of Medicine, Human Genetics Department and Bioethics Unit</p> <p>Brooks, Jennifer; University of Toronto Dalla Lana School of Public Health</p> <p>Walker, Meghan J. ; University of Toronto Dalla Lana School of Public Health; Ontario Health</p> <p>Stockley, Tracy; University Health Network, Laboratory Medicine Program</p> <p>Pashayan, Nora; University of Cambridge, Department of Public Health and Primary Care</p> <p>Antoniou, Antonis; University of Cambridge, Department of Public Health and Primary Care</p> <p>Easton, Douglas; University of Cambridge, Department of Public Health and Primary Care</p> <p>Chiarelli, Anna; University of Toronto Dalla Lana School of Public Health; Ontario Health</p> <p>Knoppers, Bartha; McGill University, Center of Genomics and Policy</p> <p>Simard, Jacques; Hopital du Saint-Sacrement, Oncology Division, CHU de</p>

	Québec-Université Laval Research Center; Laval University Faculty of Medicine, Department of Molecular Medicine Nabi, Hermann; Hopital du Saint-Sacrement, Oncology Division, CHU de Québec-Université Laval Research Center; Laval University Faculty of Medicine, Department of Social and Preventive Medicine
<b>Primary Subject Heading</b>:	Oncology
Secondary Subject Heading:	Genetics and genomics, General practice / Family practice, Health services research
Keywords:	Epidemiology < ONCOLOGY, Breast tumours < ONCOLOGY, PUBLIC HEALTH, Cancer genetics < GENETICS

SCHOLARONE™  
Manuscripts



I, the Submitting Author has the right to grant and does grant on behalf of all authors of the Work (as defined in the below author licence), an exclusive licence and/or a non-exclusive licence for contributions from authors who are: i) UK Crown employees; ii) where BMJ has agreed a CC-BY licence shall apply, and/or iii) in accordance with the terms applicable for US Federal Government officers or employees acting as part of their official duties; on a worldwide, perpetual, irrevocable, royalty-free basis to BMJ Publishing Group Ltd ("BMJ") its licensees and where the relevant Journal is co-owned by BMJ to the co-owners of the Journal, to publish the Work in this journal and any other BMJ products and to exploit all rights, as set out in our [licence](#).

The Submitting Author accepts and understands that any supply made under these terms is made by BMJ to the Submitting Author unless you are acting as an employee on behalf of your employer or a postgraduate student of an affiliated institution which is paying any applicable article publishing charge ("APC") for Open Access articles. Where the Submitting Author wishes to make the Work available on an Open Access basis (and intends to pay the relevant APC), the terms of reuse of such Open Access shall be governed by a Creative Commons licence – details of these licences and which [Creative Commons](#) licence will apply to this Work are set out in our licence referred to above.

Other than as permitted in any relevant BMJ Author's Self Archiving Policies, I confirm this Work has not been accepted for publication elsewhere, is not being considered for publication elsewhere and does not duplicate material already published. I confirm all authors consent to publication of this Work and authorise the granting of this licence.

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28

29

30

31

32

33

34

35

36

37

38

39

40

41

42

43

44

45

46

47

48

49

50

51

52

53

54

55

56

57

58

59

60

Primary care providers' experience and satisfaction with personalized breast cancer screening risk communication: A descriptive cross-sectional study

Word count: Abstract= 324; document = 3204

Total number of figures and tables = 4

For peer review only

## Abstract

**Objective:** To describe primary care providers' (PCPs) experience and satisfaction with receiving risk communication documents on their patient's breast cancer (BC) risk assessment and proposed screening action plan.

**Design:** Descriptive cross-sectional study.

**Setting:** A survey was distributed to all 763 primary care providers (PCPs) linked to 1,642 women participating in the PERSPECTIVE I&I research project in Quebec, approximately one to four months after the delivery of the risk communication documents. The recruitment phase took place from July 2021 to July 2022.

**Participants:** Primary care providers.

**Main outcome measures:** Descriptive analyses were conducted to report participants' experiences and satisfaction with receiving risk communication. Responses to two open-ended questions were subjected to content analysis.

**Results:** A total of 168 PCPs answered the survey, from which 73% reported being women and 74% having more than 15 years of practice. Only 38% were familiar with the risk-based BC screening approach prior to receiving their patient risk category. A majority (86%) agreed with the screening approach and would recommend it to their patients if implemented at the population level. A majority of PCPs also reported understanding the information provided (92%) and expressed agreement with the proposed BC screening action plan (89%). Some PCPs recommended simplifying the materials, acknowledging the potential increase in workload, and emphasizing the need for careful planning of professional training efforts.

**Conclusion:** PCPs expressed positive attitudes toward a risk-based BC screening approach and were generally satisfied with the information provided. This study suggests that, if introduced in Canada in a manner similar to the PERSPECTIVE I&I project, risk-based BC screening would likely be supported by most PCPs. However, they emphasized the importance of addressing concerns such as professional training and the potential impact on workload if the approach were to be implemented at the population level. Future qualitative studies are needed to further explore the training needs of PCPs and to develop strategies for integrating this approach with the high workloads faced by primary care providers.

**Keywords:** Risk-based breast cancer screening, risk assessment, primary care providers, polygenic risk score.

1  
2  
3  
**Strengths and limitations of this study:**

- 4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60
- This is the first study to report on the experience and satisfaction of PCPs receiving patients' personalized BC risk assessment and proposed action plans in order to adapt screening for their individual patients in a real-life scenario. By focusing on real-life scenarios, the study captures genuine feedback from PCPs, as opposed to feedback based on hypothetical situations.
  - The primary limitation of this study is the potential for selection bias, as participants with a more positive attitude towards risk-based BC screening may have been more likely to participate in our survey, leading to an over-representation of individuals who are more favorable to the approach compared to those who are less supportive.
  - Our sample is also not representative of PCPs population in the province of Quebec. We have an over-representation of women family physician and an under-representation of primary care nurses' practitioners.



## Introduction

According to the World Health Organization, more than 2.3 million women were diagnosed with breast cancer (BC) in 2020, leading to more than 600,000 deaths <sup>1</sup>. In Canada, BC is the second most commonly diagnosed cancer and more than 28,600 women were diagnosed with this disease in 2022 <sup>2</sup>. Fortunately, the death rate from BC has steadily declined since its peak in 1986 <sup>3 4 5</sup>. Such a decline in mortality can be attributed to improved treatments and more efficient screening programs <sup>6 3</sup>.

While current age-based BC screening programs have been associated with a reduction in mortality for certain populations, there are still areas for improvement <sup>7</sup>. This includes reducing BC overdiagnosis <sup>8</sup>, which can have several consequences such as unnecessary medical exams, treatments, and psychological impacts on patients <sup>9</sup>. Moreover, age-based screening recommendations ignore several BC risk factors, such as genetic susceptibility, lifestyle habits, or reproductive history <sup>10</sup>. Evidence suggests that a more personalized risk-based approach could be a cost-effective way to improve BC screening programs <sup>11 12 13 14</sup>. This personalized approach involves targeting women at the highest risk for developing BC. <sup>15</sup> First, such risk stratification is expected to allow for reducing BC mortality through early detection of tumors in high-risk patients, thereby significantly increasing the chances of effective therapeutic management, cure, and long-term survival <sup>15</sup>. Moreover, by focusing screening efforts on specific populations, this personalized approach would lead to a more rational and cost-effective allocation of limited healthcare resources, representing a significant benefit in terms of cost optimization and spending efficiency within the healthcare system <sup>13 15</sup>.

Although a personalized risk-based approach appears promising, its implementation does represent a challenge <sup>14 16</sup>. Part of this challenge concerns the coordination of health services through adequate preparation of, and efficient communication with, primary care providers (PCPs) <sup>14</sup>. Several studies indicated that PCPs seem to have positive attitudes towards the implementation of risk-based assessment for BC <sup>17 18</sup> <sup>19 20</sup>. The use of genomic technologies for multifactorial risk assessment in other types of cancer also seems to be generally well received by various PCPs <sup>21 22 23</sup>. However, they tend to report a lack of training in conducting BC risk assessment <sup>24</sup>. Other barriers to implementing a risk-based screening approach were identified, such as an increased workload, a lack of financial and human resources, and a lack of coordination

1  
2 between public and private PCPs<sup>25</sup>. Primary healthcare professionals, such as nurse practitioners and family  
3  
4 physicians, are in a prime position to facilitate the implementation of BC screening approaches tailored to  
5  
6 each patient's individual risk level<sup>20</sup>. Their role is essential on several instances<sup>20 26 27 28</sup>. First, they are  
7  
8 expected to clearly explain to their patients the advantages and disadvantages of different screening  
9  
10 methods based on personal risk assessments. Second, to be able to effectively interpret and communicate  
11  
12 each patient's calculated risk level using specific prediction tools. Finally, to advise their patients on the most  
13  
14 appropriate screening and prevention strategies based on their individual risk profile. Ultimately, PCPs in this  
15  
16 context are expected to have the crucial responsibility of educating, and guiding their patients towards the  
17  
18 screening options best suited to their personal risk of BC<sup>20</sup>.

21  
22 Most previous studies collected PCPs' opinions and attitudes on hypothetical implementation  
23  
24 scenarios<sup>17 18 21 23 29</sup>. To our knowledge, no study has evaluated the experience and satisfaction of PCPs on  
25  
26 the actual receipt of a patient's personalized BC risk category in real-life practice. This feedback is essential  
27  
28 for informing future implementation efforts<sup>30</sup>.

31  
32 The aim of our study was to describe the experience and satisfaction of PCPs regarding the receipt  
33  
34 of information about their patients' personalized BC risk category and proposed screening action plan within  
35  
36 the context of real-life practice, as part of the PERSPECTIVE Integration and Implementation study<sup>16 31</sup>.

37  
38  
39  
40 **Materials and Methods**

41  
42 **Setting**

43  
44 The present descriptive cross-sectional study is part of a major Canadian research project entitled  
45  
46 PERSPECTIVE: I&I (Personalized Risk Assessment for Prevention and Early Detection of Breast Cancer:  
47  
48 Integration and Implementation), which aims to improve breast cancer risk assessment and determine  
49  
50 optimal approaches for implementing risk-based screening and prevention within the Canadian health  
51  
52 system<sup>32 16</sup>. This project included a pre-implementation research activity recruiting more than 3,750 women  
53  
54 from Quebec and Ontario, Canada's two most populous provinces<sup>31</sup>. Participating women underwent a  
55  
56 comprehensive BC risk assessment using the Breast and Ovarian Analysis of Disease Incidence and Carrier  
57  
58 Estimation Algorithm (BOADICEA) model implemented in the CanRisk prediction tool. This tool estimates  
59  
60

participants' 10-year BC risk using the polygenic risk score (PRS) and multiple risk factors, namely age at menarche, age at menopause, number of children, age at first live birth, use of oral contraception, use of hormone replacement therapy, body mass index, height and alcohol use<sup>33 34 35,36</sup>. By identifying and combining common, low-penetrance genetic variants, PRS is considered as useful tool for estimating the genetic risk of developing disease at both individual and population levels<sup>37 36 38 39</sup>. The information from the risk assessment was then used to inform patients about their risk category and possible screening action plan. In Quebec province only, the risk communication documents were sent to both the women and their designated family physician or primary care nurse practitioner. This included a risk letter that reported on women's 10-year estimated risk, stratified into three risk categories using age-dependent risk thresholds. The remaining lifetime risks (from age 30 to 80 years) for these three categories—referred to as "average," "higher than average," and "high"—are based on percentages of less than 15%, 15%–24%, and more than 25%, respectively<sup>16</sup>. It also included the proposed screening action plan based on that risk category. Finally, it also includes a 2-page information booklet on the study, risk assessment, the importance of discussing their risk level with their patient, and a follow-up decision tree detailing the proposed action plan based on risk category<sup>16 32</sup>.

### Design and participants

All family physicians and primary care nurse practitioners designated by each of the 1,642 women participating in PERSPECTIVE I&I in the province of Quebec were sent an invitation letter and the survey. They were mailed about one to four months after the letter informing women of their risk category and the corresponding screening action plan were sent out. To increase participation, two additional reminders were sent by fax one to six months after the initial mailing. The recruitment phase took place from July 2021 to July 2022. Participants were consented by completing the questionnaire. In addition, we specified the terms of confidentiality and participation in the first paragraph of the survey, while also providing a telephone number and e-mail address for any questions concerning the study. The Ethics Review Boards of the *CHU de Québec-Université Laval* Research Center (Quebec City University Hospital) approved this study (MP-20-2020-4670).

### Survey instrument development

1  
2 The survey was based on previous work <sup>15,22,23,40-42</sup> and developed in French by a multidisciplinary team of  
3  
4 clinicians and scientists with expertise in epidemiology, social science, and medicine. This 9-item  
5  
6 questionnaire was pilot-tested with twelve clinicians and scientists not involved in the study. In the survey  
7  
8 introduction, PCPs were reminded that, as part of the PERSPECTIVE I&I research project, they have received  
9  
10 a letter reporting risk category for at least one of their patient's BC risk assessments. After this introduction,  
11  
12 PCPs were invited to share their experience and satisfaction through seven close-ended multiple-choice  
13  
14 questions related to the following aspects:  
15

- 16  
17 • Familiarity with the risk-based BC screening approach, clarity of the letter used to inform women on  
18  
19 their risk category, usefulness of the information booklet in understanding the result letter, attitudes  
20  
21 and readiness regarding the proposed screening action plan, and perceived needs for more training  
22  
23 (1 question with 8 statements);  
24
- 25  
26 • Use and appreciation of the PERSPECTIVE I&I project website, which provides further information  
27  
28 about the risk-based BC screening approach (1 question with 4 statements);  
29
- 30  
31 • Attitudes towards implementing a risk-based approach at the population level and its perceived  
32  
33 benefits (2 questions);  
34
- 35  
36 • Socio-demographic information such as profession, gender, and years of practice (3 questions).  
37

38 The survey also had two open-ended questions on possible ways to improve the risk-based BC screening  
39  
40 approach and the material provided and on additional resources that would be needed to support their  
41  
42 practice (2 questions). The questionnaire is available in the Supplementary file 2.  
43

44  
45 **Statistical analysis**

46  
47 Descriptive analyses were used to report participants' responses to the seven closed-ended questions. We  
48  
49 used Fisher's exact test with the SAS software, Version 9.4 (Copyright © 2016 by SAS Institute Inc., Cary, NC,  
50  
51 USA) for our bivariate analyses. Specifically, we examined whether participants' attitudes toward the risk-  
52  
53 based BC screening approach varied based on years of practice and gender. Years of practice were classified  
54  
55 as follows: less than 5 years, 5-10 years, 11-15 years, 16-20 years, 21 years and over.  
56

57  
58 A content analysis was performed on the two open-ended questions. Responses were coded by AO  
59  
60 and JL to group them into larger themes using an Excel spreadsheet. ASB also independently coded the data.

AO, JL, and ASB then deliberated over their respective coding to come up with intercoder agreement to assure the reliability of the identified themes<sup>43</sup>. An inductive approach was favored for the coding and analysis of our qualitative data where codes were selected without prior theoretical framework. It should be noted that the answers to our open-ended questions were generally short. To remain faithful to the perspectives of our participants, the themes identified are also presented in general terms.

## Results

Out of the 763 PCPs contacted, 168 (22%) participated in our study. Most of them (i.e., 72.4%) had only one patient participating in the PERSPECTIVE I&I project, while 27.6% had two patients or more. Among participants, 72.6% were female and 74.4% had more than 15 years of practice (Table 1).

### Quantitative results

Only 38.1% of our participants knew about screening based on personalized BC risk assessment. Despite this, 86.9% of participants believed it is appropriate to carry out BC risk assessment prior to screening. The majority also found the proposed action plan appropriate (85.7%) and were ready to follow it (88.7%). Moreover, 92.3% reported understanding the information provided in the risk letter and 89.3% of them agreed that the information booklet enabled them to understand the description of their patient's risk category. Finally, the perception of participants was mixed about the need for more training; 44.1% of them agreed that they need more training while 34.5% neither agreed nor disagreed with this statement (Figure 1).

When asked about their appreciation of the study's online resource, 158 (94%) participants reported that they did not visit the website mentioned in the risk communication documents. As for the 10 (6%) participants who did visit it, all agreed that the website answered their questions, that the information was clearly presented and easily accessible, and that they would recommend the website to their colleagues if they wish to learn more about risk-stratified BC screening approach.

When we asked participants how likely they would encourage their patients to participate in a risk-based BC screening program, 87.5% of them responded that they were likely or very likely to encourage their patients to take part in such a program (Figure 2).

1  
2 When presented with various statements about the benefits of personalized risk-stratified approach  
3  
4 for BC screening, 82.1% of the participants agreed that it could screen high risk women and 69% of them  
5  
6 responded that it could both reduce unnecessary mammograms in the future and screen women of less than  
7  
8 50 years of age. A little more than 40% agreed that it could reduce the number of false positive mammograms  
9  
10 and that it could lead to cost savings for society. Only a third (34.5%) of our participants believed that the  
11  
12 approach could reduce the number of BC deaths (Figure 3).  
13  
14

15 **Qualitative results**

16  
17 A total of 42 participants provided an answer to the open-ended question on whether there are aspects to  
18  
19 be modified in the risk-based BC screening approach and in the material provided. A few participants  
20  
21 considered the approach to be ineffective, unnecessary, or irrelevant. The main concern was related to the  
22  
23 potential increase in PCPs' workload. The importance of addressing the follow-up care for high-risk patients  
24  
25 was also emphasized, along with the need to focus on promoting the risk-based BC screening approach within  
26  
27 the general population. Concerning the documents received, some would have appreciated a shorter,  
28  
29 simplified version while others felt that it would be necessary to clarify which information should be provided  
30  
31 to patients. Lastly, some participants stated that there were no aspects to change to the risk-based approach  
32  
33 or to the risk communication documents received.  
34  
35  
36  
37

38 A total of 21 participants responded to the open-ended question on additional resources that would  
39  
40 help their practice within a risk-based BC screening approach. Participants mentioned the need to develop  
41  
42 resources for PCPs such as a mobile application. They were concerned about improving access to information  
43  
44 about genetic and mentioned the importance of developing information tools for patients. The need for  
45  
46 additional training and case discussions was also raised. Finally, participants suggested transferring the  
47  
48 follow-up role to nurses.  
49  
50  
51  
52

53 **Discussion**

54 **Summary of results and perspective of the literature**

55  
56 According to our results, risk-based approach to guide BC screening is receiving a strong support from PCPs  
57  
58 with a majority considering the approach appropriate and being likely or very likely to recommend the  
59  
60

approach if it were to be implemented at a population level. This positive appraisal of the risk-based BC screening approach echoed the results reported in previous studies<sup>17 18 19 20</sup>.

However, one of the main concerns was related to the potential impact on workload. This concern is also reported in other studies and is deemed as an important barrier to implementation<sup>44 17 45</sup>. Future risk-based BC screening initiatives will need to invest in the development and implementation of an efficient operational integration of this approach<sup>46</sup>. As our participants suggested, this could involve a greater role for nurses in assessing and communicating breast cancer risk category to patients. Several implementation scenarios such as self-management by women themselves are possible for the risk-based BC screening approach and should be considered and pilot-tested<sup>40,47</sup>.

The need for more professional training was mentioned in both our qualitative and quantitative results. As with workload, the need for training is a recurring aspect in previous studies looking at the implementation of risk-based screening<sup>18,24</sup>. This indicates the necessity of leading concerted multi-level strategies to offer adequate training in personalized risk assessment and stratification that includes genomics and precision medicine approaches<sup>22</sup>. In the context of the PERSPECTIVE I&I project, a website was available and mentioned in the documents for PCPs wishing to have additional information about the approach. It was concerning to know that only 6% of our participants consulted the website despite admitting their need for more training. Two factors may explain the limited use of the website. First, participants found the information in the documents clear and sufficient to understand the BC risk assessment and screening action plan. Second, this aligns with the concerns expressed by PCPs regarding an increased workload. In light of these results, risk-based BC screening initiatives should ensure that risk assessment letters describing the risk category be concise, clear and in an easy-to-read format. This also indicated the importance of involving all relevant stakeholders, particularly PCPs, when designing communication tools. If implementation efforts include a website or online resources, these should complement, rather than replace, the information booklet provided with the risk category letter. The website should offer complementary and detailed information for those wishing to learn more about personalized risk assessment and risk-stratified BC screening approach. As mentioned by PCPs, given the limited resources in healthcare systems, the most important objective is to ensure the simplicity and clarity of the information provided rather than quantity.



1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

**Strength and Limitations**

To our knowledge, no study has examined the experience and satisfaction of PCPs in receiving personalized BC risk assessments and proposed action plans to tailor screening for their individual patients. This makes it possible to collect real-life PCPs feedback compared to feedback based on hypothetical scenarios. Furthermore, our results are timely given that several major research projects are underway to study the implementation of risk-based BC screening approaches <sup>14 48 16 49</sup>.

The main limitation of this study is that our sample may be biased toward PCPs with more positive attitudes toward risk-based BC screening. As a result, PCPs who were less interested in this approach may have been less likely to participate in the survey, leading to their underrepresentation in our sample. Thus, such a sample cannot be considered representative of the broader population PCPs population. In addition, our sample is not representative of PCPs in the province of Quebec. Notably, there is an over-representation of female family physicians and an overall under-representation of primary care nurse practitioners <sup>50</sup>, which may impact the generalizability of our findings. However, it is important to note that the survey was launched during the midst of the third wave of the COVID-19 pandemic, a time when healthcare professionals were under significant stress and facing an increased workload <sup>51 52</sup>. Another limitation is the lack of socio-demographic data for the 78% of primary care providers who did not participate in our survey, which limits the statistical analysis and generalizability of our findings.

The inclusion of qualitative open-ended questions provided valuable context to our quantitative results by allowing participants to offer insights and suggestions that our research team had not anticipated during the development of the questionnaire. Notably, this included concerns about increased workload and the way information was presented in the letter and information leaflets. In this way, the responses to the open-ended questions enriched our findings, offering new perspectives and more detailed explanations of primary care providers' views on implementing the risk-based screening approach <sup>53</sup>.

However, open-ended questions alone do not provide a comprehensive understanding of PCPs' attitudes and perspectives. Future qualitative research is needed to gather more contextualized and detailed data on their views, particularly regarding training needs and how to integrate a risk-based BC screening approach while managing high workloads.



Despite these limitations, our findings provide valuable insights into the experiences and satisfaction of family physicians and nurse practitioners when receiving their patients' BC risk categories and screening action plans.

### Implications for clinical practice and future research

Overall, our results show that PCPs are in favour of the integration of the risk-based BC screening approach when provided with real-life information about risk category and screening action plan. With the reduction of costs associated with genome sequencing and the rapid advancement of technologies<sup>54</sup>, it is becoming increasingly feasible for healthcare systems to allocate resources in calculating patients' genomic risk to include in risk assessment tools in order to offer to patients a risk stratified approach for screening tailored to their risk category. This study contributes to the growing body of scientific evidence evaluating the potential of implementing personalized risk assessment to offer a risk-based BC screening approach. Specifically, our findings indicate the importance of considering PCPs' perspectives when planning to implement this BC screening approach. In addition, future studies with a qualitative design would probably provide a unique opportunity to further explore PCPs views about the approach and put our findings into a wider context.

References

1. World Health Organization. Breast cancer. <https://www.who.int/news-room/fact-sheets/detail/breast-cancer#:~:text=In%202020%2C%20there%20were%202.3%20million%20women%20diagnosed,years%2C%20making%20it%20the%20world's%20most%20prevalent%20cancer.>

2. Brenner DR, Poirier A, Woods RR, et al. Projected estimates of cancer in Canada in 2022. *Canadian Medical Association Journal*. 2022;194(17):E601-E607. doi:10.1503/cmaj.212097

3. Canadian Cancer Society. Breast cancer statistics. <https://cancer.ca/en/cancer-information/cancer-types/breast/statistics>

4. Dibden A, Offman J, Duffy SW, Gabe R. Worldwide Review and Meta-Analysis of Cohort Studies Measuring the Effect of Mammography Screening Programmes on Incidence-Based Breast Cancer Mortality. *Cancers*. 2020;12(4):976.

5. Duffy SW, Vulkan D, Cuckle H, et al. Effect of mammographic screening from age 40 years on breast cancer mortality (UK Age trial): final results of a randomised, controlled trial. *The Lancet Oncology*. 2020;21(9):1165-1172. doi:10.1016/S1470-2045(20)30398-3

6. Maajani K, Jalali A, Alipour S, Khodadost M, Tohidinik HR, Yazdani K. The Global and Regional Survival Rate of Women With Breast Cancer: A Systematic Review and Meta-analysis. *Clin Breast Cancer*. Jun 2019;19(3):165-177. doi:10.1016/j.clbc.2019.01.006

7. Román M, Sala M, Domingo L, Posso M, Louro J, Castells X. Personalized breast cancer screening strategies: A systematic review and quality assessment. *PloS one*. 2019;14(12):e0226352. doi:10.1371/journal.pone.0226352

8. Mandrik O, Zielonke N, Meheus F, et al. Systematic reviews as a 'lens of evidence': Determinants of benefits and harms of breast cancer screening. *International Journal of Cancer*. 2019;145(4):994-1006. doi:<https://doi.org/10.1002/ijc.32211>

9. Marmot MG, Altman DG, Cameron DA, Dewar JA, Thompson SG, Wilcox M. The benefits and harms of breast cancer screening: an independent review. *Br J Cancer*. Jun 11 2013;108(11):2205-40. doi:10.1038/bjc.2013.177

10. Nelson HD, Zakher B, Cantor A, et al. Risk factors for breast cancer for women aged 40 to 49 years: a systematic review and meta-analysis. *Ann Intern Med*. May 1 2012;156(9):635-48. doi:10.7326/0003-4819-156-9-201205010-00006

11. Shieh Y, Eklund M, Madlensky L, et al. Breast Cancer Screening in the Precision Medicine Era: Risk-Based Screening in a Population-Based Trial. *JNCI: Journal of the National Cancer Institute*. 2017;109(5)doi:10.1093/jnci/djw290

12. Pashayan N, Morris S, Gilbert FJ, Pharoah PDP. Cost-effectiveness and Benefit-to-Harm Ratio of Risk-Stratified Screening for Breast Cancer: A Life-Table Model. *JAMA Oncology*. 2018;4(11):1504-1510. doi:10.1001/jamaoncol.2018.1901
13. Sankatsing VDV, van Ravesteyn NT, Heijnsdijk EAM, Broeders MJM, de Koning HJ. Risk stratification in breast cancer screening: Cost-effectiveness and harm-benefit ratios for low-risk and high-risk women. *Int J Cancer*. Dec 1 2020;147(11):3059-3067. doi:10.1002/ijc.33126
14. Pashayan N, Antoniou AC, Ivanus U, et al. Personalized early detection and prevention of breast cancer: ENVISION consensus statement. *Nat Rev Clin Oncol*. Nov 2020;17(11):687-705. doi:10.1038/s41571-020-0388-9
15. van den Broek JJ, Schechter CB, van Ravesteyn NT, et al. Personalizing Breast Cancer Screening Based on Polygenic Risk and Family History. *JNCI: Journal of the National Cancer Institute*. 2021;113(4):434-442. doi:10.1093/jnci/djaa127
16. Brooks JD, Nabi HH, Andrulis IL, et al. Personalized Risk Assessment for Prevention and Early Detection of Breast Cancer: Integration and Implementation (PERSPECTIVE I&I). *J Pers Med*. Jun 4 2021;11(6)doi:10.3390/jpm11060511
17. French DP, Woof VG, Ruane H, Evans DG, Ulph F, Donnelly LS. The feasibility of implementing risk stratification into a national breast cancer screening programme: a focus group study investigating the perspectives of healthcare personnel responsible for delivery. *BMC Womens Health*. May 2 2022;22(1):142. doi:10.1186/s12905-022-01730-0
18. Blouin-Bougie J, Amara N, Simard J. Toward a Population-Based Breast Cancer Risk Stratification Approach? The Needs and Concerns of Healthcare Providers. *J Pers Med*. Jun 10 2021;11(6)doi:10.3390/jpm11060540
19. Kaplan CP, Karliner L, Lee A, Livaudais-Toman J, Tice JA, Ozanne E. Acceptability of an mHealth breast cancer risk-reduction intervention promoting risk assessment, education, and discussion of risk in the primary care setting. *Mhealth*. 2021;7:54. doi:10.21037/mhealth-20-82
20. Lapointe J, Buron A-C, Mbuya-Bienge C, et al. Polygenic risk scores and risk-stratified breast cancer screening: Familiarity and perspectives of health care professionals. *Genetics in Medicine*. 2022/11/01/ 2022;24(11):2380-2388. doi:<https://doi.org/10.1016/j.gim.2022.08.001>
21. O'Shea R, Rankin NM, Kentwell M, et al. How can Australia integrate routine genetic sequencing in oncology: a qualitative study through an implementation science lens. *Genetics in medicine : official journal of the American College of Medical Genetics*. Sep 2020;22(9):1507-1516. doi:10.1038/s41436-020-0838-x
22. Owusu Obeng A, Fei K, Levy KD, et al. Physician-Reported Benefits and Barriers to Clinical Implementation of Genomic Medicine: A Multi-Site IGNITE-Network Survey. *J Pers Med*. Jul 24 2018;8(3)doi:10.3390/jpm8030024

23. Hann KEJ, Fraser L, Side L, et al. Health care professionals' attitudes towards population-based genetic testing and risk-stratification for ovarian cancer: a cross-sectional survey. *BMC Womens Health*. Dec 16 2017;17(1):132. doi:10.1186/s12905-017-0488-6
24. Bellhouse S, Hawkes RE, Howell SJ, Gorman L, French DP. Breast Cancer Risk Assessment and Primary Prevention Advice in Primary Care: A Systematic Review of Provider Attitudes and Routine Behaviours. *Cancers*. Aug 18 2021;13(16)doi:10.3390/cancers13164150
25. Laza-Vásquez C, Hernández-Leal MJ, Carles-Lavila M, et al. Barriers and Facilitators to the Implementation of a Personalized Breast Cancer Screening Program: Views of Spanish Health Professionals. *Int J Environ Res Public Health*. Jan 27 2022;19(3)doi:10.3390/ijerph19031406
26. Mbuya Bienge C, Pashayan N, Brooks JD, et al. Women's Views on Multifactorial Breast Cancer Risk Assessment and Risk-Stratified Screening: A Population-Based Survey from Four Provinces in Canada. *J Pers Med*. Feb 2 2021;11(2)doi:10.3390/jpm11020095
27. Chowdhury S, Henneman L, Dent T, et al. Do Health Professionals Need Additional Competencies for Stratified Cancer Prevention Based on Genetic Risk Profiling? *J Pers Med*. Jun 9 2015;5(2):191-212. doi:10.3390/jpm5020191
28. Lapointe J, Côté J-M, Mbuya-Bienge C, et al. Canadian Healthcare Professionals' Views and Attitudes toward Risk-Stratified Breast Cancer Screening. *Journal of personalized medicine*. 2023;13(7):1027. doi:10.3390/jpm13071027 Accessed 2023/06//.   
<http://europepmc.org/abstract/MED/37511640><https://www.mdpi.com/2075-4426/13/7/1027/pdf?version=1687338552><https://doi.org/10.3390/jpm13071027><https://europepmc.org/articles/PMC10381377><https://europepmc.org/articles/PMC10381377?pdf=render>
29. Owusu Obeng A, Fei K, Levy KD, et al. Physician-Reported Benefits and Barriers to Clinical Implementation of Genomic Medicine: A Multi-Site IGNITE-Network Survey. *Journal of personalized medicine*. 2018;8(3):24. doi:10.3390/jpm8030024
30. Dent T, Jbilou J, Rafi I, et al. Stratified cancer screening: the practicalities of implementation. *Public Health Genomics*. 2013;16(3):94-9. doi:10.1159/000345941
31. Walker MJ, Blackmore KM, Chang A, et al. Implementing Multifactorial Risk Assessment with Polygenic Risk Scores for Personalized Breast Cancer Screening in the Population Setting: Challenges and Opportunities. *Cancers (Basel)*. May 31 2024;16(11)doi:10.3390/cancers16112116
32. Pashayan N, Antoniou AC, Lee A, et al. Should Age-Dependent Absolute Risk Thresholds Be Used for Risk Stratification in Risk-Stratified Breast Cancer Screening? *Journal of Personalized Medicine*. 2021;11(9):916.
33. Lee A, Mavaddat N, Wilcox AN, et al. BOADICEA: a comprehensive breast cancer risk prediction model incorporating genetic and nongenetic risk factors. *Genet Med*. Aug 2019;21(8):1708-1718. doi:10.1038/s41436-018-0406-9

34. Carver T, Hartley S, Lee A, et al. CanRisk Tool-A Web Interface for the Prediction of Breast and Ovarian Cancer Risk and the Likelihood of Carrying Genetic Pathogenic Variants. *Cancer Epidemiol Biomarkers Prev*. Mar 2021;30(3):469-473. doi:10.1158/1055-9965.epi-20-1319
35. Lakeman IMM, Rodríguez-Girondo M, Lee A, et al. Validation of the BOADICEA model and a 313-variant polygenic risk score for breast cancer risk prediction in a Dutch prospective cohort. *Genetics in medicine : official journal of the American College of Medical Genetics*. Nov 2020;22(11):1803-1811. doi:10.1038/s41436-020-0884-4
36. Mavaddat N, Michailidou K, Dennis J, et al. Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. *Am J Hum Genet*. Jan 3 2019;104(1):21-34. doi:10.1016/j.ajhg.2018.11.002
37. Mavaddat N, Ficoirella L, Carver T, et al. Incorporating Alternative Polygenic Risk Scores into the BOADICEA Breast Cancer Risk Prediction Model. *Cancer Epidemiol Biomarkers Prev*. Mar 6 2023;32(3):422-427. doi:10.1158/1055-9965.epi-22-0756
38. Lewis CM, Vassos E. Polygenic risk scores: from research tools to clinical instruments. *Genome Medicine*. 2020/05/18 2020;12(1):44. doi:10.1186/s13073-020-00742-5
39. Mbuya-Bienge C, Pashayan N, Kazemali CD, Lapointe J, Simard J, Nabi H. A Systematic Review and Critical Assessment of Breast Cancer Risk Prediction Tools Incorporating a Polygenic Risk Score for the General Population. *Cancers (Basel)*. Nov 12 2023;15(22)doi:10.3390/cancers15225380
40. Esquivel-Sada D, Levesque E, Hagan J, Knoppers BM, Simard J. Envisioning Implementation of a Personalized Approach in Breast Cancer Screening Programs: Stakeholder Perspectives. *Healthcare Policy*. Nov 2019;15(2):39-54. doi:10.12927/hcpol.2019.26072
41. Koitsalu M, Sprangers MA, Eklund M, et al. Public interest in and acceptability of the prospect of risk-stratified screening for breast and prostate cancer. *Acta Oncol*. 2016;55(1):45-51. doi:10.3109/0284186x.2015.1043024
42. Puzhko S, Gagnon J, Simard J, Knoppers BM, Siedlikowski S, Bartlett G. Health professionals' perspectives on breast cancer risk stratification: understanding evaluation of risk versus screening for disease. *Public Health Reviews*. 2019/02/28 2019;40(1):2. doi:10.1186/s40985-019-0111-5
43. Popping R. Analyzing Open-ended Questions by Means of Text Analysis Procedures. *Bulletin de méthodologie sociologique: BMS*. 10/01 2015;128:23-39. doi:10.1177/0759106315597389
44. Laza-Vásquez C, Codern-Bové N, Cardona-Cardona À, et al. Views of health professionals on risk-based breast cancer screening and its implementation in the Spanish National Health System: A qualitative discussion group study. *PloS one*. 2022;17(2):e0263788. doi:10.1371/journal.pone.0263788
45. Rainey L, van der Waal D, Jervaeus A, et al. Are we ready for the challenge of implementing risk-based breast cancer screening and primary prevention? *Breast*. Jun 2018;39:24-32. doi:10.1016/j.breast.2018.02.029

1  
2 46. Onega T, Beaber EF, Sprague BL, et al. Breast cancer screening in an era of personalized regimens:  
3 A conceptual model and National Cancer Institute initiative for risk-based and preference-based  
4 approaches at a population level. *Cancer*. 2014;120(19):2955-2964. doi:<https://doi.org/10.1002/cncr.28771>  
5  
6 47. Evans I, Thornton H, Chalmers I, Glasziou P. *Testing Treatments: Better Research for Better*  
7 *Healthcare*. Pinter & Martin  
8  
9 Copyright © 2011 Imogen Evans, Hazel Thornton, Iain Chalmers and Paul Glasziou.; 2011.  
10  
11 48. Esserman LJ. The WISDOM Study: breaking the deadlock in the breast cancer screening debate. *NPJ*  
12 *Breast Cancer*. 2017;3:34. doi:10.1038/s41523-017-0035-5  
13  
14 49. Roux A, Cholerton R, Sicsic J, et al. Study protocol comparing the ethical, psychological and socio-  
15 economic impact of personalised breast cancer screening to that of standard screening in the “My Personal  
16 Breast Screening” (MyPeBS) randomised clinical trial. *BMC Cancer*. 2022/05/06 2022;22(1):507.  
17  
18 doi:10.1186/s12885-022-09484-6  
19  
20 50. Canada S. *2021 Census of Population*.  
21  
22 51. Canada S. Impacts of COVID-19 on health care workers: infection prevention and control  
23 (IHCWIPC). Statistics Canada; 2020.  
24  
25 52. Aymerich C, Pedruzo B, Pérez JL, et al. COVID-19 pandemic effects on health worker's mental  
26 health: Systematic review and meta-analysis. *Eur Psychiatry*. Jan 21 2022;65(1):e10.  
27  
28 doi:10.1192/j.eurpsy.2022.1  
29  
30 53. Rouder J, Saucier O, Kinder R, Jans ME. What to Do With All Those Open-Ended Responses? Data  
31 Visualization Techniques for Survey Researchers. *Survey Practice*. 2021;  
32  
33 54. National Human Genome Research Institute. The Cost of Sequencing a Human Genome. 2021.  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

Protected by copyright, including for uses related to text and data mining, AI training, and similar technologies.  
Enseignement Supérieur (ABES)

## Tables

**Table 1.** Participants' characteristics (n=168)

Sociodemographic variables	Frequency n (%)
<b>Profession</b>	
Physician	163 (97.0)
Primary Care Nurse Practitioner	5 (3.0)
Prefer not to answer	0 (0.0)
<b>Gender</b>	
Women	122 (72.6)
Men	46 (27.4)
Other	0 (0.0)
<b>Years of practice</b>	
< 5 years	14 (8.3)
5-10 years	20 (11.9)
11-15 years	8 (4.8)
16-20 years	20 (11.9)
> 21 years	105 (62.5)
Prefer not to answer	1 (0.6)

## Figure legends

**Figure 1.** Participants' experience and satisfaction with the risk letter and the risk-based breast cancer screening approach.

**Figure 2.** Participants' likeliness to encourage patients to participate in programs that offer personalized risk assessment for breast cancer screening if it were to be offered at population level.

**Figure 3.** Benefits of personalized risk assessment for breast cancer screening according to participants.

## Supplementary materials

**Supplementary file 1:** Questionnaire

**Supplementary file 2:** CROSS-Checklist

## Footnotes

**Contributors:** All authors have contributed to the development of this research. HN and JS participated in the conceptualization of the research. AO, JL and PF wrote the first draft of this manuscript. All authors



1 critically reviewed and commented on drafts of this manuscripts and approved the first version submitted.  
2  
3  
4 AO, AB, PF and HN revised the manuscript following the reviewers' comments, before the final version  
5  
6 submitted. HN acted as guarantor.  
7

8 **Funding:** This work was support by the Canadian Institutes of Health Research (#155865), the "Fondation  
9  
10 du cancer du sein du Québec", Genome Canada (#13529), Génome Québec and the Ontario Research  
11  
12 Foundation.  
13

14  
15 **Ethics approval and consent to participate:** Ethics approval was obtained. Ethics Research Committees of  
16  
17 the CHU de Québec-Université Laval (MP-20-2020-4670). Participants were consented by completing the  
18  
19 questionnaire.  
20

21  
22 **Competing interests:** The authors declare that they have no competing interests.  
23

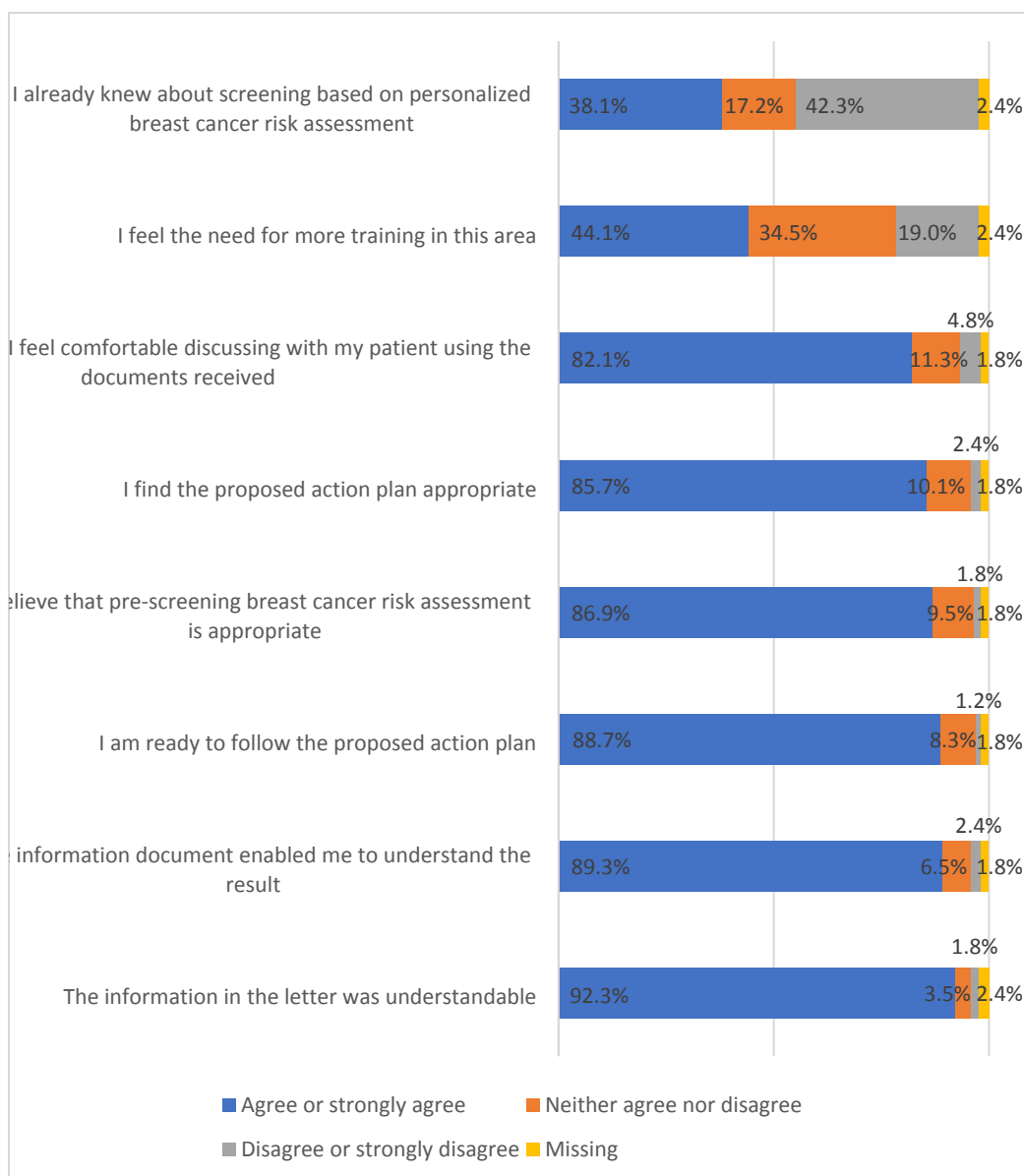
24 **Patient and public involvement:** Patients and/or the public were not involved in the design, or conduct, or  
25  
26 reporting, or dissemination plans of this research.  
27

28  
29 **Availability of data and materials:** Data are available upon a reasonable request to the corresponding  
30  
31 author.  
32

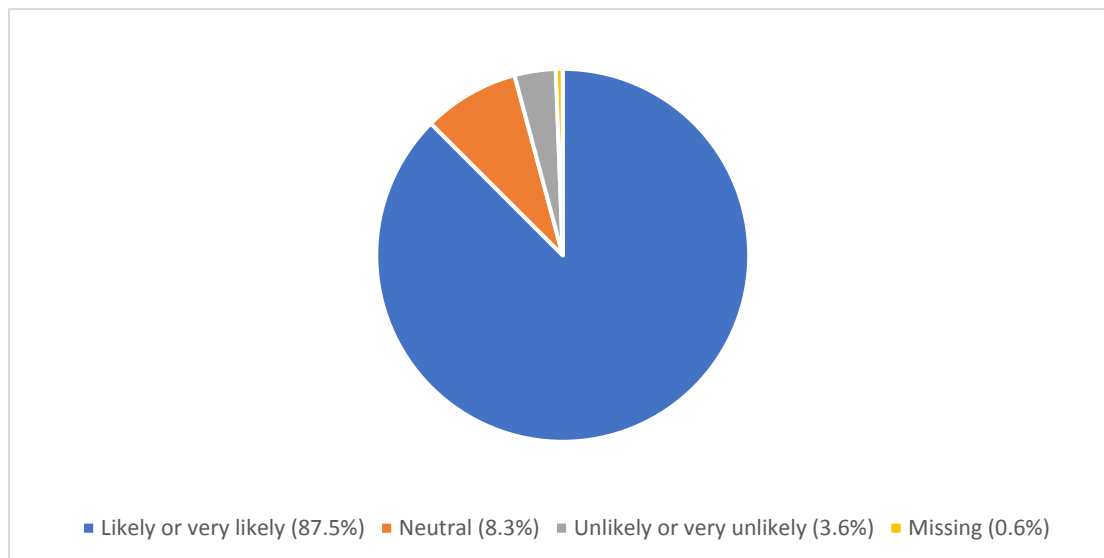
33 **Acknowledgements:** We would like to extend our sincere thanks to all the co-authors in our study, who  
34  
35 generously shared their time, experiences, and insights with us. We would also like to thank the  
36  
37 participants, without whom this study would simply not have been possible.  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

Protected by copyright, including for uses related to text and data mining, AI training, and similar technologies.  
Enseignement Supérieur (ABES)

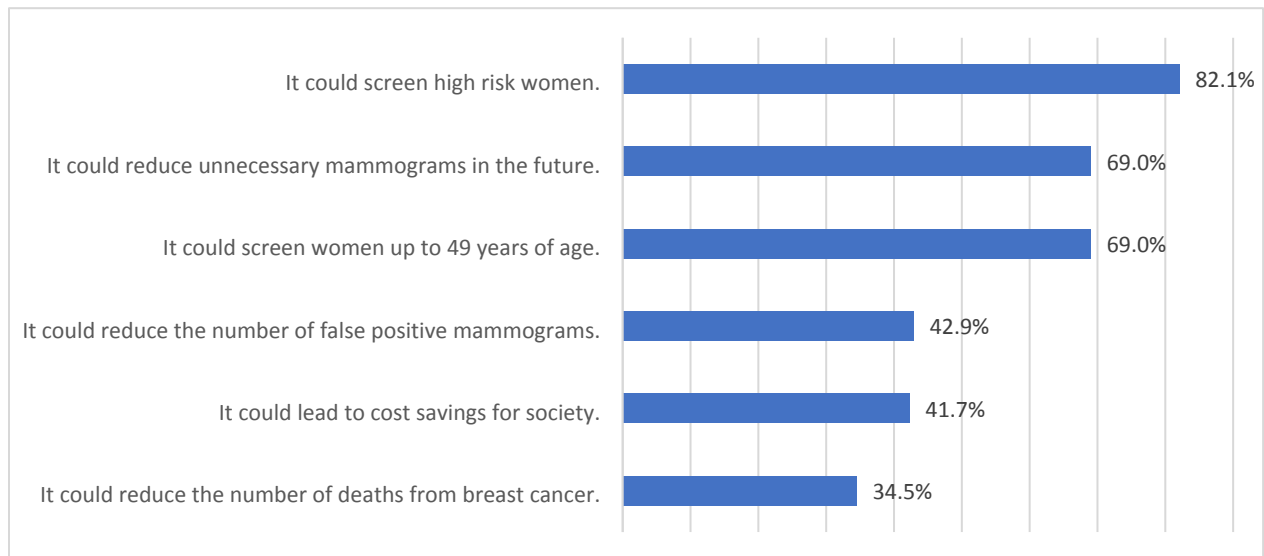




**Figure 1.** Participants' experience and satisfaction with the risk letter and the risk-based breast cancer screening approach.



**Figure 2.** Participants' likeliness to encourage patients to participate in programs that offer personalized risk assessment for breast cancer screening if it were to be offered at population level.



**Figure 3.** Benefits of personalized risk assessment for breast cancer screening according to participants.

# Experience feedback questionnaire on the PERSPECTIVE I&I study

You have recently received one or more results of the personalized breast cancer risk assessment of your patient(s), carried out as part of the PERSPECTIVE research project. To enable us to take your opinions and preferences into account in improving our processes, we invite you to complete this short survey, which will take about 10 minutes. Your answers will be anonymized: we will not collect your name or any information that could link you to your answers to the questions.

Feel free to share your comments about the study.

You may refuse to answer a question at any time.

If you have any questions about the questionnaire, please contact us by telephone at 418 682-7391 (toll-free 1 888 682-7391) or by e-mail at [info@etudeperspective.ca](mailto:info@etudeperspective.ca).

**Q1. After reading the documents received (letter and information document), please indicate your level of agreement with the following statements:**

	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly disagree
The information in the letter was understandable	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
The information document enabled me to understand the result	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
I already knew about screening based on personalized breast cancer risk assessment	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
I believe that pre-screening breast cancer risk assessment is appropriate	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
I find the proposed action plan appropriate	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
I am ready to follow the proposed action plan	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
I feel comfortable discussing with my	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

patient using the documents received					
I feel the need for more training in this area	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

**Q2. Have you consulted the website mentioned in the letter?**

- ☐ Yes
- ☐ No

**Q2a. If yes, what are the reasons? (Check all that apply)**

- ☐ I had some questions after reading the documents
- ☐ I wanted to validate my understanding after reading the documents
- ☐ I wanted to find out more about personalized breast cancer risk assessment
- ☐ Other (Please specify): \_\_\_\_\_

**Q2b. If yes, after visiting the website, please indicate your level of agreement with the following statements:**

	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly disagree
The website answered my questions	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
The information on the website was clearly presented	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
I found the information I was looking for on the website	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
I would recommend the website to my colleagues so that they can inform themselves about personalized breast cancer risk assessment.	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

**Q3. If personalized risk assessment for breast cancer screening were to be offered at population level, how likely would you be to encourage your patients to participate in such programs?**

- ☐ Very likely
- ☐ Likely

- ☐ Neutral
- ☐ Unlikely
- ☐ Very unlikely

**Q3a. If you answered 'Unlikely' or 'Very unlikely', what are the reasons for your choice? (Check all that apply)**

- ☐ This approach requires too much consultation time
- ☐ I consider this approach superfluous in relation to the PQDCS
- ☐ It is not my responsibility
- ☐ I wonder about the medical validity of this approach
- ☐ I'm not comfortable enough with this approach
- ☐ I will use this approach when others do
- ☐ Other (Please specify): \_\_\_\_\_

**Q4. Check the statement(s) with which you agree regarding the use of personalized risk assessment for breast cancer screening.**

- ☐ It could reduce unnecessary mammograms in the future
- ☐ It could screen women up to 49 years of age
- ☐ It could screen high risk women
- ☐ It could reduce number of false positive mammograms
- ☐ It could reduce the number of deaths from breast cancer
- ☐ It could lead to cost savings for society

**Q5. Are you a:**

- ☐ Physician
- ☐ Primary care nurse practitioner
- ☐ Prefer not to answer

**Q6. What is your gender?**

- ☐ Women
- ☐ Men
- ☐ Other

**Q7. For how long have you been working in your profession?**

- ☐ Less than 5 years
- ☐ From 5 to 10 years
- ☐ From 11 to 15 years
- ☐ From 16 to 20 years
- ☐ More than 21 years
- ☐ Prefer not to answer

## OPTIONAL OPEN-ENDED QUESTIONS

**Q8. In your opinion, are there any aspects of the personalized risk assessment approach and the material provided that need to be changed? If so, which ones?**

---

**Q9. What additional resources could help you?**

---

For peer review only

Checklist for Reporting Of Survey Studies (CROSS)

Section/topic	Item	Item description	Reported on page #
Title and abstract			
Title and abstract	1a	State the word “survey” along with a commonly used term in title or abstract to introduce the study’s design.	1-2-3
	1b	Provide an informative summary in the abstract, covering background, objectives, methods, findings/results, interpretation/discussion, and conclusions.	1-2-3
Introduction			
Background	2	Provide a background about the rationale of study, what has been previously done, and why this survey is needed.	4-5
Purpose/aim	3	Identify specific purposes, aims, goals, or objectives of the study.	5
Methods			
Study design	4	Specify the study design in the methods section with a commonly used term (e.g., cross-sectional or longitudinal).	5
	5a	Describe the questionnaire (e.g., number of sections, number of questions, number and names of instruments used).	6-7
Data collection methods	5b	Describe all questionnaire instruments that were used in the survey to measure particular concepts. Report target population, reported validity and reliability information, scoring/classification procedure, and reference links (if any).	6-7
	5c	Provide information on pretesting of the questionnaire, if performed (in the article or in an online supplement). Report the method of pretesting, number of times questionnaire was pre-tested, number and demographics of participants used for pretesting, and the level of similarity of demographics between pre-testing participants and sample population.	6-7
	5d	Questionnaire if possible, should be fully provided (in the article, or as appendices or as an online supplement).	Supp. Files 1
	6a	Describe the study population (i.e., background, locations, eligibility criteria for participant inclusion in survey, exclusion criteria).	6
Sample characteristics	6b	Describe the sampling techniques used (e.g., single stage or multistage sampling, simple random sampling, stratified sampling, cluster sampling, convenience sampling). Specify the locations of sample participants whenever clustered sampling was applied.	6
	6c	Provide information on sample size, along with details of sample size calculation.	6
	6d	Describe how representative the sample is of the study population (or target population if possible), particularly for population-based surveys.	11
Survey	7a	Provide information on modes of questionnaire administration, including the type and number of contacts, the location where the survey was conducted (e.g., outpatient	6-7



administration		room or by use of online tools, such as SurveyMonkey).	
	7b	Provide information of survey's time frame, such as periods of recruitment, exposure, and follow-up days.	6
		Provide information on the entry process:	N/A
	7c	→For non-web-based surveys, provide approaches to minimize human error in data entry.	
		→For web-based surveys, provide approaches to prevent "multiple participation" of participants.	
Study preparation	8	Describe any preparation process before conducting the survey (e.g., interviewers' training process, advertising the survey).	6-7
Ethical considerations	9a	Provide information on ethical approval for the survey if obtained, including informed consent, institutional review board [IRB] approval, Helsinki declaration, and good clinical practice [GCP] declaration (as appropriate).	6
	9b	Provide information about survey anonymity and confidentiality and describe what mechanisms were used to protect unauthorized access.	6
	10a	Describe statistical methods and analytical approach. Report the statistical software that was used for data analysis.	7-8
	10b	Report any modification of variables used in the analysis, along with reference (if available).	N/A
Statistical analysis	10c	Report details about how missing data was handled. Include rate of missing items, missing data mechanism (i.e., missing completely at random [MCAR], missing at random [MAR] or missing not at random [MNAR]) and methods used to deal with missing data (e.g., multiple imputation).	18-Figure
	10d	State how non-response error was addressed.	11
	10e	For longitudinal surveys, state how loss to follow-up was addressed.	N/A
	10f	Indicate whether any methods such as weighting of items or propensity scores have been used to adjust for non-representativeness of the sample.	N/A
	10g	Describe any sensitivity analysis conducted.	N/A
<b>Results</b>			
Respondent characteristics	11a	Report numbers of individuals at each stage of the study. Consider using a flow diagram, if possible.	8
	11b	Provide reasons for non-participation at each stage, if possible.	N/A
	11c	Report response rate, present the definition of response rate or the formula used to calculate response rate.	8

1				
2				
3				
4			Provide information to define how unique visitors are determined. Report number of	8
5		11d	unique visitors along with relevant proportions (e.g., view proportion, participation	
6			proportion, completion proportion).	
7	Descriptive			
8				
9	results	12	Provide characteristics of study participants, as well as information on potential	8-18
10			confounders and assessed outcomes.	
11				
12		13a	Give unadjusted estimates and, if applicable, confounder-adjusted estimates along	8-9-18
13			with 95% confidence intervals and p-values.	
14				
15	Main findings	13b	For multivariable analysis, provide information on the model building process, model	N/A
16			fit statistics, and model assumptions (as appropriate).	
17				
18			Provide details about any sensitivity analysis performed. If there are considerable	N/A
19		13c	amount of missing data, report sensitivity analyses comparing the results of complete	
20			cases with that of the imputed dataset (if possible).	
21	Discussion			
22				
23			Discuss the limitations of the study, considering sources of potential biases and	11
24	Limitations	14	imprecisions, such as non-representativeness of sample, study design, important	
25			uncontrolled confounders.	
26				
27				
28	Interpretations	15	Give a cautious overall interpretation of results, based on potential biases and	9-10-11-
29			imprecisions and suggest areas for future research.	12
30	Generalizability	16	Discuss the external validity of the results.	11
31				
32	Other sections			
33				
34	Role of funding source	17	State whether any funding organization has had any roles in the survey's design,	19
35			implementation, and analysis.	
36				
37	Conflict of interest	18	Declare any potential conflict of interest.	19
38				
39	Acknowledgements	19	Provide names of organizations/persons that are acknowledged along with their	19
40			contribution to the research.	
41				
42				
43				
44				
45				
46				
47				
48				
49				
50				
51				
52				
53				
54				
55				
56				
57				
58				
59				
60				