

# Better Together

Co-Designing Supportive Care Services with Patients and Families

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January 2026

Health  
Design  
Lab.

EMILY CARR  
UNIVERSITY  
of Art + Design

BC  
CAN  
CER

Provincial Health Services Authority

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# Land Acknowledgement

BC Cancer respectfully acknowledges that our work occurs on the traditional, ancestral, and unceded territories of First Nations across British Columbia.

This work has been done in collaboration with Emily Carr University of Art and Design, located on the unceded territories of the xwməθkwəyəm (Musqueam), Skwxwú7mesh Úxwumixw (Squamish), and səliilwətaʔt (Tseil-Waututh) Nations.

We practice these land acknowledgments not only as acts of recognition, but also as a reminder to attune ourselves to the real, and lived impacts of colonial systemic forces, as we work towards engaging in processes that look to better support people who have lived experience with cancer.





# Overview

This report presents insights gathered in collaboration with BC Cancer patients and families about the current landscape of Supportive Care Services across the province, and their vision for improved awareness and access to services. The project was a partnership between *BC Cancer's Patient and Family Partnerships team*, *BC Cancer's Supportive Care Program*, and the *Health Design Lab at Emily Carr University of Art + Design*. It was supported by 2 Patient Advisors and 39 patient and family partners who played a key role in both phases of the project. We are deeply grateful to these individuals with lived experience of cancer treatment, either as patients or as loved ones of patients, who directly contributed to this work.

Despite the critical role Supportive Care Services play in quality of life during cancer treatment and recovery, many patients and families experience barriers when accessing them. Using participatory, co-design research methods, our aim was to identify actionable opportunities to increase awareness of and improve the accessibility and quality of Supportive Care Services. This work occurred from November 2024 to January 2026, helping to shape the *BC Cancer Supportive Care 5-year Strategic Plan* and prioritize actions within its 3-year roadmap.

***“There is a big  
difference between  
treatment and care.”***

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*Listening to the voices of people with lived experience is critical to improving their care experiences.* All quotes included in this document are from people with lived experience of BC Cancer services as patients, family members or loved ones, who participated in the workshops described on the following pages

# Background & Rationale

BC Cancer delivers evidence-based, world-class cancer treatment to people living across British Columbia, while also recognizing that quality health care extends beyond treatment alone. An approach that values the lived experiences of patients and families is central to this work. Through ongoing engagement, patients and caregivers have consistently shared that many supportive care services, such as counselling, nutrition support, and financial navigation, are often not well known or easy to access. Some people did not realize these services existed, while others struggled to navigate how to access them, which contributes to uneven experience and unmet needs across the province. This feedback aligns with the priorities in B.C.'s 10 Year Cancer Action Plan, which emphasizes improving timely access to equitable, patient centered care across the cancer journey and strengthening supportive and survivorship services as essential parts of high-quality cancer care.

In response, BC Cancer Patient and Family Experience, and Supportive Care teams partnered on this project to better understand patient and family experiences and to identify opportunities to improve awareness and access to supportive care across all BC Cancer regional centers. The goals of this initiative were twofold: to raise awareness of existing supportive care services and to identify practical improvement ideas that make these services more accessible, navigable, and responsive to diverse patient and family needs. Central to this effort was moving beyond potential health care team biases, and internal assumptions, and learning directly from people with lived experience to inform service improvement and to ensure alignment with the provincial cancer care priorities.

To guide this work, we grounded our approach in a few key questions from the patient and family perspective that became the starting point for this project. We wanted to better understand:

**What supportive care services people had actually accessed and how they found them?**

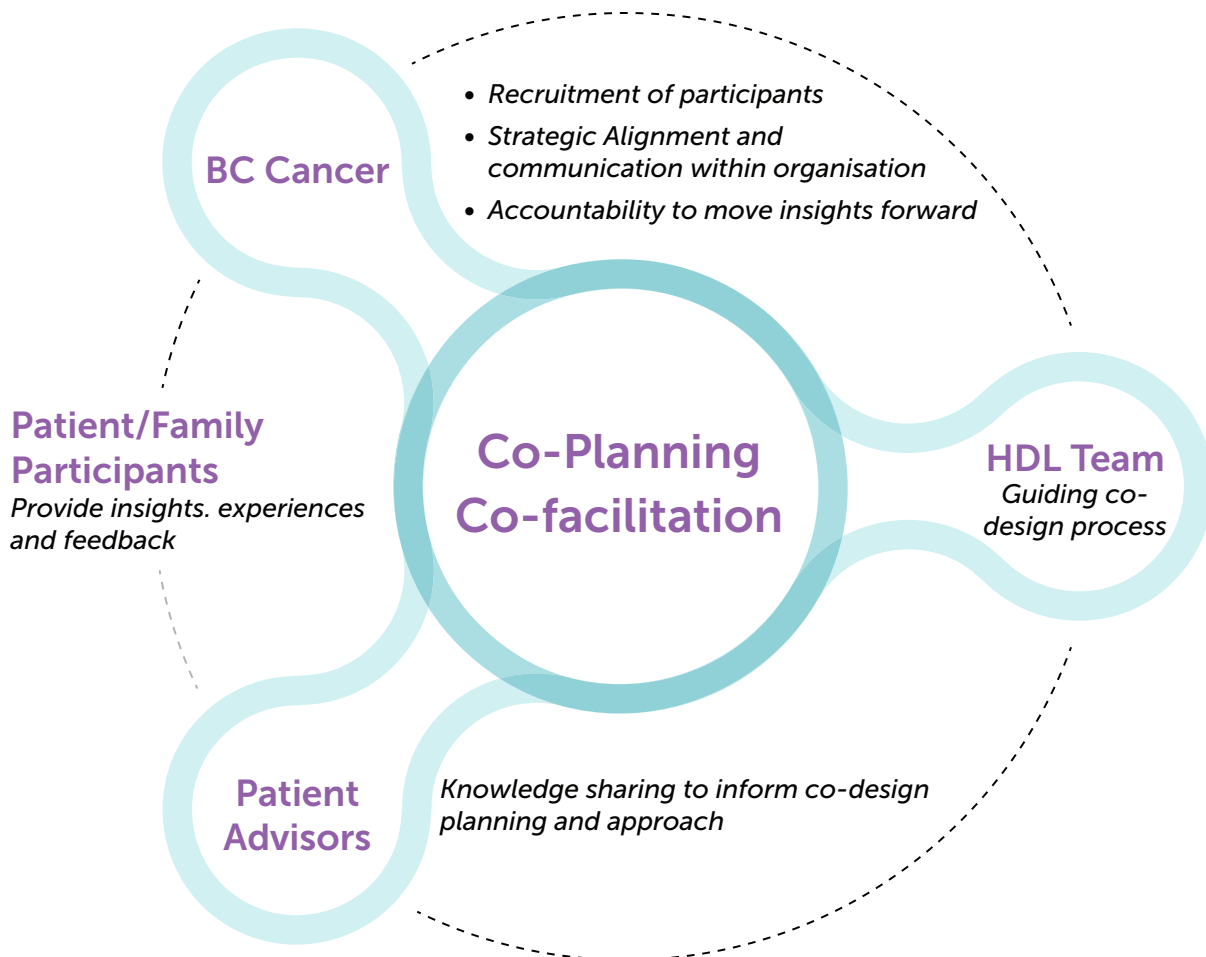
**What helped or hindered their awareness of these services?**

**And most importantly, what ideas they had for improving access moving forward?**

# A Co-Design Approach

We chose a co-design approach because we wanted patients and families to be leaders in shaping solutions, not just participants in a process. That meant creating intentional space for them to influence future outcomes. In this case, how supportive care services are accessed and experienced across BC Cancer.

To support this, we partnered with the Health Design Lab at Emily Carr University of Art & Design, whose expertise in participatory design helped us use creative and collaborative methods to surface insights, build shared understanding, and explore solutions together. We also embedded lived experience directly into the project team by recruiting two Patient Advisors, as paid members of the team ensuring the patient perspective informed every stage. The team formed in November 2024 and worked together until January 2026.



## What are Supportive Care Services?

People with cancer can face significant challenges related to both symptoms of the disease and the adverse effects of treatment. This might occur before beginning treatment, during surgery or system/radiation therapy, after treatment ends or throughout long-term treatment. Challenges can be physical, informational, emotional, psychological, social, spiritual, and practical. Supportive care services are any program or service that helps improve quality of life throughout the cancer care experience for patients and family. For example, supportive care services can include support with symptom and pain management, therapy and counselling, and nutrition support. Though focused on quality of life, supportive care generally also improves people's overall experience of care and often helps them get the best treatment possible, leading to better outcomes.



PRE-DIAGNOSIS

DIAGNOSIS

SEXUAL HEALTH  
OR FERTILITY

MENTAL & EMOTIONAL  
SUPPORT

***“This is a full-time job, having cancer, I just want someone to tell me what is reasonable to expect. I expect my healthcare team to put me in touch with the people I need. Someone bringing information to me so that I can make a decision.”***

# Project Overview

The BC Cancer project was conducted in two phases, using a mix of virtual and in-person formats.

**Phase 1** began with a province-wide survey in *December 2024*, which initiated engagement with patients and families. This was followed by two 90-minute virtual workshops and one three-hour in-person workshop in *April 2025*. This phase concluded with a validation session in *June 2025* where participants reviewed workshop findings.

**Phase 2** focused on turning ideas from Phase 1 into actionable initiatives. Building on these insights, and in alignment with the BC Cancer Strategic Planning Initiative, two virtual workshops were held in *October and November 2025*. As compared to Phase 1's engagement, which focused exclusively on patients and families, Phase 2 expanded participation to include BC Cancer staff.



# Phase 1: Our Approach

We began patient and family engagement with a province-wide survey aimed at patients, caregivers, and family members across BC Cancer. The goal was to broadly understand people’s awareness and experiences with supportive care services. The survey also functioned as a recruitment pathway for future engagements.

Following the survey, we hosted two workshops: one virtual and one in-person, with 20 participants invited to each. The aim was to understand the survey findings in greater depth, using participatory methods. We discussed their experiences accessing supportive care services and co-generated ideas for improving them. Participants from across all BC Cancer centres attended. Each session was facilitated by the project team which included Patient Advisors, BC Cancer staff and the Health Design Lab team. Both the virtual and in-person workshops were structured in two parts:

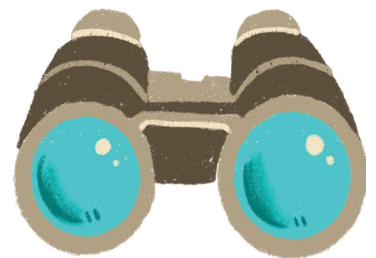
## Part 1: Mapping Current Experiences

Participants were invited to self-reflect on their experiences within the BC Cancer system by creating a map of their cancer care journey. They were asked to use supportive care service cards to show which services they accessed or wished they had accessed, and when these needs came up during their journey. These maps were then shared in smaller groups giving participants an opportunity to discuss their experiences.



## Part 2: Imagining Future Possibilities

Shifting the focus to a more future-oriented lens, participants were presented with illustrated characters and asked to choose a “sidekick” that they wish could have supported them in their cancer journey. This activity served as a warm-up icebreaker for further ideation.



Next, the group reviewed key themes that had emerged from the previously completed survey to ensure the ideas generated were grounded in the experiences of survey respondents, bringing those 500 voices into the room.

Participants were then invited to imagine an ideal system and think about what it would look like, sound like, and who would need to be involved to achieve this vision. These conversations led to rich discussions about how supportive care services could be improved to better meet people's needs.

Following the workshop, the project team synthesized the insights from these workshops to identify key themes related to current experiences and future opportunities.

Both workshops were designed with safety, diversity, accessibility and inclusivity in mind, ensuring meaningful participation for people who may face challenges

with technology or virtual formats, as well as remote location limitations. We aimed to create a meaningful, reciprocal and trauma-informed environment for everyone involved.

*“The service I wish I could have received, probably from the beginning, was the emotion support for me and for my family. We didn't know who to talk to for any information. I started to look around on the Canadian sites and found the information was overwhelming, because individual services have their own website.”*





## Phase 2: Our Approach

Phase 2 focused on transforming the ideas generated in Phase 1 into actionable and clearly prioritized solutions for BC Cancer through a second round of co-design workshops. In this phase, our goal was to gather BC Cancer Staff with patients, and families, to enable meaningful conversation about the essential role that staff play in the execution and improvement of these services. All 39 previous workshop participants were invited to attend two workshops, along with 13 staff from BC Cancer regional centers.

At this stage, patient and family ideas from Phase 1 were synthesized with initiatives in BC Cancer's Supportive Care Services Strategic Plan to create a comprehensive list of recommendations. Before the workshops began, the project team assigned each idea an initial feasibility rating, based on estimated systemic limitations and timelines. Patient Advisors were integral to this process and ensured that patient priorities remained centered throughout. To prepare for the workshops, all participants received a pre-workshop survey asking them to select the four ideas that they felt were the most impactful, and worthy of discussion. This allowed us to layer perceived impact onto the feasibility assessment. Based on this, we selected 2–3 related ideas for each breakout room to explore further during the two 90-minute virtual workshops.

**Workshop 1** was held in October and focused on *Goal 1: Improve Communication and Navigation Experience for Supportive Care Services* and *Goal 2: Provide Opportunities for Mental and Emotional Support as Part of Cancer Treatment*.

**Workshop 2** was held in November and focused on *Goal 3: Increase Availability, Access, and Quality of Supportive Care Services Across the Province*, as well as going deeper into Goal 2.

In each workshop, participants were divided into smaller breakout rooms, with each room tackling and a different set of ideas for further discussion. These discussions were centered around the following questions:

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Do these top ideas resonate with you?

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How would this look in action? What resources might we need? What barriers might we face?

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Are there other ways to meet the needs behind the ideas? How might we overcome challenges to feasibility?



## WORKSHOP 1

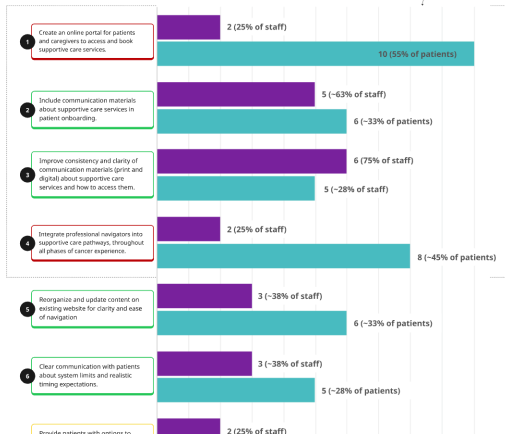
GOAL 1

### Goal #1: Improve Communication and Navigation Experience for Supportive Care Services

High Feasibility  
Medium Feasibility  
Low Feasibility

Staff: 8  
Patients + Families: 16

Discussion ideas for Friday



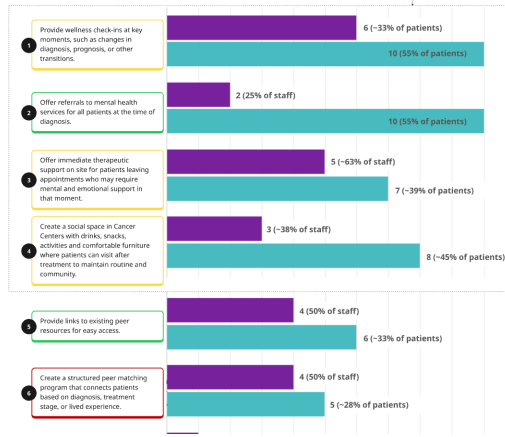
GOAL 2

### Goal #2: Provide Opportunities for Mental and Emotional Support as Part of Cancer Treatment

High Feasibility  
Medium Feasibility  
Low Feasibility

Staff: 8  
Patients + Families: 16

Discussion ideas for Friday



miro BCC Workshop Breakout Rooms (11/07/2025)

### How might we integrate mental and emotional support into the patient visit experience?

- support needs to be built into every visit
- the importance of someone coming around and asking "how are you?"
- Having somebody come around and talk in the chemo room to check-in about mental and emotional support
- need continuous check-ins- not just a one-time thing
- the importance of timely/initial dialogue with family partners/family should be eligible to get the support they need
- It's a longitudinal journey, but the system itself is episodic
- strategy for early connection to supportive care
- support over time
- BCC staff needs to be aware of what else is available outside BCC so that people can access support over time
- have someone debrief after appointments or radiation treatment to support the stress related to what has just happened- could be a nurse with medical knowledge
- different people have different needs and different needs over time
- phone call the day before asking what would be helpful
- offer of appointment times beforehand
- having a space that you can go to beforehand where you can talk about what to expect
- doing counselling afterwards would not have been helpful because of the drugs during appointments- also having to move on to next appointment
- referrals to Inspire Health
- phone calls from counsellors
- living 5 hours away, relying on phone calls for doctor's visits, setting up appointments
- BC Cancer lodges- continuous counselling through peers

S. Srividya

# Phase 1: Insights



Three clear themes about people's experiences with supportive care services emerged in Phase 1. Overall, we heard that people faced challenges with:

**Awareness of Services:** How people learn about services and how to access them.

**Mental and Emotional Wellbeing:** The mental & emotional strain of navigating cancer treatment.

**Service Offerings:** The relevance and accessibility of existing supportive care services.

***“What I’ve discovered since I dealt with all of this is that it’s not consistent for everyone. That’s an issue, that everyone gets different information.”***

## CURRENT EXPERIENCES WE HEARD

### Awareness of Services

#### **1. Insufficient Access to Informational**

**Resources:** Patients who are typically able to navigate services still struggle to find clear, consistent information about what supportive care services are available and how to access them. Many feel overwhelmed by the volume of materials they receive at diagnosis, and the resources shared vary widely from patient to patient.

#### **2. Mistimed Delivery of Information**

**About Supportive Care:** Patients emphasized that when supportive care information is shared, is just as important as what is shared. They need this information early and repeated at diagnosis, during treatment, after treatment, and especially at key transition points, including transition into survivorship or palliative care outside BC Cancer.

**3. Dependence on Healthcare Team:** We heard that without clear and easy-to-find information, patients and family rely heavily on their healthcare team to learn about supportive care services, making it crucial that this information is shared in a timely and helpful way.

#### **4. Lack of Individualized Navigation:**

Patients and families face many barriers to navigating supportive care services, including the physical and emotional toll of cancer, complex life circumstances, and limited support networks. This leaves patients who need a higher level of support in the stressful position of coordinating their own care.

*“I knew the services existed somewhere, but I didn’t know how much was available. Can I even access the services? Is it okay to reach out? I was so desperate for anything, and I don’t know if anyone cares.”*

#### **5. Limited Access to Peer Support:**

Patients and families often express the value of connecting with peers who have shared similar experiences. Peer support offers both emotional comfort and practical guidance that can help people navigate the healthcare system more effectively.



***“I think we also must consider that some of us will never be off treatment. There will never be an after for me, and some services I have been told I only have a certain number of sessions. They assume there is an after, but there is no after for me.”***

## Mental & Emotional Wellbeing



**1. Mental and Emotional Strain of Navigating Cancer Care:** We heard that the experience of cancer treatment can take a heavy toll on a person's mental and emotional wellbeing, especially at moments of transition, like diagnosis or prognosis changes. When timely, meaningful mental health support isn't available, it becomes harder for patients to cope with treatment and access the services they need. Support for caregivers is also lacking. It is essential to recognize and address these unique needs as well.

*"I lost 15 pounds in about 2 weeks because the diagnosis was so devastating to me and I just couldn't eat... That was the worst time for me in the whole treatment period. For me personally, I would have liked to be able to access mental and emotional support right at the start of diagnosis."*

## Service Offerings & Accessibility

### **1. Insufficient and Generic Service**

**Offerings:** We heard that many supportive care services people need aren't available at BC Cancer, and existing services don't always reflect the diverse needs of patients and families.

### **2. Regional Disparities in Services:**

We heard that a patient's location in BC significantly shapes their access to supportive care. Those living far from a Cancer Centre face major financial and logistical barriers to attending in-person services. Those living near a center may also have limited options, since many programs are offered only at specific sites and aren't available virtually across the province.

### **3. Limited Access to Support Services:**

We've heard that limited access to supportive care services due to short-term availability, long waitlists, and unclear scheduling, has a major impact on patients and families.

*"I tried to access in person counselling services but was told that it would have to be via zoom or phone. I am hearing impaired and require in person (face to face) for effective communication."*

*"There is too much inconsistency between regions and services. It feels like it's up to luck."*

*"The wait to get help through support services was a couple of months. I had forgotten I had requested help!"*

# We heard that patients and families want:

## **Awareness of Services:**

1. Accurate and Accessible Information Resources
2. Healthcare Team Supported Navigation
3. Peer-Supported Navigation
4. 1:1 Navigation Support

## **Mental and Emotional Wellness:**

1. Increase Opportunities for Mental and Emotional Support as Part of Cancer Treatment

## **Service Offerings:**

1. Evaluation of Regional Programs to Improve Equitable Access
2. Expand and Improve Service Quality and Offerings
3. Coordination with External Supportive Care Services



Once the workshop series concluded, the project team synthesized all the notes captured in the sessions. These notes were sorted to create the themes and statements for this report. The statements derived from the synthesis, have been laid out in this report linearly for ease of reading. This linear nature, however, does not reflect the way the feedback was shared in the workshops or the interconnectedness of the themes and insights.

***“You are empowering the patient  
by providing all the information  
available.”***

# Phase 2: Insights



In Phase 2, patients and families engaged in conversations to further expand their ideas for improvement. This included a survey to prioritize all the ideas gathered from Phase 1, and two workshops that enabled patients, families and Supportive Care Services staff to further discuss ideas. The following is a summary of the key ideas that emerged.



# Awareness of Services

## 1. Information Resources:

- Create clear information resources about supportive care services.
- Reorganize and update content on the existing website for clarity and ease of navigation.
- Design consistent print materials for supportive care services including posters, pamphlets, booklets, etc.
- Improve the use of the existing e-bulletin as a dissemination platform for ongoing communication of timely events and opportunities.
- Increase supportive care services' social media presence to advertise services.
- Map pathways for how patients receive cancer treatment in BC and are referred to BC Cancer to ensure people are connected to necessary information about supportive care services.

## 2. Navigation Support

- Create a pathway for care teams to introduce supportive care options.
- Integrate professional navigators, like a nurse or social worker.
- Set up a peer navigator program at orientation.
- Support patients who want to set up their own peer navigation and support groups when there isn't a formal program provided.

*“Every month there could be a newsletter with clubs and groups. Maybe I don’t have any pain right now, but maybe 6 months from now I might have pain. Repetition might be helpful. [...] Everyone is opted in automatically and patients can choose to opt out. Being able to choose in the way you want to be communicated to.”*

*“When you hear the cancer word, you stop listening. Having something like a brochure to look at later is nice.”*



# Mental & Emotional Wellbeing

## 1. Navigation and Awareness of Mental and Emotional Support :

- Reach out to new patients to ensure they have knowledge, resources and access to internal and external mental and emotional supports.
- Refer patients to counselling at diagnosis.
- Advertise non-BC Cancer support groups to improve access to appropriate support.
- Review existing models for peer support and navigation to identify best practices and adapt them to BC Cancer's context.

## 2. Mental and Emotional Support At Any Time:

- Create opportunities for peer support.
- Create a comfortable drop-in space for peer connection before, after and outside of appointments.
- Reinforce healthcare provider training on the value of emotional support for patients at diagnosis. (e.g., listening, holding a hand, offering comfort). Emotional support needs to be built into every interaction.

*“The stage 4 community knows so much. We have lived through so many medications and have so much knowledge, but it’s so isolated to stage 4. I want to share it.”*

*“I was in the Victoria center recently and there was a girl with a cart with books and puzzles and stuff, and I turned to her and said that was just amazing. [...] It made me happy to see that.”*

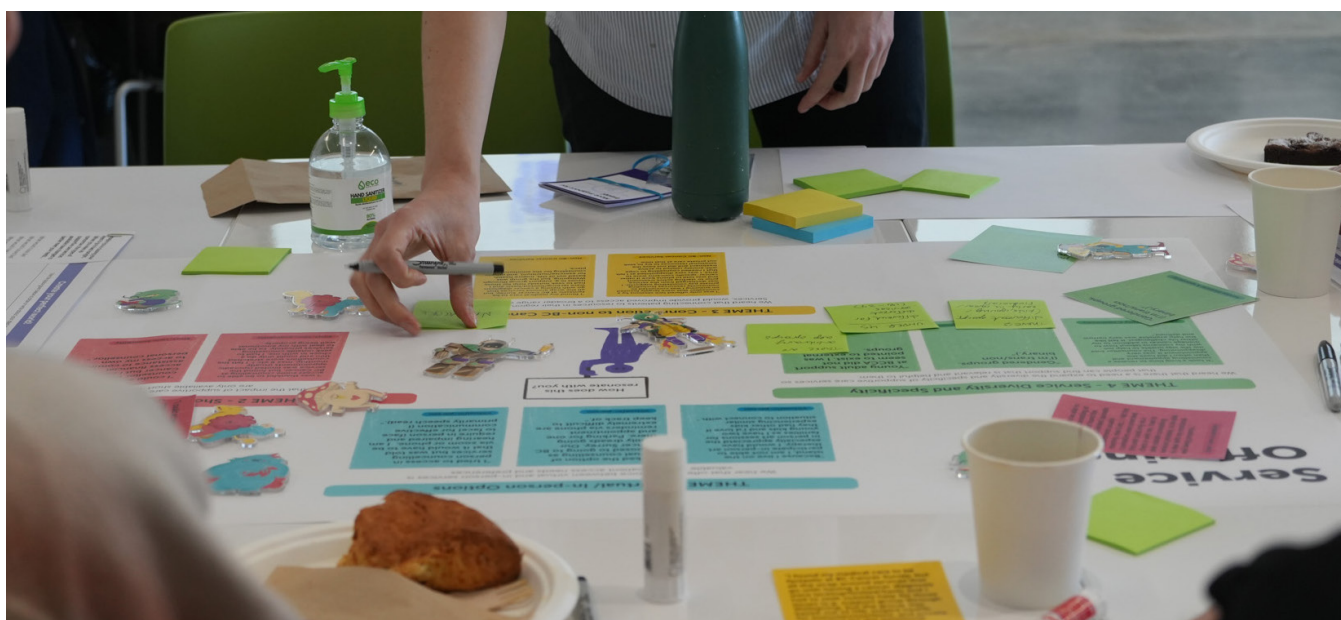


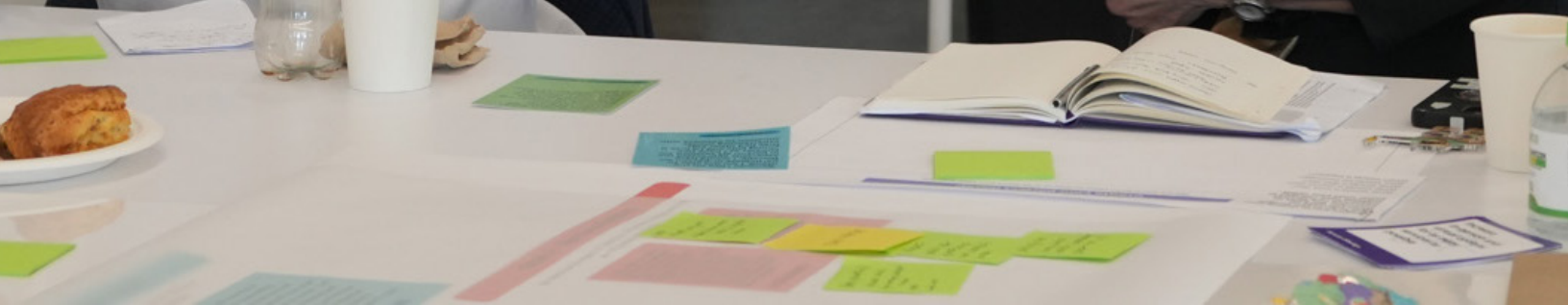
# Service Offerings & Accessibility

## 1. Service Offerings:

- Create pre-packaged patient education on subjects like nutrition, exercise, and procedures.
- Expand identity-specific support groups based on community needs (e.g. age, cultural, language-specific, etc.). Patient and family partners engaged on this project are not wholly representational of people seeking supportive care services for cancer across B.C., therefore deeper engagement will be required to develop more identity specific services.
- Provide financial support options to help patients access private services when BC Cancer cannot provide the same service or quality of service
- Expand service availability beyond standard working hours to improve access for working patients and family members.

*“I have heard in other provinces that there’s sometimes childcare options where there’s nannies that you can access. That would have been wonderful for us, because then my husband could have come with me to treatments or surgeries, instead of having to stay home and look after kids.”*





## 2. Service Access:

- Create an online booking system that patients can use to self-refer to supportive care services, track waitlists and manage their appointments. Consider designing a mobile app, given that many people track and organize their lives on their phones
- Provide patients with multiple delivery options to allow choice depending on the appointment topic or their accessibility needs. (Ex. in person, video call, or phone call)
- Offer a hybrid service delivery option where some appointments occur in person with virtual follow-up. Patients spoke to the benefit of a preliminary in person meeting, or having a critical once-a-year meeting in person, with more routine follow-ups occurring virtually for efficiency and ease of access.

## 3. Communication and Coordination:

- Improve communication between patients and families, supportive care providers and medical care providers using a trauma-informed approach to ensure smooth referrals and coordinated care.
- Improve communication and service coordination between BC Cancer and non-BC Cancer service providers.

*“Because of where I’m at in my treatment journey, I would prefer virtual because I have to be very, very careful with infection and not being around others that aren’t in my circle.”*



*“A difficult part of my cancer journey was being told that I wouldn’t be able to have children at all. That was very, very hard. [...] there was no communication between the fertility clinic and the oncologist when that happened. Then I went to an oncologist appointment, and he was gung-ho for the idea of me to have children. He wasn’t aware.”*

# Next Steps: 3 Year Roadmap

Insights gathered through this collaborative work have directly shaped the BC Cancer Supportive Care 5-Year Strategic Plan and informed the priorities outlined in its 3-year roadmap, as highlighted in the report. Over the next three years, BC Cancer's supportive care program will focus on improving awareness of services, expanding opportunities for mental and emotional support as part of cancer treatment, and strengthening service offerings across the province.

Several projects have already been implemented across each of these priority areas, including but not limited to the following:

## **Awareness of Services through information resources and navigation support**

An internal review of all supportive care information resources, including digital content on the BC Cancer website and paper-based materials available across the six regional cancer centres has begun. This work aims to reorganize and update content for clarity, consistency, and ease of navigation. It also includes improving the use of the existing e-bulletin and how supportive care content is presented on the BC Cancer website.

In a tangential project, the program is also designing an AI-enabled navigation tool to help patients more easily identify, understand, and access supportive care services based on their individual needs.

## **Mental and Emotional Wellbeing**

BC Cancer's Supportive Care program is in the process of developing a short, engaging orientation video to be disseminated across BC Cancer physicians, strengthening awareness and confidence, supporting timely and appropriate referrals, and reinforcing the integration of mental and emotional wellbeing into cancer care across the patient experience (i.e. at diagnosis, during treatment, post-treatment and long-term follow-up, and living with advanced disease).

## **Service Offering and Accessibility**

BC Cancer's Supportive Care program is in the process of developing a peer navigation group for patients and family members. This work aims to create structured opportunities for participants to explore real world navigation challenges, including understanding available supports, sharing lived experience, and identifying strategies for overcoming system barriers.

BC Cancer has recently expanded its cancer rehabilitation program, as a part of that, we're refreshing patient education materials related to nutrition, exercise, and other common procedures.

*Together, these initiatives will advance a more accessible, consistent, and compassionate supportive care system across the province, and BC Cancer will continue to look for additional opportunities to partner with patients and caregivers to support both the initiatives outlined here and additional improvements that emerge over time.*

# Acknowledgements

We are deeply grateful to the 39 individuals with lived experience of cancer treatment, either as patients or as loved ones of patients, who directly contributed to this work by sharing their insights and ideas through workshops held in April 2025, follow-up validation meetings and email correspondence. We sincerely thank each of them for their time, thoughtfulness, and enthusiastic participation in helping improve the cancer treatment experience for others. The following contributors gave their consent to be acknowledged by name:

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