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**KŌKUA MAU**  
“Continuous Care”  
**A Movement to Improve Care**

The Conversation Project is dedicated to helping people talk about their wishes for end-of-life care.

We know that no guide and no single conversation can cover all the decisions that you and your family may face. What conversations can do is provide a shared understanding of what matters most to you and your loved ones. This can make it easier to make decisions when the time comes.

Name: \_\_\_\_\_

Date: \_\_\_\_\_

# Your Conversation Starter Guide

How to talk about what matters to you and have a say in your health care.



the **conversation** project

## We can't plan for everything. But we can talk about what is most important — in our life, and in our health care — with those who matter most.

Talking with the important people in our life can bring us closer together. It also helps us create the foundation of a care plan that's right for us — a plan that will be available when the need arises.

The Conversation Project wants to help everyone talk about their wishes for care through the end of life, so those wishes can be understood and respected. We created this guide to help you start a conversation (and keep talking) so you can have a say in your health care — today and tomorrow.

It's also important to choose what's known as a health care agent or health care power of attorney — someone who would make health care decisions on your behalf if you became unable to voice those decisions yourself. Visit [Your Guide to Choosing a Health Care Agent](#) for guidance on picking an agent.

If you are completing this document on a computer, first save it to your desktop with a name you can easily find again. Then open your saved document and type in your answers. (Otherwise, what you type will not be saved.) Completing it on your computer will create a digital document that you can easily share with others.

### We'll help you take it step by step.

You can take your time! There's no need to say everything that matters in one conversation — you can start talking, then keep talking. It's all about what works best for you.

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## STEP 1

# Think About What Matters to You



To get ready to talk about what matters to you and your wishes for care through the end of life, it's helpful to gather your thoughts as a first step. You don't need to have the conversation just yet. Here are some helpful ways to think about what matters to you and prepare for your conversation.

> What does a good day look like for you?

SOME IDEAS Is it time with family or friends? Enjoying favorite everyday activities? What do you need to enjoy a good life — through the end of life?

> What or who supports you during difficult times?

SOME IDEAS Your faith, culture, family, friends, pets

> Try finishing this sentence:  
What matters to me through the end of my life is...

SOME IDEAS Being able to recognize my children; being independent; being able to spend time with the ones I love

That's your "what matters to me" statement.

Sharing it with people you trust could be a big help if they need to communicate with your health care team one day. They may need to share what's important to you and what you need to be able to have a good day. They also may need to decide what type of treatment you'd want to receive. Completing this guide will help you refine what you want them to know about what matters to you.

STEP 2

# Plan Your Talk



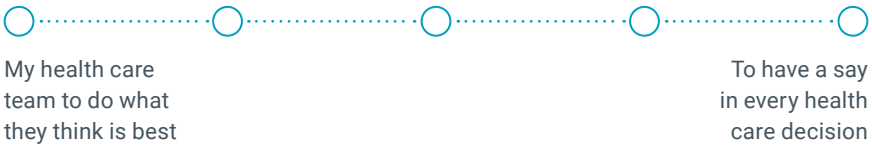
Having a say in your health care is more likely if you share how you feel about certain situations that could arise now, in the future, and toward the end of life.

For each statement below, mark the place on the line that is closest to what you think or believe about each statement now. There are no “right” or “wrong” choices — your answers are about what works for you.

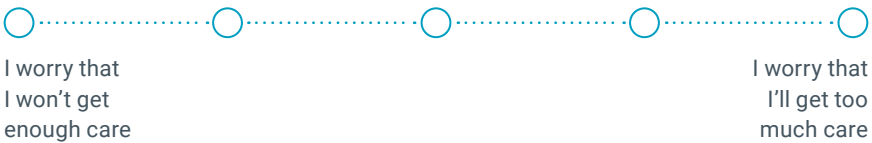
> As a patient, I'd like to know...



> When there is a medical decision to be made, I would like...



> What are your concerns about medical treatments?



> If I am diagnosed with a serious illness that could shorten my life, I would prefer to...



Not know how quickly the disease is progressing

Understand how quickly my disease is progressing

Any other notes you want to add?

>

> If you were seriously ill or near the end of your life, how much medical treatment would you feel was right for you?



I would want to try every available treatment to extend my life, even if it's uncomfortable

I would not want to try treatments that impact my quality of life in order to extend my life

> Where do you prefer to be toward the end of life?

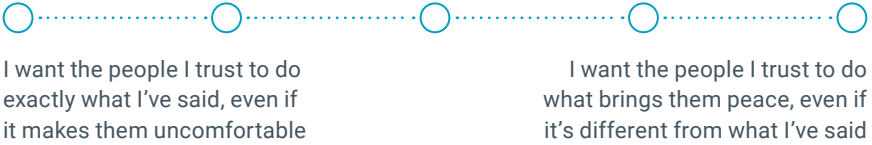


I strongly prefer to spend my last days in a health care facility (hospital, assisted living, or nursing facility)

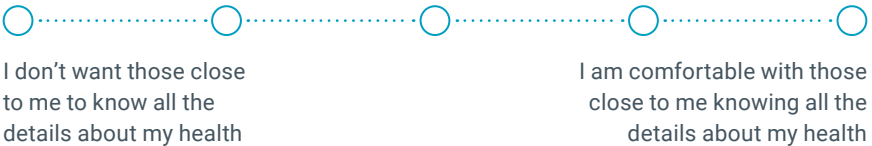
I strongly prefer to spend my last days at home

> Now, look at your previous answers. What do you notice about the kind of health care you said is right for you?

> If you weren't able to speak for yourself, would you want people to follow all your wishes or do what they think is best in the moment?



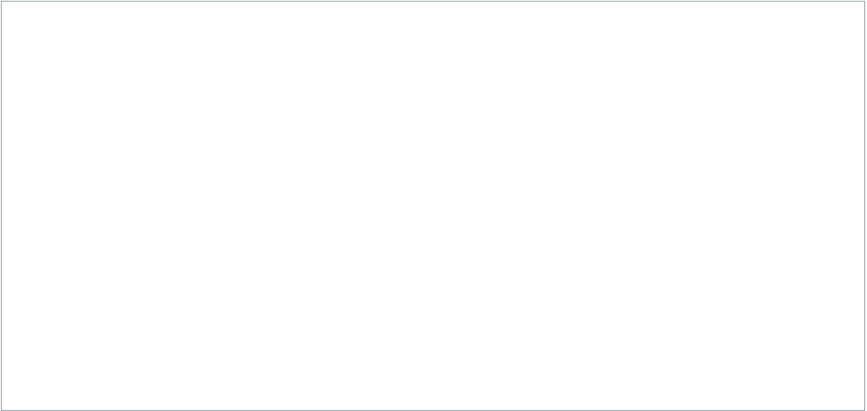
> When it comes to sharing information about my health with others...



> When I die...



- > What specific information would you want (or not want) shared with certain trusted people?



- > Look at your previous answers. What are the most important things for your friends, family, and health care team to understand about what matters most to you through the end of life?



### STEP 3

## Start Talking



How much do the people who matter to you know about what matters most to you? There may be some things they already know, and other things that you need to tell them. Sometimes we might think others know how we feel, but they don't. Conversations help make what we think and how we feel as clear as possible.

> Who needs to know what matters to you in your health care?

Check all that apply:

- |  |   |
|--|---|
| <input type="radio"/> Parent(s)  | <input type="radio"/> Trusted friend(s)           |
| <input type="radio"/> Spouse/partner(s)                                  | <input type="radio"/> Doctor(s)                   |
| <input type="radio"/> Chosen family member(s)                            | <input type="radio"/> Nurse practitioner/nurse(s) |
| <input type="radio"/> Adult child/children                               | <input type="radio"/> Social worker               |
| <input type="radio"/> Faith leader (minister, priest, rabbi, imam, etc.) | <input type="radio"/> Other: _____                |

> Where would you feel comfortable talking?

- |  |  |
|--|--|
| <input type="radio"/> At the kitchen table     | <input type="radio"/> Video chat or phone call |
| <input type="radio"/> At a favorite restaurant | <input type="radio"/> At my place of worship   |
| <input type="radio"/> In the car               | <input type="radio"/> Other: _____             |
| <input type="radio"/> On a walk                |  |

> The Conversation Project uses the saying, "It always seems too soon, until it's too late."

When will you start this conversation?



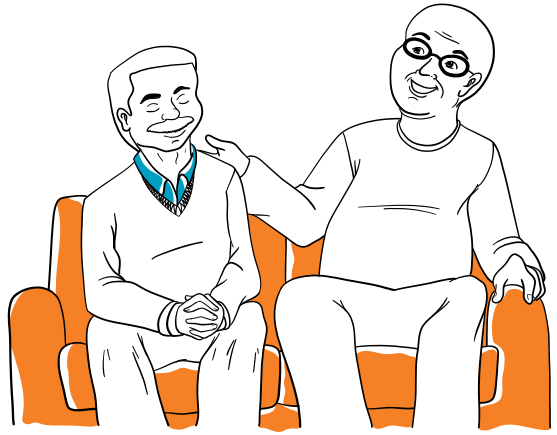
You've gathered your thoughts, written down your ideas, and picked your trusted people. Now, how do you begin a conversation?

> This list doesn't cover everything, but here are some things you can say to start talking.

- "I need your help with something."
- "Can you and I have a conversation about \_\_\_\_\_?"
- "I was thinking about what happened to \_\_\_\_\_, and it made me realize \_\_\_\_\_."
- "Even though I'm OK right now, I'm worried that \_\_\_\_\_, and I want to be prepared. Can we talk about some things that matter to me?"
- "Will you help me think about my future?"
- "I heard about the Conversation Project and answered some of their questions about things that matter to me when it comes to my care through the end of life. I'd like to talk to you about it."
- "When \_\_\_\_\_ died, do you think their wishes and priorities were respected toward the end of their life?"

> Here is a list of some other things you may want to cover when you talk.

- Do you have any worries about your health?
- What do you need to address to feel more prepared (examples: finances, property, legal documents, relationships, health care situations)?
- Do you have any fears, concerns, or mistrust about where or how you receive health care?
- Who do you want (or not want) to be involved in your health care?
- When you look ahead to the future, are there important events or dates you hope you're there for?
- Are there kinds of treatment you would want or not want (examples: resuscitation attempts, ventilation, feeding tube)?
- If your health condition changed, when would it be OK with you to shift from trying to cure an illness to trying to enjoy the end of life as much as possible?



## Tips for your talk

Imagine the conversation in your mind first. You can even write a letter that explains your values about the kind of care that works for you to figure out words that feel comfortable for you to use.

- 
- You can also consider having a practice conversation, so you feel as prepared as possible to have a “real” conversation.
  - You don’t have to talk about everything or talk to everyone in the first conversation. In fact, we suggest you keep talking over time!
  - Be patient. Some people are nervous or may need time to get ready to talk. Every time you start a conversation, it helps you come closer to making your wishes fully known. Keep trying.
  - You don’t have to lead the whole conversation; it’s important to also listen to what the other person says so you can build trust.
  - Nothing you say is permanent. You can always change your mind as things change in the future.
  - You may find out during these conversations that you and your trusted people disagree. That’s OK (no judgment!). The important thing is that you’re talking now and to keep talking — so you’re prepared in case your health changes.
  - You can share this guide, with or without your thoughts included, with your trusted people.
  - Think of these conversations as relationship-building opportunities to learn from each other about how to support and honor one another.

## STEP 4

# Keep Talking

Now that you've started the conversation, keep going! Talk to more people who may have a say in your health care. The more you talk, the more people you are close to will know what matters to you. And that makes it more likely that you'll get the kind of health care you want — now and through the end of life. Here are some things you can think about to keep the conversation going.

➤ When would be a good time to talk again?

**SOME IDEAS** It's a good idea to have another conversation when life changes happen, such as the birth of a baby, when family and friends are together for a holiday or visit, before a trip, or when a health issue is getting harder to manage.

➤ What might you want to repeat or explain again, so you're sure your trusted people understand what's important to you?

➤ Who do you want to talk to next time? Are there people (such as family members who may disagree) who should hear things from you at the same time?

➤ What do you want to make sure to ask or talk about next time?

# What to do next

Now, it's a good idea to record your conversation with an important legal document to be sure your choices are followed. This is called an advance directive. It has two parts.

## 1. Choose Your Health Care Agent, also called Health Care Power of Attorney

This is the part of the advance directive where you name the person you have chosen to make health care decisions on your behalf, if needed, as well as an alternate if your first choice is unavailable. As explained in this guide, be sure to have a conversation — and keep talking — with these people to be sure they understand what matters to you. You can find more information and suggestions in [Your Guide to Choosing a Health Care Agent](#)

## 2. Your Advance Health Care Directive: Instructions for my agent

This is the part of the advance directive where you describe your preferences and wishes for your health care if you cannot speak for yourself. These are many of the same things that you have thought about and discussed throughout this guide. Every state and most countries have their own advance directive forms. In the United States, the NHPCO (National Hospice and Palliative Care Organization) can help you find the right forms in your state ([nhpco.org/advancedirective](http://nhpco.org/advancedirective)). In Hawaii, visit [www.Kokuamau.org](http://www.Kokuamau.org). It's important to share your advance directive with more than your Health Care Agent alone. For example, if you pick an adult child to be your agent and have other children, they should all be aware of what matters to you in your health care and know who you have chosen as your agent. Talk to anyone who can help you have a say in your care through the end of life and provide copies of your advance directive to anyone who may need them. Share your advance directive with your healthcare provider. If you want tips on talking about what matters to you with your health care team, visit [Your Guide for Talking with a Health Care Team](#).

### Learn more and share

VISIT US  
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