Peace of Mind

Design Specification Audience: Web designers and developers



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HCDE 518, Autumn 2017

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Design Problem

America has a fast-growing aging population and with older age, the risk of getting dementia is elevated. Research shows that the number of people living with dementia in the US is projected to increase over time (Alzheimer's Association, 2016). People living with dementia need caregivers to take care of them, but caregiving is no easy task. Caregivers face many stressors in their lives such as social isolation, lack of time for self, friends, and friends, interruptions in their careers, financial stress, and unrelieved physical labor in caregiving (Archbold, 1982).

The stress faced by caregivers could also worsen over time; a one-year longitudinal study found that dementia patients had a significant worsening of self-care, memory, and behavioral problems and with that the average hours per week spent caregiving increased from 62 to 83 hours (Pratt, Schmall, Wright, & Cleland, 1985). Caregivers without social support and good coping mechanisms are also at an increased risk for poor mental and physical health outcomes (Pratt, Schmall, Wright, & Cleland, 1985; Goode, Haley, Roth, Ford, 1998; Neudorfer, 1991; Haley et al, 1996).

Design Goal

Build a website to help improve the quality of life among caregivers of people with dementia and help them better assist their care receivers.

Research Approach

In order to learn about the experiences and difficulties of caregivers and care receivers, we used two different types of research method. Due to the sensitivity and legal procedures for conducting research to our target audience, we were not able to expand our research methods to a broader audience. We focused on studying target users as people with dementia from early to mid stage and their direct / indirect caregivers including family members, hired caregivers, and volunteers.

Research Methods

• **Interviews** with 4 caregivers and 1 care receiver. We chose one-on-one interviews to learn about caregivers' and care receivers' experiences first-hand. We conducted the interview with several caregivers and one care receiver in an open-ended conversation style to foster trust and

empathy. In order to find interviewees in a short time span, we used convenience sampling.

- Literature Review to identify stressors and coping mechanisms amongst caregivers and people with dementia. We chose this method because there is a rich amount of empirical data from well-cited studies published in peer-reviewed scientific journals in this area. Our research aimed at answering the following two questions:
 - 1. What are the stressors faced by caregivers of people with dementia and their coping methods?
 - 2. What are the stressors faced by people with dementia and their coping methods?

The studies we found ranged from being cited 95 times to 745 times, which indicates that they have a high impact factor on the field. The sample sizes in the studies ranged from 60 to 240 caregivers of people with dementia and 92 to 135 people with dementia, which is a lot more people than we can recruit for our first study method given the short amount of time and sensitivity of the topic.

Research Findings

• Interview Findings

Caregivers

In our interviews with caregivers, it was apparent that they all had unique solutions to helping out their care receivers. For example, two caregiver used a word document to keep track of their care receiver's symptoms and medical results. Another found a solution online and used pleasant sounding door chimes to alert her when her husband left the home. One tech-savvy caregiver used Philips Hue lights to change the color of his mom's room whenever he wanted her to call him. However, not all of the caregivers we interviewed had high-tech solutions; one of the caregivers only used a television and radio to support their care receiver.

Person with Dementia

In our interview with an early-stage dementia patient, we found that she had to stop using most electronics over time due to their complexity. For example, her telephone and TV remote have become too complex for her to operate. The only electronic device she uses are a talking watch that says the time out loud at regular intervals and an audiobook player that has a very large "play/pause" button that she can find by touch.

• Literature Review Findings

Caregivers

The stressors faced by caregiver include helping their care recipient with self-care tasks that they're unable to perform independently anymore (e.g., bathing, feeding, and managing finances) and dealing with their care recipient's behavioral problems (e.g., angry, dangerous, or embarrassing behavior) and memory impairments (disorientation, wandering, and repetitive questions) (Goode, Haley, Roth, & Ford, 1998).

Caregivers who use good coping mechanisms to handle their stress are often better off. In a longitudinal study, researchers found that who count their blessings and have a positive outlook on life had less caregiver burden (Vitaliano, P.P., Russo, J., Young, H., Teri, L., & Maiuro, R.D.). Having social support, especially from family, was associated with better physical health outcomes over time and lower caregiver burden (Pratt, Schmall, Wright, & Cleland, 1985; Goode, Haley, Roth, Ford, 1998).

People with Dementia

Problems in daily functioning often are the reason for a decrease in quality of life in dementia patients (Graff et al., 2007). In addition, people with dementia often have psychiatric problems such as anxiety, depression, and psychosis - up to 90% of people with dementia have psychiatric comorbidities (Plassman et al., 2007).

Information and emotional support improves the quality of life of patients and their caregivers (Graff et al., 2007). Researchers also found that cognitive stimulation could help improve dementia patient's cognitive outcomes (e.g. delayed memories and problem solving) (Quayhagen et al., 2000).

Design Decision

The research findings helped us gain insights on both caregivers and care receivers. While we originally decided to design a solution for both caregivers and their receivers, we changed our design direction because it was apparent through our research that care receivers' daily life function depends greatly on their caregivers' support and performance. Helping to improve the quality of life for caregivers can consequently help them provide better care for their care receivers. How to design something that better suits caregivers' needs was the biggest design challenge we faced.

Our Solution

Peace of Mind is a website that provides caregivers of people with dementia with an online platform to connect and share solutions with each other. In the website, caregivers can gain and give social support and advice about unique caregiving situations.

It has a community forum where caregivers could ask and answer questions and a social support feature where caregivers can send private messages to each other. The website also has a "How To" feature where caregivers can learn about unique technical solutions and projects to better assist their care receivers. It also has a store section where caregivers can read about and purchase products for their care receivers. Additionally, it has a journal for caregivers to keep track of their care receiver's medication, monitor their homes, plan fun events, record their own daily mood and to-dos, and write about what they are grateful for each day (gratitude diary). Users are encouraged to use the website with "Karma Points" - the more they use the website, the more points they can gain.

Users

Primary Users - Familial caregivers of people with dementia. This is most commonly a spouse or adult daughter who oversees day-to-day caregiving.

Secondary Users - Hired caregivers who are recurrent participants in caregiving, volunteers who are recurrent participants in caregiving, or caregivers in assisted living homes.

UI Explanation

Because our audience skews older, the UI of the website is intended to be bright, and legible, and easy to navigate. We chose an easy-to-read Arial black font against a white background, large headers, and simple, square buttons throughout our site. Photographs are neutral to positive in tone, and feature older families and individuals. Iconography is intended to be playful and inviting. A playful and relaxed cursive font is used for certain title fonts, including the font of our logo. Accent colors are intended to be cool and calming. Copy is intended to be warm and inviting without appearing artificial and contrived.

Project Scope

Our final product contained basic functionality of our core features that met our user needs. Core features included creating a profile, a community section for interacting with other caregivers; a store, how-to, and search section for solving problems; and a journal section for expressing gratitude.

Advanced functionality within these features was not included due to lack of available production time and lower prioritization.

Features In Scope

- **General:** Navigate to main areas of the site, create an account, upload a profile photo, view profile
- **Community:** View the community forum, view individual community forum posts, create a community forum post, message another user
- Store: View Store home, view individual product page for Philips Hue light
- **How to:** View How To home page, view How To category page, view individual How To page for setting up a Philips Hue light
- Journal: View medical records, home monitor, planning (calendar), gratitude diary, to-do list, and mood.
- Search: Search for a sample topic and view result page.

Features Out of Scope

- **General:** Adjust settings, edit profile, premade avatar options, screen name generator, privacy features, animated interactions
- **Community**: Add a friend, earn Karma points, "like" a forum post, comment on a forum post, full chat functionality,
- Store: Sort by "works with" or use case, affiliate marketing checkout
- How to: Add a How To article, view articles by tag, sort articles
- Journal: Add and edit entries, share gratitude journal entries
- Search: Filter search results

Design Rationale

While our initial idea was to build an artificial intelligence system for dementia patients, our user research led us to understand that we could better help dementia patients by helping their caregiver(s). In our research, it also became obvious that dementia patients struggle with using technology. Their symptoms

change so dramatically and suddenly that designing an AI system that meets the needs of most dementia patients would be difficult, if not impossible.

This led us to the idea of building a support community for dementia caregivers so that they can empower and learn from one another. A website (as opposed to a smartphone app) was necessary to meet our user needs because only 42% of seniors over the age of 65 own a smartphone, while 67% of those same seniors have a home internet connection (Anderson & Perrin, 2017). Because our primary audience is familial caregivers, especially spouses of patients, we wanted to make our product as accessible to older audiences as possible.

Based on research findings, we created five primary sections of the website to meet the user needs and six sub-features under Journal, shown below in the following charts.

Core Feature Chart

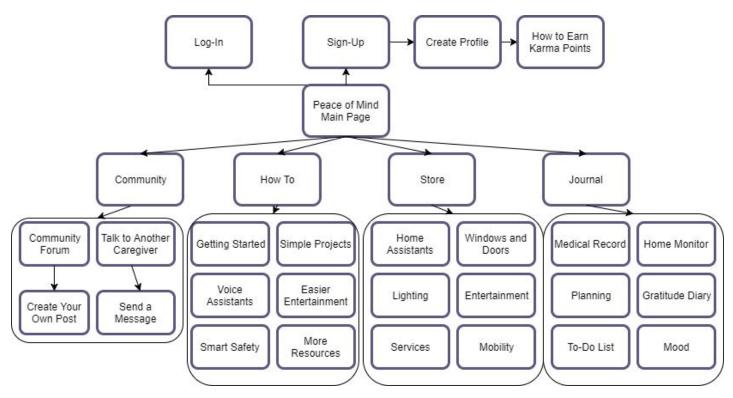
Feature	User need met	Supporting Research
Community	Emotional support	One of the caregivers we interviewed was not in any emotional support groups, but she found strength by attending church weekly, praying, and having support from her husband and hired caregiver.
		Furthermore, literature review findings showed that caregivers with high levels social support initially had better physical health outcomes over time. (Goode, K. T., Haley, W. E., Roth, D. L., & Ford, G. R.)
How to	Problem solving	Based on our interviews with the caregivers, we found out that caregivers have their own solutions for specific problems that their care receivers had.
		Caregivers needed a platform where they can exchange information and share experiences on ways to solve specific questions.
Store	Problem solving	One of the caregivers we interviewed used the Philips Hue as a tool to remind his care receiver for specific task. We were inspired by this finding and wanted to introduce products that could potentially help our target users such as smart home assistant, lighting therapy.
Journal	Problem solving	Based on our interview findings, caregivers need to technology products that will help them track their care receiver's activities.
		For example, one of the caregivers that we interviewed was using Word Document to keep her notes and filled in more details she could remember later. From then on, she included anything and everything that could be relevant, highlighting particularly important points.
Search	Problem solving	Based on our interviews with the caregivers, we found out that caregivers have their own solutions for specific problems that their care receivers had. Caregivers needed a feature to search for specific problem related information.

Journal Feature Chart

Feature	User need met	Supporting Research	
Medical Record	Problem solving - Tracking	Based on our interview findings, caregivers need to technology products that will help them track their care receiver's activities.	
Home Monitor	Solve problems in daily functioning	Compensatory strategies were used to adapt activities of daily living to the disabilities of patients, and environmental strategies were used to adapt the patient's' environment to their cognitive disabilities.	
		Community occupational therapy should be advocated both for dementia patients and their caregivers, because it improves their mood, quality of life, and health status and caregivers' sense of control over life. Effects were still present at follow-up. (Graff, M. J. L., Vernooij-Dassen, M. J. M., Thijssen, M., Dekker, J., Hoefnagels, W. H. L., OldeRikkert, M. G. M.)	
Planning	Problem solving - Tracking / Reduce depression	Behavioral therapy: pleasant event which involves planning, and executing pleasant events showed significant improvement in depression symptoms. (Teri, L., Logsdon, R.G., Uomoto, J., McCurry, S.M.)	
Gratitude Diary Counting blessing to reduce caregivers burden		Based on our literature review findings, counting one's blessings (i.e., comparing myself to others who are less fortunate) and having a positive outlook on life were associated with less caregiver burden at both baseline and follow-up. (Vitaliano, P.P., Russo, J., Young, H., Teri, L., & Maiuro, R.D.)	
To-Do List	Problem solving - Tracking	Based on our interview findings, caregivers need to technology products that will help them track their care receiver's activities.	
Mood	Problem solving - Tracking	Based on our interview findings, caregivers need to technology products that will help them track their care receiver's activities.	

Process Flows

The process flow diagram below shows the high-level navigation of the website.



User Tasks

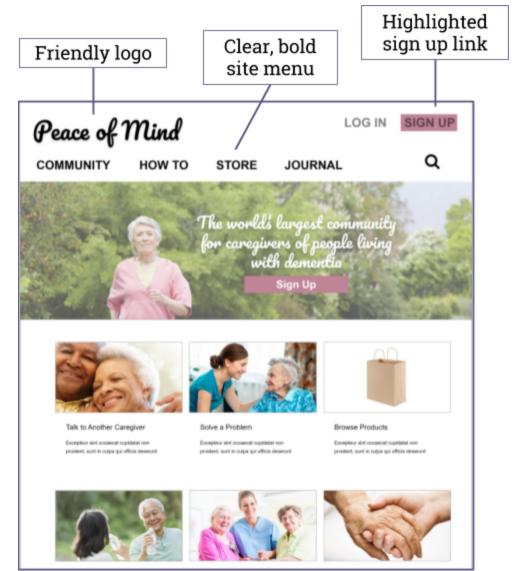
- 1. Sign up and create a profile
- 2. Find and "Like" Jane Smith's Community Forum post
- 3. Create a Community Forum Post
- 4. Visit the Community Talk to Another Caregiver page and send a message to Jane Smith
- 5. Learn to set up a Philips Hue light in the Getting Started area of How To
- 6. View Philips Hue in the Store
- 7. Visit the Store Home
- 8. Visit Journal and explore the six types of logging
- 9. Return to the homepage and perform a search

Functional Prototype

The prototype can be found at this link: <u>https://xd.adobe.com/view/d987d949-d3b0-476f-9e39-32e61f2c277c/</u>

Annotated Webpages

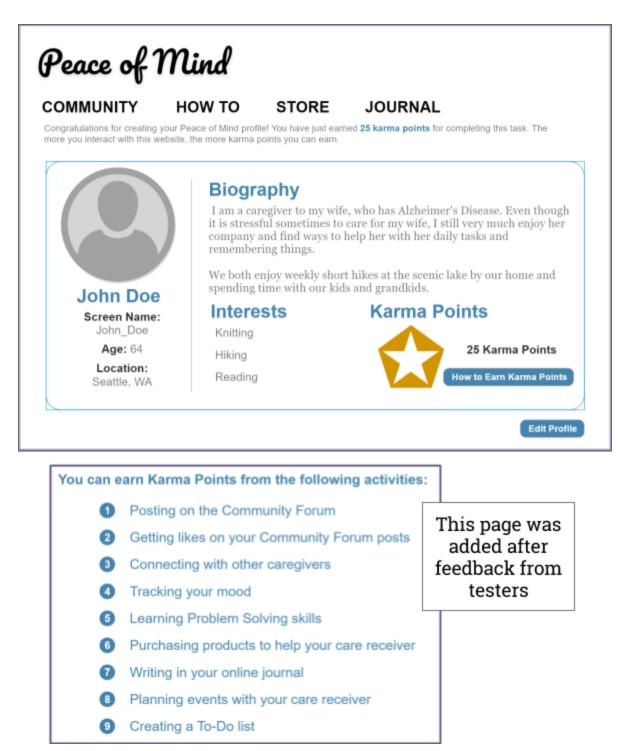
Peace of Mind Homepage



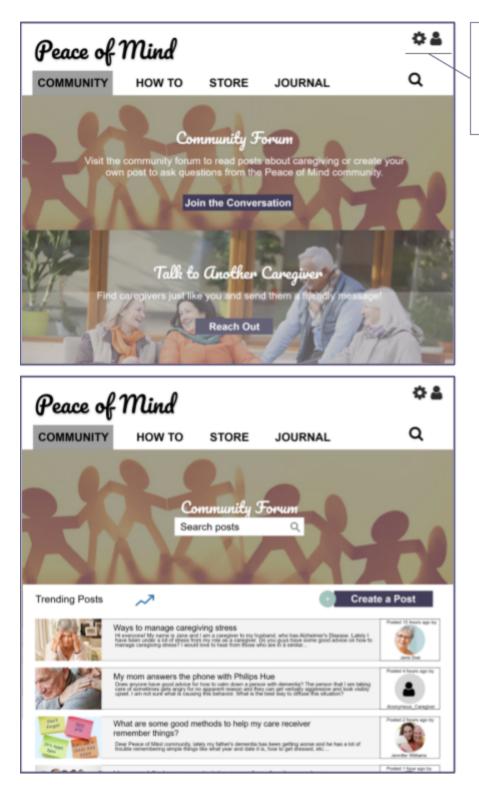
Sign Up Process

Peace of Mind	LOG IN SIGN UP
COMMUNITY HOW TO STORE JOUR Home > Sign Up Sign Up Name 	RNAL Q
Password	info to share
Select a Profile Picture (optional) Select a picture for your profile photo. Drag and Drop any photo from your como circle on the left to upload your profile photo. Back	

User Profile



Community Home & Forum

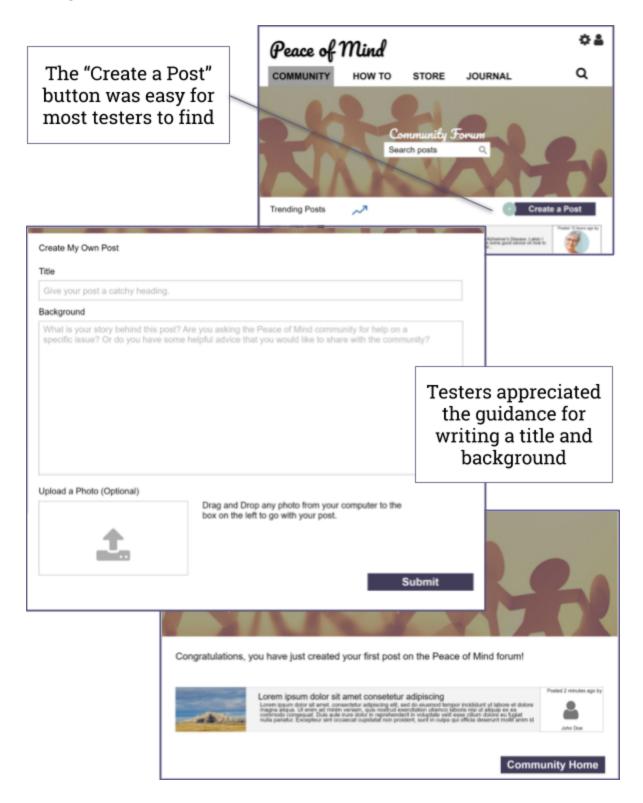


Once logged in, settings and profile links are available

Community Forum Post

Peace	ofγ	Nind			¢ a	
COMMUN	-	ноw то	STORE	JOURNAL	Q	
		ommund earch posts	system help	elps promote	-	
Disease. Lately some good adv in a similar posi Right now, I am joy in doing the the time. Even t to Friday. If I do expenses. 10 Comments	y name is J I have bee ice on how tion as I am dealing wit things that hough I am n't, then I c Hi Jane, it si health. I am in the past to spiritual guid If you're not up yoga clas	lane and I am a c in under a lot of si to manage careg n. th a lot of depress I used to enjoy a n past the average ant affort to pay 100 Likes 100 Likes 0 deal with you h also a caregive religious, meditation ses with your busy of	tress from my role iving stress? I wo sion and anxiety i nd I think that mig e age of retirement for my husband's Add C 101 Likes ane church as my br and yoga is also rea- schedule, but you cai	sband, who has Alzheimer's as a caregiver. Do you guys uid love to hear from those w n my day-to-day life. I no long ht be because I'm so exhaus t, I still work a 9-5 job from N medical bills and our living omment ht now with caregiving, work, and s mid-stage dementia. Some helpf ek social support from my family m others and sisters and praying help liv helpful for dealing with stress. It n start slow and try out a class to sr ant to try doing yoga at home.	who are Jane Smith ger find sted all Monday dealing with your mental ul methods that I have tried sembers and to seek as me to manage my stress. may seem like a lot to take	
Martha Wilson	caring for m caregiving a write down t their blessin on this site t	y father, who is appro a lot more stressful. V three things I'm grate ags and have a positi	caching late-stage de Vhat I found helpful fe ful for each day. This ve outlook on life are online journal. I use	mment you lately. Caregiving is not an ea mentia. I, too, have gone through or me is to keep a daily journal whe is backed by research showing th less likely to experience caregiver this often and I find it calming to wr	depression and it makes are I track my mood and at caregivers who count burden. There's a feature	
	0	80 Likes	📮 Add Co	mment		

Creating a Forum Post



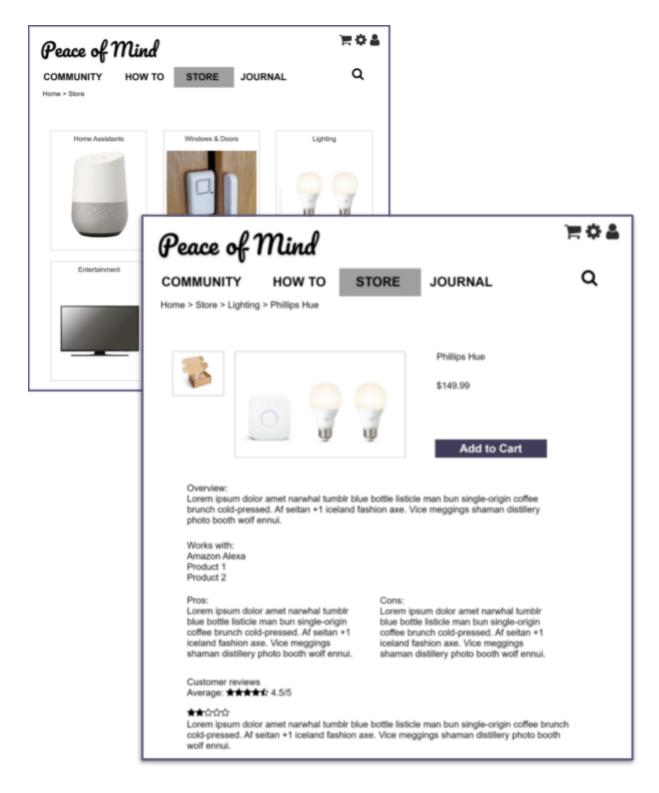
Talk to Another Caregiver



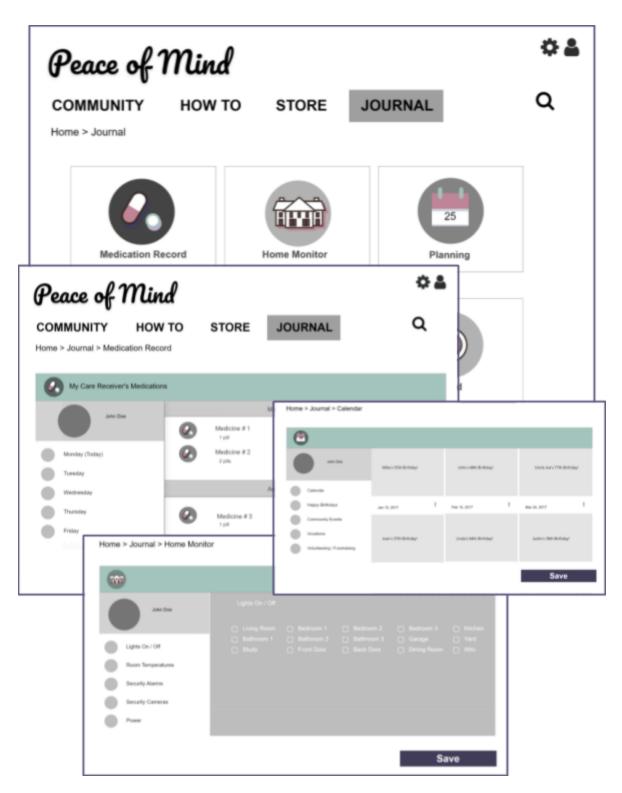
How To Section



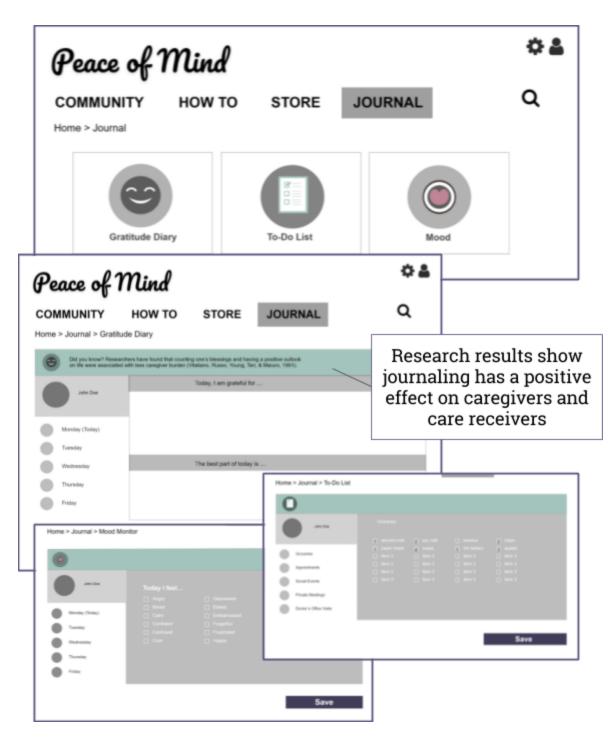
Store Section



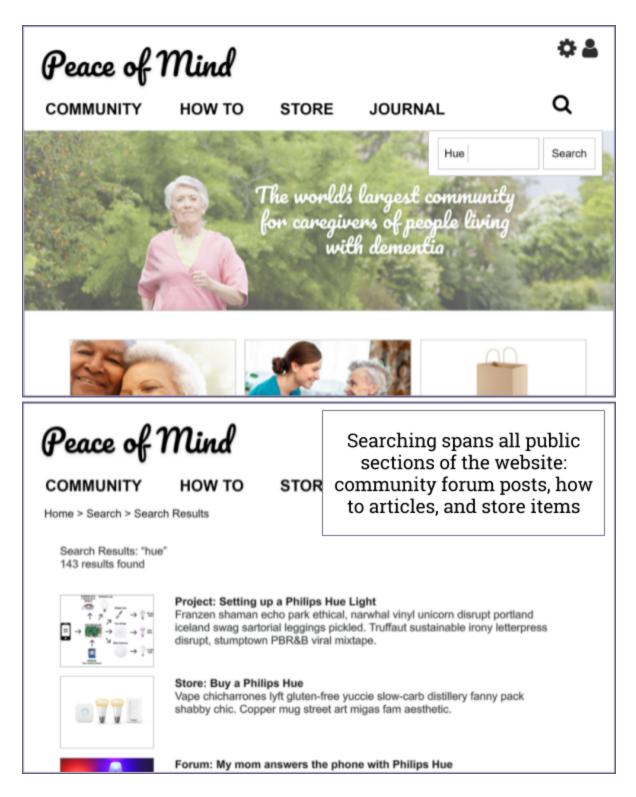
Journal Section 1



Journal Section 2



Search



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