| 1 | The Experience of Gender in Spousal Caregiving: A Phenomenologica |
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| 2 | Psychological Study (Greece) |
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| 21 | ABSTRACT |
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| 22 | Purpose/Objective: To explore how spousal caregivers of older people undergoing |
| 23 | rehabilitation experience gender within the Greek community. |
| 24 | Research method/Design: A psychological phenomenological design and analysis |
| 25 | were used to illuminate the unique meanings eleven spousal caregivers attribute to |
| 26 | their experience of gender by gathering qualitative data via interviews. |
| 27 | Results: The data provided an insight into the structure of the experience of gender |
| 28 | for the spousal caregivers as a normative diachronic identity in a succession of |
| 29 | phases: normative constitution, alienation, and reparation. |
| 30 | Conclusions/ Implications: The findings highlight the influence of gender |
| 31 | stereotypes on spousal caregivers' self-concept, agency, caregiving evaluations, and |
| 32 | practices, emphasizing the importance of adopting an intersectional perspective in |
| 33 | future research and interventions, considering various factors such as ethnicity, |
| 34 | gender, sexuality, age, power dynamics, and cultural norms. Spousal caregivers |
| 35 | experience alienation on entering the caregiving journey, with gender-related |
| 36 | vulnerabilities affecting their psychological well-being. Addressing these |
| 37 | vulnerabilities can improve caregivers' mental health and foster effective coping |
| 38 | strategies. The study emphasizes the moral aspect of caregiving, highlighting the |
| 39 | relationship between a sense of obligation, feelings of guilt, gender norms, and |
| 40 | motivations calling for challenging self-sacrificial morals and societal norms |
| 41 | associated with them to empower caregivers to prioritize their well-being while |
| 42 | maintaining their caregiving motivations. This shift in perspective can lead to a more |
| 43 | positive and fulfilling caregiving experience. |
| 44 | Keywords: gender, spousal care, rehabilitation, psychological well-being |

45 Impact and implications statement:

Innovative framework for understanding spousal caregiving's psychological effects
from a gender perspective.

• Highlights the role of gender norms in shaping caregivers' experiences, affecting their well-being, agency, emotional responses, coping and moral decision-making.

• Advances gender and care knowledge, informing research and therapy to enhance caregivers' well-being and caregiving experiences.

52 Introduction

A large and robust literature documents higher rates of psychological morbidity involving emotional distress, depression, anxiety, and social isolation among informal caregivers compared with no caregiver comparison groups, suggesting that caregiving is a significant public health issue (Schulz et al., 2020b). Research demonstrates significant well-being declines as the person enters the caregiving role, further deterioration in well-being as care demands increase, and gradual recovery when the demands of care provision decline or cease (Schulz et al., 2016). Additionally, an abrupt change in lives, lack of sense of agency and lack of perceived choice are linked with increased levels of burden and depression (Schulz et al., 2012; Zygouri et al., 2021). Given that the availability and continuity of informal caregiving is a global requirement, providers and policymakers must have access to comprehensive caregiving research that provides meaningful subgroup analyses exploring the subjective experiences of caregivers, including their attitudes, values, preferences, feelings, and expectations, helping caregivers sustain their motivation for caregiving (Harvath et al., 2020; Schulz et al., 2016; Sharma et al., 2016).

Gender and caregiving

In the caregiving context, gender is a central phenomenon that warrants examination. While research provides conflicting evidence, some studies suggest that

being a woman and a care recipient's wife predict adverse psychological effects, but others find no gender differences (Bom et al., 2019; Sharma et al., 2016; Xiong et al., 2020; Yee & Schulz, 2000). Moreover, apart from the equivocal and inconsistent evidence, there is an overemphasis on women caregivers, neglecting data on men, who increasingly assume caregiver roles (Sharma et al., 2016). Evidence suggests that it is not the objective conditions as more hours of care and more caregiving tasks are performed by women, but the subjective evaluation of the caregiving workload as well as the subjective evaluation of its effects that may explain gender differences in psychological morbidity (Pinquart & Sörensen, 2006; Savundranayagam & Montgomery, 2010; Swinkels et al., 2019). The influence of gender in caregiving is structural and not easy to discern as it intersects with several other variables such as culture, ethnicity, age, family relations and socioeconomic status, but these have seldom been considered in research studies highlighting the complexity of this phenomenon (Sharma et al., 2016; Swinkels et al., 2019).

Context of research

In Greece, family caregiving is prevalent (Katrougalos & Lazaridis, 2016), with an estimated 34% of the population providing informal care (EQLS, 2016). A lack of long-term care facilities has hindered women's participation in labor force, leading to part-time work or early retirement (Ziomas et al., 2018). Traditional patriarchal beliefs in Greek society have historically justified gender inequality, although there is a transition towards more individualistic values that will potentially impact women's position in family and caregiving arrangements (Georgas, 1989; Tsiganou J., 2021).

Aim of the study

Given the above, this study uses psychological phenomenological methods and analysis to capture the subjective perspectives of caregivers on the influence of gender on their sense of self and agency to gain valuable insights into the challenges presented by caregiving roles and their connection to the well-being of caregivers. The research question is: How do spousal caregivers of older people undergoing rehabilitation experience gender within the Greek community? The objectives to support the inquiry are a) to describe the gendered patterns of perception, thought, feelings and behaviour of spousal caregivers, b) to investigate the normative structures that constitute the experience of gender, and c) to explore the sense of agency in the experience of gender.

This research is exploratory and not grounded on any hypothesis or prediction. The research objectives' content, formulation and rationale are based on a constructionist epistemological position that challenges the notion of universal and objective truth, instead emphasizing the socially constructed nature of knowledge and the importance of multiple perspectives in understanding and interpreting reality (Schwandt, 1994). Under this epistemological stance, a fundamental assumption of this study is the belief that culture exerts influence on people's lives and that knowledge is inherently dependent upon communities and thus governed to a large degree by normative rules that are historically and culturally constituted (Gergen, 1985; Guba & Lincoln, 1994). From this perspective, gender is viewed as a dynamic, socially constructed concept rather than a fixed identity (West & Zimmerman, 1987). Contemporary theorists highlight the complex interplay of biology, culture, power dynamics, and individual identity in shaping our understanding of gender (Butler, 2011; Fausto-Sterling, 2020). They advocate for a more nuanced understanding of masculinity and femininity, recognizing a spectrum of gender expressions (Connell,

2005). In line with this perspective, our research examines the dynamic nature of gender and its effect on caregiving experiences.

122 Methods

We followed the Journal Article Reporting Standards (Levitt et al., 2018) and the Consolidated Criteria for Reporting Qualitative Research (Tong et al., 2007) to conduct this study and report the findings. All data is available from the corresponding author upon reasonable request. This study is not preregistered.

Design

We used a psychological phenomenological design to illuminate the unique perspectives individuals attribute to their experiences of gender by gathering qualitative data via interviews (Langdridge, 2008). We employed phenomenological psychological reduction to suspend our biases and assumptions and remain receptive to participants' experiences, adopting a descriptive approach to thoroughly analyze the data (Englander, 2016; Giorgi et al., 2017).

Participants

The sample consisted of 11 spousal caregivers of older individuals needing rehabilitation due to brain injury-related disorders. Seven participants are female, four are male, ten are Greek, and one is Albanian. Of the 11 participants, seven identified as women, while four identified as men. Minimum age 62 years old, maximum age 83, mean average age 69.9. The minimum time for providing care is nine months, and the maximum is twenty-five years. The sample predominantly consist of retired individuals from working-class socioeconomic background. All participants are Greek residents who speak and write in Greek (see supplementary 1 for participants' demographics).

Purposive sampling was employed to select participants who met predetermined criteria and could offer comprehensive insights into the phenomenon under investigation (Patton & Schwandt, 2014). The study aimed to include caregivers from diverse socio-economic and ethnic backgrounds and at different stages of the caregiving journey but also sought some homogeneity to ensure relevance and personal significance for the respondents (Moustakas, 1994). To achieve homogeneity, considering the highly adverse impact of caregiving on spouses and partners' carers and the intersection of caring practices with the caregivers/ care receiver relationship and care receiver medical needs, the study selected caregivers who lived with and provided primary care to their spouse or partner in need of rehabilitation (Lafiatoglou et al., 2022; Zygouri et al., 2021). Although the initial aim was to include caregivers from various gender and sexual orientation backgrounds, the final sample consisted of cisgender heterosexual individuals due to the demographics of the rehabilitation clinic and the nature of the caregiving context.

Data saturation was achieved with 11 participants (Strauss & Corbin, 1998).

Procedures

The recruitment setting was a specialized public rehabilitation clinic in an urban area providing free-of-charge services to individuals requiring rehabilitation. All the participants were recruited through invitation flyers and participant information sheets that included information about the principal researcher and her reasons and interest in the topic, the research purpose, risks, rights, confidentiality, dissemination and contact details. There was no prior relationship between the researchers and the participants. All interviews were conducted once, face to face, by the principal researcher, a white woman psychologist and a PhD researcher trained in qualitative interviews. No one else was present in the interviews. One interview was

conducted at the clinic, one at the University and nine in the participants' houses and lasted from 50 to 90 minutes, ensuring sufficient participant engagement. The interviews were audio-recorded with high-quality equipment. The interview phase lasted three months (09/21-11/21). The interviews were transcribed verbatim immediately after completion to ensure data reliability, along with reflective notes (Creswell, 2012). No participant withdrew from the study. Bevan's (2014) interview method for descriptive phenomenological research was used to ensure consistency across phenomenological theoretical assumptions, strategies and techniques (see supplementary 2 for the interview protocol). Not all questions were asked, as questioning was based on the participant's responses. Gender identification was ascertained by open-ended questions (Nunner-Winkler et al., 2007). The pilot interview was included for data analysis due to the richness of the information collected.

Data Analysis and Credibility of the Study

Giorgi's phenomenological method guided the data analysis process (Giorgi, 2009). Initially, raw data was repeatedly read to grasp its overall essence. Everyday descriptions were transformed into third-person meaning units, preserving participants' experiences (Giorgi, 2009). Reflection and imaginative variation were employed to express these units in psychological language, focusing on the study's investigated phenomenon, the experience of gender (Giorgi, 2009). Specific meaning units related to the experience of gender were identified and organized in temporal order, representing the experiential structure: Normative Constitution, constructing gender identity, alienation: disruption of the gendered sense of self, and reparation: reconstitution of gender identity (Giorgi, 1985). These transformed units were synthesized into coherent descriptions of individual experiences and a general

overview based on multiple cases, encapsulating the phenomenon of gender among older spousal caregivers (see supplementary 3 for a data analysis example). To ensure validity, individual structures were compared (Giorgi, 1985). Data credibility was ensured through prolonged engagement, accurate transcription of high-quality recordings, detailed documentation, and peer debriefings. The study's credibility was further supported by purposeful sampling, a structured interview protocol, and phenomenological reduction (Giorgi, 2009).

Ethical considerations

All research activity complied with the Declaration of Helsinki, followed good practice guidance (E.U. Reg no. 536/2014), and adhered to the Charter of Fundamental Rights of the European Union, 2000/C 364/01. This study has received approval from the University's Research Ethics Committee, where the principal investigator is affiliated. All the participants provided informed consent for participation in the research (see supplementary 4 for participant consent form).

208 Results

The Structure of the Experience of Gender in Spousal Caregiving

The experience of gender, grasped by the participants at the time of the interview, supports insight into the structure of the experience of gender for the spousal caregivers as a normative diachronic identity characterized by an intricate interplay between gender identity and agency. Through a temporal lens, the phenomenon unfolds across three discernible phases: Normative Constitution, Alienation, and Reparation. Participants cross these phases, revealing how gender norms shape their identities and caregiving roles, resulting in periods of estrangement and endeavours to reclaim agency, reaffirming their gendered sense of self in the caregiving context. The phases are described below, including illustrative participants'

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quotations. The participants are quoted as 'wp' for women participants and 'mp' for men participants, reflecting their self-identified gender identities, followed by a case number based on their participation order.

Normative Constitution: Constructing Gender Identity

The participants' experience of gender begins with the anticipated gender roles within the heterosexual marriage institution. These roles are considered normative, involving specific evaluative criteria shaping ideals of gendered existence. One caregiver passionately stated, "God created us to have a family. This is the destiny of humanity" (mp2), emphasizing how spousal caregivers perceive family creation within heterosexuality as divine and central to personhood. The heterosexual marriage institution operates as an organizing principle in life, legitimizing gender identities, roles, and hierarchies through socialization, as illustrated in a caregiver's narrative: "I did a lesson to my teenage granddaughter: You will marry soon, and the man will return from work, and he will not find dinner. He will pardon you once but tell you again: return to your mother; you do not know anything" (mp2). Gender recognition with marital roles assigns discrete duties and liberties to each gender, with one participant sharing her perspective, stating, "The men do their chores out of the house. Always, the woman is at home and with the children. The woman is the 'other'" (wp9). For men caregivers, their self-worth is deeply intertwined with their role as breadwinners, where honor, independence, and good citizenship hold value. One caregiver articulated, "Be right, decent, a man who looks at himself and not what the other person is doing... I have earned respect in my work, and even today, people speak to me respectfully" (mp3). In contrast, womanhood's essence is profoundly connected to the roles of a wife and mother, celebrated as a natural gift and the foundation of femininity. One participant passionately declared, "The woman has all

the gifts. She is both a wife and a mother! Women are valuable!" (wp4), illustrating how women are expected to fulfil these traditional roles regarded as essential to their identity.

demonstrating love within their families. Participants emphasized this, affirming,

"When a woman is right, she must stand by her family" (wp8). Another participant

The significance of women further revolves around nurturing relationality and

echoed a similar sentiment: "As a woman, I care for my husband as I cared for my father. I was caring for the elders, and I was nurturing the kids" (wp9). relationality While women volunteer for the feminized caregiving roles, men's participation is excused. One participant proudly remarked, "We have taught the men to abstain from household chores! What can men do to you? Well, they cannot do anything!" (wp9). By emphasizing women's distinctiveness, caregiving solidifies as unnatural for men. A participant captured this sentiment: "As a man, morally, I support my wife. But physically, I do not. I cannot do the laundry of my wife's underwear. From within, I cannot, I do not underestimate the woman, but I cannot" (mp6).

While gender norms influence both genders, women are disproportionately affected, shaping their self-perception and agency contingent on men's validation. In a defining moment, one participant shared her thoughts on being a woman, stating, "As a woman, I love to have next to me a man who loves me, respects me, does not talk to me badly, and does not offend me. That bothers me. That may kill me" (wp10). A participant further contributed to this perspective, asserting, "The woman is in you. It is the nature, the position of the woman" (mp6),

In the accounts, the construct of gender situates the participants' cognition in understanding their self and actions and understanding of the world and others,

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structuring a continuous, rational, and coherent sense of self while shaping individuals' agency.

Alienation: Disruption of the Gendered Sense of Self

The emergence of the caregiving journey introduces a profound sense of alienation, disrupting the participants' understanding of themselves as gendered subjects. As one participant expressed, "I feel imprisoned... My life is over... Sometimes my brain gets foolish, and I want to get the cell phone, and I get the bread... I suffer and am alone" (mp1), revealing the emotional instability experienced by men. Once firmly attached to their masculine identities, these men are detached from traditional masculine activities and deeply immersed in feminized caregiving roles. This sense of alienation takes various forms, including loneliness, isolation, desperation, emotional vulnerability, depression, and introspection, which significantly impact cognitive functions, as the participant suggested. Men caregivers wrestle with a tension between their desire for personal freedom and the preservation of their reputation. As one caregiver emotionally shared, "I am in prison... I have been excluded... I could have left her to die on Saturday by not giving her oxygen... I have been told to hire a woman caregiver twenty-four hours a day. No, I cannot do that. If another woman is sleeping here, you will be tempted one day... Everything I do on my own. I have taken it upon myself" (mp3). Women caregivers also experience alienation, which manifests as a loss of the traditional marital relationship and heightened nostalgia. Their desires centre on a return to the established gender order, as one participant expressed, "Everything has changed... He was a hyperactive man, and of course, I was right behind him...

Sometimes I tell him: become the man you were" (wp5). This sense of longing is

echoed by another participant, who added, "I sense myself being left behind... I grew old... I do not even want to attend celebrations anymore" (wp4).

Women caregivers, too, engage in heightened emotional and physical labor that leads to experiences extending beyond depression as fear, hyperarousal, restlessness, and medication reliance. These challenges are further intensified by rumination. As one participant distressingly described, " How can I leave him? He calls me all the time. If I am not there, he feels insecure... I am constantly afraid of dealing with these challenging situations because I am alone. Fear, anxiety, sadness—I have lost 10 kilos, insomnia, I take Xanax every day, I cry in silence " (wp7).

Another participant disclosed, "A little bit to hear him move, I jump up because I have much anxiety and fear. I need to be well to serve him... I cannot escape this thing; I do not know why. It is now in my body... A neurologist has told my children that I am in a worse condition than their father (wp5). The internalized dependency and inadequacy, coupled with an unexpected new form of motherhood, force these caregivers to exhibit heightened empathy as a sense of sharing their husbands' vulnerability and hypervigilant concern.

In the accounts, alienation disrupts the participants' sense of belonging in the world as gendered subjects. Formerly skilled at enacting their gender identity, participants struggle to exert control over this conception. An altered sense of self hinges on agency, as the internalized norms of gender force them to adapt to an alien world.

Reparation: Reconstitution of Gender Identity

As the caregiving journey unfolds, the participants in the study justified their caregiving role by (re)constituting themselves as gendered agents. This process was a work of reason involving consciously using the gender norm demands.

| Men participants actively reconstruct their masculinity, aligning it with values |
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| such as being a law-abiding citizen and a protector who embodies decency, reliability, |
| virtue, and respectability. Upholding honor necessitates sacrifices and self-denial, |
| fortified by courage and strength. By distinguishing themselves from less honorable |
| men, these participants emphasize the moral significance of their caregiving acts and |
| are willing to execute them. One participant expressed this sentiment: "She is my wife, |
| and I hold an obligation to her and society. I cannot leave my spouse alone and |
| helpless at home while I go out. I am a man, and there are good men and bad men |
| who take a divorce" (mp1). Another participant added, "I do my duty as I should. I |
| have positive feelings. I understand what I need to do. Okay, I like it (mp2). |
| The public acknowledgement of their heroism, encompassing traditionally |
| masculine attributes like strength, bravery, perseverance, and commitment to |
| exceeding expectations, serves to solidify their gender identity as unquestionable. A |
| participant proudly stated: "Everyone here wonders how a man can do all these |
| things. They call me a hero, and others call me a rock!" (mp3), highlighting how |
| participants actively redefine their masculinity within the caregiving role. |
| In contrast, women caregivers navigate their reparative journey by embracing |
| caregiving with empathy, benevolence, and an understanding of morality as a form of |
| self-sacrifice. One participant compassionately expressed: "I feel sorrow for my |
| husband now because he was an active man, and he is now plagued by sadness I |
| prioritize his well-being over my own I have willingly relegated myself to a |
| secondary position." (wp4). She continued, "Since I was 14, I have been raising my |
| siblings and children alone People often ask me: How do you endure? My response |
| is simple: What choice do I have? I have been doing this for years. It is like having a |

baby. I felt inner strength. I felt empowered. I do not perceive it as a burden; I undertake it willingly" (wp4).

However, this perception of the feminine identity, defined by caregiving and nurturing, while empowering, also restricts women caregivers from asserting their self-determination within the caregiving role, resulting in an implicit sense of guilt and a denial of their subjectivity. In a reflective moment, one participant stated, "Okay, I might experience moments of stress. At times, I may wonder why shall I stay here? But I do not take it seriously" (wp8). Another participant contemplated her choices: "I know a woman who did not provide care; she maintained her career and had others attend to her husband. In contrast, I did not work. I had nothing. I left myself in ruins." (wp7).

However, it is crucial to recognize that the apparent self-sacrifice, seemingly devoid of personal gain, conceals a more nuanced survival strategy employed by women caregivers. A participant's statement vividly illustrates this complexity: "I believe that one must be willing to sacrifice oneself to aid someone in such dire circumstances. I needed him to stay alive, even paralyzed. My longing was intense. I felt a profound need to be with him " (wp10).

Reparation involves a deliberate engagement in normative reflections and practices to restore their gender identity while justifying their existence in the caregiving role. In this phase, individuals recognize the need to align their agency with their gender self-concept and utilize available resources to infuse their caregiving actions with gender significance, thereby mending their identity. Critical moral self-conscious emotions, including guilt and pride, are pivotal in guiding their actions. Meanwhile, self-sacrificial acts contribute to a heightened sense of worth associated with their gendered identity.

In visualizing the phenomenon, it becomes evident that normative gender roles are deeply embedded within the spousal caregiving experience. These roles exert influence through anticipated behaviors, ideals, and expectations, shaping how individuals perceive themselves and their roles within the caregiving journey. As the participants contend with their evolving self-concepts, this interplay between societal norms and personal identity leads to alienation. The pervasive influence of societal gender norms is experienced as a constraining force, limiting individual agency and shaping their experiences. Reparation, marked by normative reflections and self-sacrifice, becomes a mechanism for restoring agency and identity alignment. This phase reveals not only the process of identity repair but also sheds light on the complex psychological structure of the phenomenon. This reparation process signifies the malleability and resilience of gender identity, highlighting how participants negotiate their sense of self within the broader context of caregiving and societal norms (see supplementary 5 for an illustration).

Discussion and Implications

The overarching aim of this study was to explore how spousal caregivers in the Greek community experience gender while caring for older individuals in rehabilitation. In the phenomenological analysis of the eleven participants' transcripts, the experience of gender emerged as a normative diachronic identity with distinct phases: normative constitution, alienation, and reparation.

In phase one, normative constitution, it is seen how the participants' experience of gender develops within a culture implicitly permeated by heteronormative principles. The participants' mental representations of gender consisted of two distinct gender categories, women and men, encompassing specific evaluative criteria of being that formed the archetypes and ideal members of each of

the gender categories and against which individuals were evaluated as better or worse examples of the category (Hampton & Reimer, 2015; Rosch, 1975). For the participants in the study, the perception of two separate and opposing genders was associated with the 'natural' roles that match their assigned sex, making sexual orientation essential to their conceptualization of gender.

The findings follow research showing gender and sexuality to be inextricably tied together and inseparable constructs in the mind of the everyday perceiver, supporting that the general categories of 'women' and 'men' often assume heterosexuality emphasizing this constraint on generalizability for researchers and practitioners when employing gender categories of women and men in future research, interventions, or communication with caregivers y (Henry & Steiger, 2022; Kitzinger, 2010; Klysing, 2023).

The stereotype content associated with women and men in this study follows a complementary structure found in various cultural contexts where women are stereotyped as high in relatedness and interdependence but low in agency, while men are stereotyped as low in relatedness but high in independence and agency (Ellemers, 2018; Guimond et al., 2006). The level of internalization of stereotypical gender characteristics affected the fluidity of the individual's self-concept and sense of agency, shaping independent and relational selves (Cuddy et al., 2009; Guimond et al., 2006). As in this study, the literature shows the relational sense of self as central to women's identity, with women's agency to be manifested as a relational and collective phenomenon rather than an individual (Charrad, 2010; Gallagher, 2007).

The findings emphasize the significance of understanding how internalized gender stereotypes and the intersection of gender, sexuality, age, division of labour, power dynamics, cultural norms, and values shape caregiving behaviours and

relationships. Recognizing these factors can help explain the stressors associated with the caregiving experience and the strategies employed to cope with the stressors (Calasanti et al., 2021; Onorato & Turner, 2004).

In phase two, the study reveals the experience of alienation among caregivers with the emergence of the caregiving journey. There is a lack of research on alienation in informal caregiving, with comparative studies exploring variants of alienation such as loneliness, social isolation, and powerlessness (Seeman, 1959). The findings support research in older adults, suggesting that subjective loneliness is related to social isolation (Wenger & Burholt, 2004). Although women and men experienced social isolation in terms of an objective decline in social interactions, the subjective sense of dwindling social connectedness and rejection was profound among men linked to changes in their masculine identity and engagement in feminized caregiving acts, leading men carers to strive to maintain masculinity avoiding seeking support, further exacerbating their social isolation (Milligan & Morbey, 2016).

Primary prevention of loneliness is necessary to preserve social networks and promote resilience among older carers, acknowledging further men's vulnerability who may struggle with evolving perceptions of masculinity and recognizing that an environment that respects their autonomy may be necessary to engage some men in psychological support (Willis et al., 2020). Respite services, daycare, institutional care services or the assignment of a case manager as a nurse to the caregiver and care recipient dyad may benefit the older caregivers with a temporary break from caregiving duties to engage in desired activities (Schulz et al., 2020a). Findings emphasize the importance of recognizing diverse expressions of masculinity in caregiving, the range of emotions, and caregivers' psychological vulnerability, regardless of gender. (Campbell & Carroll, 2007; Giesbrecht et al., 2017).

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For women caregivers, alienation involves a loss of happiness in the performance of traditional gender roles in marriage and motherhood (Ahmed, 2020; Suppes, 2020). Consistent with the literature on gender differences in psychological morbidity among caregivers, women reported poorer mental health than men in terms of anxiety (Pillemer et al., 2018; Yee & Schulz, 2000). A heightened affective empathy may partially explain women's psychological vulnerability, possibly influenced by gender stereotypes (Zahn-Waxler and Van Hulle, 2012; Michalska, Kinzler and Decety, 2013). Although affective empathy, in contrast to cognitive empathy, is connected to emotional distress, anxiety and depression, few studies have examined the connection between both facets of empathy and mental health outcomes in caregivers (Tone & Tully, 2014). The findings point to affective empathy as a therapeutic target for caregivers with anxiety and depression symptoms, considering risk factors such as spousal caregiving, age and gender and the need to explore if supporting carers to regulate their emotions by maintaining a clear distinction between the self and the other would be beneficial for their well-being (Hua et al., 2021). Meditative intervention strategies may be beneficial as they encourage reflection on what is and is not achievable in helping a loved one and also providing respite to ease care provision and treatments to decrease the suffering of the care recipient (Collins & Kishita, 2019; Schulz et al., 2020). Moreover, understanding caregiver distress considering factors such as care recipients' disabilities is essential to tailor support and interventions for caregivers. For example, in stroke caregiving, the demanding nature of assisting in rehabilitation and the hope of recovery may intensify stress levels, whereas in dementia caregiving, progressive cognitive decline may induce more prolonged and chronic stress (Schulz et al., 2016).

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Heightened empathy was also connected to hypervigilant monitoring observed in other studies for women carers and discussed as a strategy to maintain control in unfamiliar situations (Green & King, 2009). It can be argued that this hypervigilant monitoring is associated with a form of intensive mothering driven by internalized prejudices and the 'good mother' stereotype consisting of un-reflected guilt and maladaptive reparation efforts that alleviate distress and depression symptoms (Liss et al., 2013; O'Connor et al., 2007). Men caregivers also experienced guilt characterized by emotional ambivalence, absence of self-disclosure, loneliness and alienation (Bruno et al., 2009). Literature shows that caregivers experience guilt for various reasons: actions, limitations, negative emotions, relationship changes, and for neglecting other areas, connecting guilt with emotional distress (Gallego-Alberto et al., 2022). Research also suggests that women feel guiltier for leaving dependents alone or neglecting other areas, while men feel guilty for not performing domestic tasks and losing patience (Brea et al., 2016). In this study, caregivers' guilt had two components: interpersonal guilt, arising from caregiving motivations, prompting reparatory acts of attentiveness to the care recipient's needs, and intrapsychic guilt that appeared as a dysphoric feeling associated with personal distress and a fear of transgressing moral standards related specifically to gender norm violation (Carnì et al., 2013). These findings suggest examining strategies for managing the two types of guilt and understanding their differences to inform therapy. Exploring experiences to identify vulnerability factors that contribute to each type of guilt could also enhance understanding of psychological processes and guide targeted interventions (Mancini & Gangemi, 2021).

In phase three, reparation, the findings highlight the influence of gender norms on participants' moral motivation for caregiving, showing moral motivation to result

from the interaction between individual levels of gender identification and the content of shared gender stereotypes (Nunner-Winkler et al., 2007). Women's care orientation predisposes them to adopt a moral of self-sacrifice led by internalized selflessness (Shabot, 2022). Men's justice orientation, by encompassing excess altruism, also involves a moral of self-sacrifice that is often perceived as heroic and commendable, allowing them to reform their masculinity without displacing their hegemonic vision (Campbell & Carroll, 2007; Connell & Messerschmidt, 2005).

It is argued that self-sacrificial acts are praiseworthy, though non-obligatory, in that their omission is not blameworthy (Urmson, 1958). However, this study questions the extent of free will in these acts for carers, as they seem to be influenced by internal and external factors and driven by societal expectations. Compliance with these expectations allowed individuals to affirm their gendered sense of self and self-worth, as deviating from them bred self-doubt, guilt, and alienation, showing that the self-sacrificial caregiving acts were not only pursued solely for the sake of the other but for the sake of avoiding negative consequences for the self. Self-sacrificial acts gave the caregivers a tremendous opportunity for worth gain in their harmed gender identity more than other neutral or pleasurable acts could give (Dugas et al., 2016). Research on daily sacrifices in intimate romantic relationships shows that when the cost is high, or sacrifices are driven by avoidance motivation, they are harmful to the well-being of both partners and determinantal for relationship maintenance (Day & Impett, 2017; Impett et al., 2013).

The findings highlight the need for therapeutic and preventive measures to address self-sacrificial morals in caregiving. Differentiating between commitment in a caregiving relationship and self-sacrifice, as well as altruism, is essential. Goal commitment involves persistence and effort, self-sacrifice involves focusing on a

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cause and neglecting alternative goals, whereas altruism entails acting for others without personal gain and does not involve necessary significant loss (Bélanger et al., 2018).

Feelings of guilt and lack of perceived choice in the caregiving role are connected with increased burden and psychological morbidity among caregivers (Schulz et al., 2016). Numerous research highlight the influence of social and cultural factors on caregiver motives and choices (Zarzycki et al., 2022). It has been suggested that social norms and expectations impose a sense of obligation on individuals, which is thought to be the primary caregiving motive (Corey & McCurry, 2017). The current findings add to the literature by addressing the internal, individual, context and gender-based caregiving experience whilst also considering moral and ethical aspects of caregiving, showing how culture and society-dependent factors provide context to psychological factors that shape the perceived obligation to provide care. Understanding caregivers' motivational approach and assessing emotional impact can provide insights into their level of joy and pleasure, determining the genuineness of caregiving acts. Interventions may support caregivers' assertiveness skills to effectively express their needs and desires, communicate boundaries and preferences, and engage in activities that bring them fulfilment. There is a need to support caregivers to think and challenge beliefs that reinforce the necessity of self-sacrifice either for their worth or the happiness of others, empowering them to prioritize their well-being for sustaining their motivations in caregiving. This shift in perspective can lead to a more positive and fulfilling caregiving experience.

Limitations

This study aimed to provide an in-depth analysis of the experiences of a specific subgroup of caregivers; therefore, the purposive sample strategy inherently

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limits the generalizability of the findings to the under-study population (Palinkas et al., 2015). Greece's sociocultural context, unique historical factors, and specific gender norms shaped participants' experiences. Notably, all participants identified as cisgender and heterosexual, further restricting generalizability to other gender identities and sexual orientations. The sample primarily represented a working-class demographic from one urban clinic, limiting applicability to diverse socioeconomic backgrounds, rural settings, varied clinical contexts, and caregiving dynamics. Nevertheless, this study provides a framework for examination in different populations. Future research should expand on these findings to understand gender stereotypes, caregiving experiences, and moral decision-making across a broader spectrum of caregivers, encompassing diverse cultural, ethnic, and socioeconomic backgrounds, gender and sexual orientations. Qualitative research findings, reliant on subjective analysis and narrative descriptions, may be subject to multiple interpretations influenced by researcher biases, participant responses, and translation issues. Informants may provide unreliable data due to a desire to please or hidden intentions, emotions, principles, or viewpoints (Tongco, 2007). Our research team maintained rigor through reflexivity and transparent data collection, methods, and analysis to mitigate these limitations.

Conclusion

The study offers a framework for examining the caregiving experience and psychological outcomes for spousal caregivers from a gender perspective. It highlights the influence of stereotypical attributes linked to femininity and masculinity on self-concept, agency, and the formulation of caregiving evaluations and practices, highlighting the importance of incorporating an intersectional perspective in future research and interventions with caregivers, considering factors

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such as gender, sexuality, gender relations, age, division of labour, power dynamics, cultural norms, and values. The results indicate that caregivers feel alienated upon embarking on the caregiving journey, with specific gender-based vulnerabilities that impact their psychological well-being. By recognizing these vulnerabilities and addressing gendered expectations and societal pressures, interventions can foster caregivers' mental health and facilitate the development of effective coping strategies. The findings shed light on the complexity of empathy, the multifaceted nature of guilt, and their connection to the caregivers' gender self-concept, emphasizing the central role of emotions in the experience of stressors guiding caregivers' thoughts and behaviours. The research highlights the moral dimension of caregiving and its relationship with gender norms and motivations, calling for challenging selfsacrificial morals and societal norms associated with them to empower caregivers to prioritize their well-being while sustaining their motivations in caregiving. By recognizing gender-based vulnerabilities in the spousal caregiving journey, rehabilitation psychologists can create a supportive environment for caregivers, fostering a more positive, fulfilling and rewarding caregiving experience. References Ahmed, S. (2020). The Promise of Happiness. *The Promise of Happiness*. https://doi.org/10.1515/9780822392781 Bélanger, J. J., Schumpe, B. M. M., Menon, B., Ng, J. C., & Nociti, N. (2018). Self-Sacrifice for a Cause: A Review and an Integrative Model. In *The SAGE* Handbook of Personality and Individual Differences: Volume II: Origins of Personality and Individual Differences (pp. 465–485). https://doi.org/10.4135/9781526451200.n25

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