# **Impact of COVID-19 on Cancer Caregivers:** Results from the Global Carer Well-being Index

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# **HIGHLIGHTS:** Cancer caregivers were more likely to report...



# 71 vs 63%

..that COVID has made caregiving harder than caregivers in other groups (e.g.; dementia, Parkinson's disease)

67 vs 61%

.that COVID has worsened their emotional/mental health than caregivers in other groups (e.g.; spinal cord injury, congestive heart failure)



# 61 vs 56%

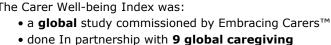
. a positive impact on the relationship with the individual they are caring for because of the pandemic



26 vs 13%

.becoming a caregiver in the last year than caregivers in other groups (e.g.; muscular dystrophy, multiple sclerosis)

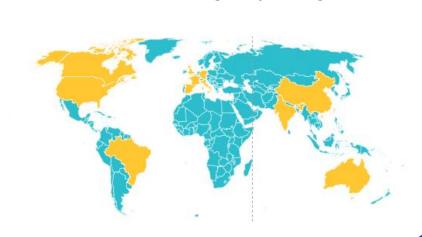




organizations

BACKGROUND

- conducted in 12 countries (United States, Canada, Brazil, United Kingdom, France, Germany, Italy, Spain,
- designed to explore the **impact of the pandemic** on the health and well-being of unpaid caregivers







- The survey was conducted:
  - via online and phone methodologies
  - from September 3 October 27, 2020
  - with 9,000 unpaid (informal) caregivers caring for someone
    - Long term illness (e.g.; cancer)
    - Physical disability (e.g.; spinal cord injury)
    - Cognitive mental condition (e.g.; Alzheimer's)
- Sample size for **cancer caregivers** was **N=1035**, including 109 from USA and 102 from Canada
- · Statistically significant differences between cancer carers and all other carers were evaluated using the Chi-square test with  $p \le 0.05$



- 63% of all caregivers were women, with a **60/40 female to male** split in the cancer caregiver group.
- Average age of starting caregiving was 39.0 in cancer caregivers, and 37.2 in non-cancer caregivers.
- Cancer caregivers were more likely to be in the 60+ age group (15 vs 11%) than non-cancer caregivers.



### **RISING DEMANDS**

## Juggling responsibilities:

29% report more difficulty balancing work with their care duties

# Increased dependence:

42% say person they care for is relying on them more than ever before; which is higher than the average across all caregivers surveyed (39%)

### Time increase:

Pre-pandemic, spent an average of 16+ hrs/week providing care; 34% expect an increase to 30+ hrs/week, even after the



#### **CHANGED RESPONSIBILIITES**

# **Increased technology use:**

- 53% have increased responsibility managing medical appointments virtually
- 63% need additional training on telehealth and other online tools

#### **Emotional Support:**

- 58% are providing more emotional support than
- 91% state they are putting the needs of those they care for above their own

# Work / care balance:

58% feel that being an unpaid caregiver has negatively impacted their paid work responsibilitie



# **TOLL ON CAREGIVERS**

#### **Emotional health:**

- 67% say pandemic has worsened emotional health
- 28% feel they have no-one to turn to for support

# Physical health:

- 50% say the pandemic has worsened physical health
- 61% report lack of sleep
- 77% report unprecedented levels of burnout

### **Financial health:**

58% say the pandemic has worsened their financial status - household and medical bills, further impacted by reduced working hours and a lack of financial support



# **INEQUITIES IMPACING** CAREGIVERS

### Public and private sector support:

- 73% have never received support from local government, which is higher than other caregivers (66%)
- 84% never from private companies, 77% never from non-profit organizations

### Understanding benefits:

92% report that help navigating the health and social services / care system is important to ensure they can provide the care needed.

# **Employer support:**

- 96% feel that employers should provide additional flexibility policies and support for caregivers
- 63% have never received support from their employer





#### Address Emotional / mental well-being:

Provide more respite services that are accessible, flexible and easy to secure, especially for unpaid caregivers of people with cancer who have complicated requirements and very specific needs



# **Provide training and support:**

Provide supportive training on access and effective use of telehealth and other online resources, especially for those managing medical appointments and ongoing treatment for those with cancer



# Reduce inequalities:

Incentivize employers and educational environments to develop support programs for employees and students who support people living with cancer (e.g.; flexible working/study hours, paid medical leave)



There are ways HCPs and their team can help recognize and support cancer caregivers:

CARE PROVIDER

**ROLE** of the HEALTH

- 1. Formally recognize family caregivers as part of the patients inter-disciplinary healthcare team.
  - Does the caregiver have the information they need to support their loved one?
- 2. Inquire about the family caregiver's health and **well-being** and if possible access if changes over
  - Are they having trouble sleeping, issues with balancing work and caring? Do they need to talk to someone?
- 3. Have resource materials specific to family caregivers unique needs.
  - Are their local support groups, respite care resources, etc. that you can refer a family to?

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