Acceptance, grief and meaning

Prigerson & Maciejewski\(^1\) assert that the resolution of grief coincides with increasing acceptance of loss, mainly cognitive and emotional acceptance. The role of spiritual acceptance has not been mentioned directly, although experiences like inner peace, tranquillity and letting go, or regaining what is lost or being taken away, are more spiritual rather than emotional or intellectual. Moreover, some of the features which can be considered spiritual are included as criteria for prolonged grief disorder,\(^2\) such as confusion about one’s identity and feeling that life is empty and meaningless since the loss. Issues related to culture and the meaning and value of death\(^3\) are relevant to both grief and acceptance, and I wonder whether these should also be considered.

Patients diagnosed with terminal cancer often confront existential issues. Experiences with patients with advanced or terminal cancers indicate that not only is cognitive and emotional acceptance important, but that spiritual aspects are equally relevant. Spiritual acceptance of grief will help the grieved to understand the meaning and purpose of the loss. As Frankl\(^4\) states ‘suffering ceases to be a suffering as soon as it finds a meaning’. Longitudinal studies should clarify not only the way in which grief’s suffering ceases to be a suffering as soon as it finds a meaning. It may heighten an individual’s sense of anomie (i.e. a feeling of disorientation and alienation from society caused by the perceived absence of a supporting social or moral framework) and affect a person’s will to live. The meaning derived from spiritual beliefs may buffer individuals from the emptiness that follows a major interpersonal loss. Still, we do not consider spiritual beliefs to be components of grief. Rather spirituality may be a powerful antidote (perhaps, social support and social integration are others) to the pain of grief and elixir promoting emotional acceptance.

There is some small change besides the two sides of the coin:


Authors’ reply: We thank Dr Chaturvedi for highlighting the potentially important role that spirituality plays in the acceptance of loss. Recent research attests to the powerful influence of spirituality and religious beliefs in shaping patients’ cognitive acceptance of terminal illness, treatment preferences, and even in determining the receipt of intensive, life-prolonging care in the last week of life.\(^3\)

Nevertheless, we wish to differentiate between components of grief (e.g. yearning) and factors affecting the intensity and course of grief (e.g. spirituality). We posit that grief is on the same continuum as emotional acceptance – opposite poles of a unitary dimension. We contend that both spirituality and cognitive acceptance are distinct from, but related to, emotional acceptance and grief. Spirituality might foster emotional acceptance; cognitive acceptance might exacerbate grief. Identifying factors affecting grief and emotional acceptance may suggest ways to enhance an individual’s mental health and well-being in the face of death, and offer ways to minimise loss-related suffering.

As a further distinction, we consider the loss of meaning in the context of prolonged grief disorder\(^2\) to represent the emptiness experienced by the absence of an attachment figure. It is not intended to refer to a broader existential crisis. The sense of emptiness felt in grief may well lead a person to question the meaning of life. It may heighten an individual’s sense of anomie (i.e. a feeling of disorientation and alienation from society caused by the perceived absence of a supporting social or moral framework) and affect a person’s will to live. The meaning derived from spiritual beliefs may buffer individuals from the emptiness that follows a major interpersonal loss. Still, we do not consider spiritual beliefs to be components of grief. Rather spirituality may be a powerful antidote (perhaps, social support and social integration are others) to the pain of grief and elixir promoting emotional acceptance.

3. Holly G. Prigerson, Paul K. Maciejewski Brigham and Women’s Hospital, Harvard Medical School, and Statistics Core of the Center for Psycho-oncology & Palliative Care Research, Dana-Farber Cancer Institute, 44 Binney Street, Smith 268, Boston, MA 02115, USA. Email: Holly.Prigerson@dfci.harvard.edu doi: 10.1192/bjp.194.6.561

Taking an internet history

Cooney & Morris\(^1\) argue that we should consider taking an ‘internet history’ to help assess young people’s risk of self-harm, suicide and presumed psychopathologies such as ‘internet addiction’. Although an understanding of how a client uses the internet may be important, the authors caricature what we know about the risks of the internet.

Although information on suicide methods is available online,\(^2\) there is currently no clear evidence that the risk of self-harm or...
suicidal behaviour is raised by ‘pro-suicide’ internet sites, as we lack all but the most preliminary studies in this area. Those studies that have been completed, in line with earlier research on ‘pro-anorexia’ sites, reported that ‘pro-self-injury’ boards relay mixed messages – clearly providing social support, coping methods and understanding, but also tending to minimise the significance of self-harming behaviour.\(^3\) On the basis of current evidence, we might hypothesise that the use of such websites could equally be a protective factor or a risk factor.

The authors also mention internet addiction but seem unaware that the existing research is based on inconsistent criteria, is subject to widespread sample bias, relies almost entirely on correlative studies,\(^4\) and that the concept itself lacks conceptual validity.\(^5\) I challenge the authors to find any empirical studies to support their claim that in Asia ‘cardiopulmonary-related deaths and even game-related murders in internet cafes are now regarded as serious public health issues’.

I wholeheartedly support the authors’ contention that clinicians should consider the role of the internet in the lives of patients, but I would stress that this needs to be done with an understanding of the relevant research literature and a working knowledge of both the technology and culture of the medium.

We ask no less in other areas of clinical work and this is particularly important in a time when fears about the internet are amplified by the media with little regard to the evidence, particularly important in a time when fears about the internet.

We welcome Dr Bell’s interest in our letter and his agreement with him. He makes some crucial points which we too would emphasise. In particular, we all share the ‘contention that clinicians should consider the role of the internet in the lives of patients, but I would stress that this needs to be done with an understanding of the relevant research literature and a working knowledge of both the technology and culture of the medium.’ Sadly, there is too little sound evidence to inform our attitudes. It has been as a result of some distressing clinical experiences, as well as concerns raised sensation rather than scientifically in the media, that we have been moved to highlight the issue and to embark on our own preliminary studies.

None of this is to re-ignite a debate on whether the internet is helpful or harmful. As Dr Bell has observed, such a reductionist approach belies the complexity and variety of internet-based activities, any of which may have an influence in either direction.\(^1\) We instead reflect that without empirical data to inform us, and where there is the possibility of either risk or benefit, careful and sensitive questioning of patients with high internet use may be a valuable component of a full psychiatric assessment.

The internet has taken a central place in modern culture particularly among younger people. Although we may not fully understand the complex interactions of the web and mental health, and while we await research to enlighten us, we are left with the choice to either ignore or engage with this phenomenon. Legislators, mental health advocates,\(^7\) concerned parents and media journalists have all focused their efforts. It is time for scientists and clinicians to follow suit. In our view, this begins with the careful taking of an internet history.

---


---

Dementia: suicide by drowning

Purandare et al’s article on suicide in dementia is a valuable contribution to suicide research in the elderly, particularly in those with dementia.\(^1\) The authors have already dealt with a number of methodological limitations quite succinctly. One important limitation in particular is the choice of controls. As the authors rightly stated, a control group of patients with dementia who had not died by suicide would have been more appropriate.

In the Method section, the authors referred to ICD–10 only and not ICD–9. As far as I am aware from my own experience dealing with the Office for National Statistics (ONS), ICD–10 has been used by ONS only since 2001. Prior to this date and for the first 5 years of Purandare et al’s study period (1996–2000), the ONS used ICD–9. If the authors applied the same criteria in their selection of suicide and open verdicts in cases reported between April 1996 and December 2000, then I assume they would have selected: ICD–9 E950–E959 for suicide and E980–989 excluding E988.8 for open verdicts respectively in a similar manner as they did with ICD–10 (p. 175). However, this very relevant fact does not appear to have been mentioned or explained by the authors, and was quite possibly omitted from the manuscript in error. However, this omission, which covers 5 years of a 9-year study, ought to be acknowledged and duly corrected.

I am grateful that the paper provides the opportunity to make one or two comments on some issues relating to drowning as a method of suicide in the elderly. Suicide by drowning accounted for 13.5% of total elderly suicide, being the third commonest cause of death in elderly suicide in England and Wales during 1979–2001 (16% for women as the second commonest cause of death in elderly suicide in England and Wales during 1979–2001) (16% for women as the second commonest cause of death in elderly suicide in England and Wales during 1979–2001)