Modelling complex pathways between late-life depression and disability: evidence for mediating and moderating factors

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Previous research has consistently shown an association between depression and disability in the elderly but little is known about the mechanisms linking the two. Recent longitudinal population studies have shown considerable inconsistency in the criteria used to establish causality and terms such as mediation and effect modification have been frequently applied incorrectly in terms of the inferences drawn. We underline the necessity to adopt more stringent theoretical criteria for the establishment of intermediary effects in the relationship between depression and disability to better identify cross-validated potential intervention points for reducing the risk of disablement and depression.

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Introduction

Although a large body of literature confirms the high co-occurrence of depression and disability in the elderly (Lenze et al. 2001), much less is known about the mechanisms linking the two. The increasing number of longitudinal observations now available makes the exploration of temporal pathways possible; however, such studies have also identified several intervening factors that potentially modulate the relationship between the two, making health outcomes following either depression or disability difficult to predict or prevent. Failure to take into account the innate complexity of the cause–effect relationships involved has led to confusion in our present understanding of the relationship between depression and disability. The conceptual framework (if any) used by most studies to establish mediating effects is seldom discussed in publications, thus leading to considerable confusion in terminology and the use of inappropriate statistical methods.

Theoretical criteria for the establishment of mediation and moderation

Depression, like many other mental disorders, cannot be attributable to a single cause but is increasingly recognized to lie along multiple causal chains involving environmental, social and biological risk factors that mediate the relationship with activity limitations. Causal chains are often complex in the case of psychiatric disorders as the pathology may both cause and be the result of the same mediating factors such as social, biological disturbances or at-risk behaviours. Kraemer et al. (2001, 2008) have proposed a theoretical model of this type of complex causal chain that may be used to validate causality and risk factor mediation within longitudinal data sets. In this model a mediator or intermediate variable is defined as a variable that occurs in a causal pathway from an independent to a dependent variable. It ‘causes’ variation in the dependent variable (Y), and itself is subject to variation by the independent variable (X). The independent variable precedes the mediator (Me), it is correlated to Me, and when X and Me are jointly analysed either Me ‘dominates’ X (total mediation) or X and Me co-dominate (partial mediation). Proving that X precedes Me is thus a key point in demonstrating mediation. A moderator or effect modifier refers to variations in the magnitude of the effect of the independent variable across levels of another variable. Mo moderates the association between X and Y, if Mo precedes X, Mo and X are not correlated and Mo and X co-dominate. Mo identifies subpopulations in which different causal chains operate or one causal chain operates differently (Kraemer et al. 2001). Moderators may thus be considered in relation to susceptibility or resiliency or...
buffering factors. A mediator is in a causal sequence between two variables whereas a moderator is not part of this causal sequence (Fig. 1).

Establishment of mediating and moderating effects
Of the 24 longitudinal population studies that have been published to our knowledge since 2001 examining the association between depression or depressive symptomatology and activity limitation, 11 examined depressive symptomatology as a risk factor for activity limitation, 11 examined activity limitations as a risk factor for depressive symptomatology and two examined the reciprocal relationship. We found considerable inconsistency in the criteria used to establish causality and terms such as mediation and effect modification were frequently incorrectly applied in terms of the inferences drawn. Adjusting for a covariable measured at baseline and observing a decrease in the strength of the association between the main factor and the outcome does not prove that this covariable represents a mediating effect. Mediating effects should be defined as new events occurring during follow-up or as changes since baseline, and associations should be established by appropriate statistical models such as linear mixed models, generalized mixed models or multi-state models applied to longitudinal data. However, it should be noted that the use of linear mixed models may be questionable as the scales used to measure depression or disability are seldom normally distributed in the general population. We also observed that bias due to cohort attrition has rarely been taken into account in research in this area. Depression and disability may induce early drop-outs and sensitivity analyses need to be undertaken to test the robustness of the results.

To illustrate the problem of defining effect mediators let us consider concomitant illnesses. Depression in late life is frequently influenced by other conditions present at baseline: depression is common in older persons recovering from myocardial infarction or stroke and in those suffering from cancer, diabetes, hip fracture and cognitive impairment (Blazer, 2003). Depression may also induce physical illness including cardio- and cerebrovascular disorders. Depressed individuals may have detrimental health habits (poor nutrition, smoking, alcohol, lack of exercise) that are risk factors for vascular illness. Immune dysfunction associated with depression may also be associated with increased risk of other pathologies such as cancer (Lenze et al. 2001). To demonstrate that diseases are mediating effects in the pathway leading from depression to disability, only new or worsening diseases occurring after depression onset and before disability should be considered. Pre-existing depression risk factors cannot be considered as mediators but rather as confounding factors. Such a distinction is important to explain the mechanism leading to disability and to find targets for intervention to prevent or postpone disability in depressed elderly persons.

Mediating and effect modification between depression and activity limitations
Several potential factors have been proposed by previous studies but all require more stringent evaluation of their mediating effect: namely, loss of motivation leading to self-neglect, decrease in physical activity, and poor nutrition inducing sarcopaenia. Depression is also observed to induce lack of compliance with medical treatment (Bruce, 2001), isolation, and difficulties in coping with life events, which in turn determine depression. Conversely, beneficial intermediate effects are observed due to psychiatric care, antidepressant prescription, improvement of social support and correction of sensorial impairment, all of which may break the ongoing chain to disability.

Several effect modifiers were found to be significant: pre-existing chronic disease (Geerlings et al. 2001), time (Lenze et al. 2005), cognition (Mehta et al. 2002) and baseline functional limitations (van Gool et al. 2005). Other factors have to be considered: gender, education, genetic and environmental factors, and history of pathologies. In particular, women are known to have more depressive symptoms than men but seem to be less liable to depressive symptoms when suffering from physical health problems (Piccinelli & Wilkinson, 2000). A high level of education is also associated with reduced rates of depression, but loss of functional capacities may be less accepted by educated people. However, people with a high income may more easily access help and home adaptations, thus decreasing the consequences of functional impairment. A history of depression may also be an effect modifier, rendering people less able to cope with loss of functional activity and, conversely, the impact on disability of chronic depression being possibly stronger.

![Mediation and effect modification between depression and activity limitations](https://doi.org/10.1017/S0033291709005273)
Mediating and effect modification between activity limitations and depression

Certain factors, namely availability of a confidant, satisfaction with support, sense of control and self-esteem, have been validated as mediators between activity limitations and depression incidence using appropriate analytic techniques (Yang, 2006). Dependency in daily tasks is a persistent chronic stressor leading to negative psychological outcomes, such as loss of self-esteem, that may also be symptoms of depression. Disability may also lead to restriction in social and leisure activities, and isolation, which are important precipitants of depression. Anticipated support may be the key element for buffering the effects of this stressor and perceived support has in some cases been demonstrated to have a more significant mediating effect as opposed to objective support (Yang, 2006). With ageing and increased risk of multiple chronic disorders, the role of the caregiver may become more onerous and time-consuming, creating tension between ageing parents, adult children and spouses, with possible role inversion. Other potential mediators require further research; in particular, physical rehabilitation was anticipated to change the course of both disability and depression (Bruce, 2001).

With regard to effect modifiers and possible stratifications, time or age (Schnittker, 2005; Anstey et al. 2007) and gender, race and socio-economic status (Schieman & Plickert, 2007) were found to be significant. However, most of the effect modifiers suggested above for the reverse association should be considered, providing that they precede activity limitation onset.

Conclusion

Recent longitudinal studies have identified a large number of factors that may impinge on the causal relationship between depression and disability. These mediating and moderating factors may vary according to the direction of the relationship being examined. Given the persistent variability in measures used for both depression and disability, it is not currently possible to validate findings from one study by other observations or to conduct meta-analyses. We underline the necessity for future studies to adopt more stringent theoretical criteria for the establishment of intermediary effects in the relationship between depression and disability, to adopt a common language with regard to mediating and moderating effects and to work towards uniformity in case identification to better identify cross-validated potential intervention points for reducing the risk of disablablement and depression.

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Declaration of Interest

None.

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