Bibliography and Suggested Reading:

Robin Morris CE Workshop.

The Wellbeing of Caregivers of People with Dementia Considered from a Neuropsychological Perspective.

International Neuropsychological Society Meeting, New York, 2019

Subitha Baghirathan S, Cheston R, Anndeloris C, Shears P & Currie K (2018). A grounded theory analysis of the experiences of carers for people living with dementia from three BAME communities: Balancing the need for support against fears of being diminished. <https://doi.org/10.1177/1471301218804714>

Clare L, Wu Y-T, Quinn C, Jones IR, Victor CR, Nelis SM, Martyr A, Litherland R, Pickett JA, Hindle JV, Jones RW, Knapp, M, Kopelman MD, Morris RG, Rusted JM, Thom JM, Lamont RA, Henderson C, Rippon I, Hillman A. & Matthews FE, on behalf of the IDEAL study team. (in press).  A comprehensive model of factors associated with capability to ‘live well’ for family caregivers of people living with mild-to-moderate dementia: findings from the IDEAL study. Alzheimer’s Disease and Associated Disorders*.* [Accepted October 2018]

Dickinson C, Dow J, Gibson G, Hayes L, Robalino S & Robinson L (2017). Psychosocial intervention for carers of people with dementia: What components are most effective and when? A systematic review of systematic reviews. International Psychogeriatrics, 29 (1), 31-43.

Feast A, Orrell M, Charlesworth G, Melunsky N, Poland F & Moniz-Cook (2018). Behavioural and psychological symptoms in dementia and the challenges for family carers: systematic review. British Journal of Psychiatry, 208, 429–434. doi: 10.1192/bjp.bp.114.153684

Kishita N, Hammond L, Dietrich M & Mioshi E (2018). Which interventions work for dementia family carers?: An updated systematic review of randomised controlled trials of carer interventions. International Psychogeriatrics, 30 (11), 1679-1696.

Kumfor K, Teo D, Miller L, Lah L, Mioshi E, Hodges JR, Piguet O and Irish M (2016). Examining the relationship between autobiographical memory impairment and carer burden in dementia syndrome. Journal of Alzheimer's Disease, 51 (1). pp. 237-248. ISSN 1387.

Morris R G, Morris LW & Britton PG (1988). Factors affecting the emotional wellbeing of the caregivers of dementia sufferers. British Journal of Psychiatry, 153, 147-156.

Perales J, Turró-Garriga O, Gascón-Bayarri J, Rene-Ramirez R & Conde-Sala J L (2016). The longitudinal association between a discrepancy measure of anosognosia in patients with dementia, caregiver burden and depression. Journal of Alzheimer’s Disease, 53, 1133-1143.

**Quinn C, Jones IR, Clare L** (In Press). Illness representations in caregivers of people with dementia. Aging Ment Health, 21(5), 553-561. <https://ore.exeter.ac.uk/repository/handle/10871/27297>

Uflaker A, Edmondson, MC, Onyike CU & Appleby BS. (2016). Caregiver burden in atypical dementias: comparing frontotemporal dementia, Creutzfeldt-Jakob disease, and Alzheimer’s disease, International Psychogeriatrics, 28 (2), 269-273.

Yu D, Cheng S-T & Wang J (2018). Unravelling positive aspects of caregiving in dementia: A integrative review of the research literature. International Journal of Nursing Studies., 79, 1-26.