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| **Title** | |
| Tan SB, Williams AF, Morris ME. Experiences of caregivers of people with Parkinson's disease in Singapore: a qualitative analysis. J Clin Nurs. 2012;21(15-16):2235-46. | |
| Agreement | 109/891 |
| **Agreed** | |
| *It’s quite hectic… he takes up all my rest day. I have to send him to his appointments or therapy. It’s tiring and I’m always busy. After we moved house three years ago, some of the cartons are still not opened yet (CG03) Qcarer*  This theme revealed the needs of caregivers for improved caregiver support. Caregivers wanted to have more information available about how to manage PD. Some caregivers suggested that the healthcare system could be improved if a more integrated approach to providing PD services were adopted. A caregiver stated:  *I keep saying we need better services and solutions. What we have now is not enough and it’s all over the place. Hospital could package services into one centre with one slot of time where you can do speech (speech therapy), physio (physiotherapy) and all that (CG13) Qcarer*  Some caregivers noted that transportation needs became more difficult as the care recipient’s disease progressed. | |