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| **Title** | |
| Dekawaty A, Malini H, Fernandes F. Family experiences as a caregiver for patients with Parkinson's disease: a qualitative study. Journal of Research in Nursing. 2019;24(5):317-27. | |
| Agreement | 78/402 |
| Final Agreement | 275/402 |
| **Agreed** | |
| All participants stated that they went to the hospital regularly (once a week or once a month). There is no type of service provided by a health worker other than checking the general situation and giving medicines to the patient. Annoyance was expressed by the third participant, who stated that she was upset due to always having to wait a long time for medication, and sometimes she felt hesitant to visit the hospital unless the patient had relapsed.  Psychologically, participants felt stressed and sometimes expressed their irritation at the patient’s attitude during treatment, as in the following:  *Well, I was just annoyed he did not want to follow the rules. He was prohibited to eat this and to do that, but he still ignored it. (P1) Qcarer*  The type of family support that caregivers received was related to finances, which helped the caregivers in buying medication and meeting personal expenses for both patients and carers.  For example:  *Yes, I have a child who also likes to send me money to buy medication and to pay for my expenses. (P4) Qcarer*  In addition to support from their family, participants also received moral and material support from neighbours:  *Sometimes, there are neighbours who ask about the condition. They provide us with food or money (P1) Qcarer*  *God also does not want to help us if we do not fight right. It is impossible for God to give us money immediately . . . let’s sell things, many neighbours buy them here. (P2) Qcarer*  *Whether it is enough or not, there is help from the people who rent a stall in front of the house.*  *Thanks to God. (P4) Qcarer*  *Because doctors say she should eat a lot of fruit for digestion, sometimes neighbours also give us*  *fruit. (P5) Qcarer* | |