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| **Title** | |
| Williamson C, Simpson J, Murray CD. Caregivers' experiences of caring for a husband with Parkinson's disease and psychotic symptoms. Soc Sci Med. 2008;67(4):583-9. | |
| Agreement | 119/818 |
| **Agreed** | |
| Caregivers often attempted to gain understanding and search for information themselves before approaching professionals involved in their partner’s care. They described feeling responsible for finding out about psychotic symptoms themselves because they did not have regular contact with professionals. In addition, some caregivers felt that professionals were not completely open about the complications of Parkinson’s disease and did not readily share information about the extent of the illness:  *One thing about Parkinson’s they don’t tell you anything.*  *I: Is that how it felt, that you weren’t told anything?*  *They don’t tell you, definitely do not. They don’t let you know that. It’s like it’s all hidden somehow. I don’t know. That’s why I didn’t know about the hallucinations*  *(P2). Qcarer* | |