

Annotation

The Bachelor thesis discusses the quality of life of carers of people with the Dementia Syndrom. The theoretical part deals with the description of the Dementia Syndrom, the personality of the carer, risks of the care and the theory of the life quality focussed on the carer.

The focus of the thesis is a research based on the quality analysis of the semistructured interviews. Ten carers of different age, sex and domicile performed as the examined sample.

The practical part is focussed on the attitude of a carer to the patient and on the problematics of the cultural life of the carers. Further, on the possibility of attending a job for carers and what knowledge about the health-social services do the carers have. And at last on the answer how are these services accesible. The verbatim transcriptions of the interviews with the particular respondents are included.

The conclusion is that the carers are exhausted due to the care, that they cannot attend their job and have shortage of the information about the social and especially the health services.