Does the risk of developing dementia increase with the number of episodes in patients with depressive disorder and in patients with bipolar disorder?

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Objective: Several findings suggest that some patients with depressive or bipolar disorder may be at increased risk of developing dementia. The present study aimed to investigate whether the risk of developing dementia increases with the number of affective episodes in patients with depressive disorder and in patients with bipolar disorder.

Methods: This was a case register study including all hospital admissions with primary affective disorder in Denmark during 1970–99. The effect of the number of prior episodes leading to admission on the rate of readmission with a diagnosis of dementia following the first discharge after 1985 was estimated. A total of 18,726 patients with depressive disorder and 4,248 patients with bipolar disorder were included in the study.

Results: The rate of a diagnosis of dementia on readmission was significantly related to the number of prior affective episodes leading to admission. On average, the rate of dementia tended to increase 13% with every episode leading to admission for patients with depressive disorder and 6% with every episode leading to admission for patients with bipolar disorder, when adjusted for differences in age and sex.

Conclusion: On average, the risk of dementia seems to increase with the number of episodes in depressive and bipolar affective disorders.

Although the literature is sparse, we hypothesised that the number of prior affective episodes (depressive or manic) is associated with increased risk of developing dementia. The aim of the present study was to investigate whether the number of prior admissions for patients hospitalised with depressive or bipolar disorder predicted increased risk of subsequently getting readmitted with a diagnosis of dementia using a national case register of all admissions to psychiatric wards.

Only patients who had been hospitalised at least once on a psychiatric ward and only affective episodes leading to admission were included in the study; these patients thus represent the more severely influenced proportion of patients. Data on treatment or outpatient status were not available.

METHODS

The register

In Denmark, all psychiatric admissions have been registered in a nationwide register for the 5.1 million inhabitants. All inhabitants of Denmark have a unique person identification number that can be logically checked for errors, so it can be established with great certainty if a patient has been admitted previously, irrespective of changes in name etc. Censoring due to death and causes of death can also be established with equal certainty as the same identification number is used across all public registration systems.

Between 1 April 1970 and 31 December 1993 the International Classification of Diseases (ICD), 8th revision was used in Denmark. For various reasons, and to achieve better diagnostic reliability over time, it was decided not to change to the ICD, 9th revision in 1978. Since 1 January 1994 the ICD-10 has been used.

The sample

Patients who had had their first ever discharge in the period between 1 April 1970 and 31 December 1999 were identified.
were chosen as the reference group. The hazard ratios presented thus indicate the rate of dementia for patients with a given number of prior episodes leading to admission compared with the rate for patients with one episode. Patients with two prior depressive episodes leading to admission had the same rate of dementia (HR 1.00, 95% CI 0.25 to 4.06) compared with patients with one prior depressive episode. Patients with three prior depressive episodes had a 2.89 times increased rate of dementia (95% CI 0.64 to 13.02) as patients with one prior depressive episode. Patients with three manic episodes did not have increased rate of dementia compared with patients with one depressive episode, whereas patients with four or more episodes had increased rate of dementia. The rate of dementia varied significantly with the number of episodes for patients with depressive disorder (χ² = 31.9, df = 4; p < 0.001) and for patients with bipolar disorder (χ² = 31.9, df = 4; p < 0.001). The difference between the effects of episodes in depressive disorder and bipolar disorder was significant according to a likelihood ratio test (χ² = 11.4, df = 4; p < 0.05). The proportional hazards assumption was found to be reasonably fulfilled.

Secondly, the number of episodes leading to admission was included as an ordinal variable. In this model, the hazard ratios indicate the factor change in the rate of dementia for each affective episode. Patients with depressive disorder and patients with bipolar disorder were considered separately. The episode number significantly predicted the rate of dementia (χ² = 31.9, df = 4; p < 0.001). The rate of dementia increased on average 13% (95% CI 9% to 16%) for patients with depressive disorder and 6% (95% CI 0.25 to 4.06) as patients with one depressive episode. Patients with two depressive episodes had a 2.89 times increased rate of dementia (95% CI 0.64 to 13.02) compared with patients with one depressive episode. Patients with three depressive episodes had a 2.89 times increased rate of dementia compared with patients with one depressive episode, whereas patients with four or more episodes had increased rate of dementia. The rate of dementia varied significantly with the number of episodes for patients with depressive disorder (χ² = 31.9, df = 4; p < 0.001) and for patients with bipolar disorder (χ² = 31.9, df = 4; p < 0.001). The difference between the effects of episodes in depressive disorder and bipolar disorder was significant according to a likelihood ratio test (χ² = 11.4, df = 4; p < 0.05). The proportional hazards assumption was found to be reasonably fulfilled.

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3% to 10%) for patients with bipolar disorder. To explore whether these associations also apply to patients with an interval of one year or more between the index episode and dementia, the analysis was repeated including only those cases in which time under risk was greater than one year. Similar significant effects were found (depressive disorder: 11% (95% CI: 6% to 17%); bipolar disorder: 8% (95% CI 3% to 13%)). However, a likelihood ratio test suggested that the number of episodes expressed as an ordinal variable gave an inferior fit to the data than a categorical variable (χ² = 17.0, df = 6; p = 0.01) and therefore these should be taken as a rough indication of a general increasing trend.

To explore whether the probability of getting a diagnosis of dementia simply increased with the number of admissions, the proportion of patients with a diagnosis of dementia at each admission was calculated. Among all patients admitted twice, 10.1% had a diagnosis of dementia at the second admission. The corresponding proportions were 16.1% for patients admitted three times and 15.8% for patients admitted four times. Among patients admitted five times or more, 9.3% got a diagnosis of dementia at the fifth or at a later admission. Thus, no systematic association was found between the number of admissions and the proportion of patients who were diagnosed as having dementia.

**DISCUSSION**

The present study showed that the risk of getting a diagnosis of dementia at readmission was affected significantly by the number of prior episodes leading to admission in depressive disorder and bipolar disorder. The effect of episodes leading to admission was significantly greater for patients with depressive disorder than for patients with bipolar disorder. The rate of dementia increased on average 13% (95 CI 9% to 16%) with every episode leading to admission for patients with depressive disorder and 6% (95 CI 3% to 10%) with every episode leading to admission for bipolar patients.

**Methodological considerations**

It should be acknowledged that the study only included patients who had been hospitalised at least once and that affective episodes were included only if they resulted in hospitalisation and that the outcome (a diagnosis of dementia) only related to patients who were readmitted. The latter is illustrated by the relatively low incidence of dementia found in the study. Thus, patients may have been diagnosed as having dementia in places outside psychiatric and somatic wards following first admission without this being recorded in the registers used as the source for data for the present study.

**Validity of the diagnoses of affective disorders**

The diagnoses in the register are made by different clinicians throughout Denmark and are not standardised for research purposes. Diagnostic shifts due to transfer to ICD-9 are avoided, since ICD-8 was used in Denmark until ICD-10. The validity of the affective diagnoses has been found to be good in a validity study of the register. In a random subsample of 100 patients from the register with clinical ICD-8 diagnoses of manic-depressive psychosis at first admission, 95 patients received a lifetime ICD-10 diagnosis of affective disorder according to research diagnostic criteria.

Part of the problem with the diagnostic validity is the diagnostic concordance between ICD-8 and ICD-10. From the present register, the concordance between the two diagnostic systems has been found to be as high as 84% for patients with affective disorder and 87% for patients with dementia. Thus, 13% of patients who received an ICD-8 diagnosis of dementia received an ICD-10 diagnosis of another organic mental disorder or alcohol induced mental disorder (F06-F10).

**Misclassification of the diagnosis of dementia**

It is well known that depression may be misclassified as dementia when symptoms of cognitive dysfunction are prominent—that is, some of the diagnoses of dementia in the present study in reality represent a pseudodemented state. However, pseudodementia is in general milder and more fluctuating in nature, while diagnoses of dementia that are given on discharge from psychiatric departments following admission for days or weeks, presumably describe a moderate to severe, progressive, demented condition. One may thus presume that the risk of misclassification is low. To investigate this further, 30 patients with primary affective disorder and a subsequent main or auxiliary diagnosis of dementia were randomly selected from the register (Kessing, unpublished data). The majority of these patients scored within the demented range on two different scales of cognitive assessment (70% of patients scored below 100 on the CAMCOG (Cambridge Cognitive Examination) rating scale (range 0–120) with an average of 89.5 (SD 19.7) and 96.2% of patients scored below 1 on the Global Deterioration Scale (range 1–6) with an average of 3.9 (1.0)). One may argue that patients who eventually got a diagnosis of dementia might have had several admissions previously at which dementia might have been misdiagnosed as depression. This possibility did, however, not explain our results as additional analyses revealed that the association between the number of episodes leading to admission and the risk of dementia was also found among patients with more than one year between last admission for depression and admission with a diagnosis of dementia. Anyway, it should be acknowledged that there is a need for validation of the diagnosis of dementia through criterion based diagnoses made at repeated assessments over time.

**Detection bias**

We find no reason to believe that doctors in psychiatric wards may be more observant of symptoms of dementia in patients who have been admitted with depression or mania many times than in patients who have seldom been admitted.

On the other hand, it has previously been found that the risk of getting readmitted increases with the number of prior episodes leading to admission for patients with affective disorders in Denmark. This may by itself increase the probability for patients with many prior affective episodes of getting a diagnosis of dementia at admission as these patients have a greater chance of seeing a doctor in a psychiatric ward than patients with fewer admissions. However, arguing against this possibility, it was not found that the probability of getting a diagnosis of dementia simply increased with the number of admissions (the proportion of patients who got a diagnosis of dementia at the fifth or at a later admission was less than the proportion who got a diagnosis of dementia at the second, third, or fourth admission, respectively). Thus, we do not believe that such putative detection bias explain our finding of an association between the number of affective episodes and the risk of dementia.

**Possible confounders**

It is unlikely that the demonstrated association between affective disorder and dementia was a result of intracranial infection, cerebral arteriosclerosis, cerebrovascular accidents as stroke, haemorrhage, epilepsy, intracranial neoplasm, degenerative diseases of the central nervous system, brain trauma, endocrine disorders, metabolic and nutritional disorders, systemic infections, drug or poison intoxication, or other physical conditions, since dementia related to these disorders was coded elsewhere in ICD-8 (code 292.0 – 294.9) and ICD-10 (code DF00-09). Additionally, patients who,
following the first discharge, at a subsequent admission received a discharge diagnosis of brain disorders other than dementia were censored.

Also, in our previous studies we found that the association between affective disorder and dementia could not be explained by alcoholism or other substance abuse.22 23

**Conclusion**

Although we cannot exclude methodological considerations, we do not believe that the finding of an increasing risk of getting a diagnosis of dementia with the number of affective episodes leading to hospitalisation is simply a result of methodological pitfalls.

With regard to longitudinal studies of depressive disorder, our findings are in accordance with the finding of an association between the number of depressive episodes and the risk of developing Alzheimer’s disease in the follow-up study by Andersen1 but contrast with the finding in the four-year study by Paterniti et al.24 It is possible that a follow-up time of four years is too short, as suggested by the studies of Speck et al25 and Palsson et al26 depression may act as a predictor of dementia particularly when there is a large interval between onset of depression and onset of dementia (10 years or more between the onset of depression and the onset of dementia). It is further possible that the association is more pronounced or is found only in patients with severe affective disorder for example as reflected by the need of hospitalisation as in the present study.

No prior longitudinal study of bipolar disorder has investigated the association between the number of episodes and the risk of developing dementia.

**Implications**

Data on treatment were not available in the present study. If treatment should explain the finding of an effect of the number of affective episodes on the risk of developing dementia, then this treatment should be given continuously for long periods of time for both patients with depressive disorder and patients with bipolar disorder. Antidepressants are usually only given for shorter periods in patients with bipolar disorder,26 however anxiolytics may often be given to both patient groups for a longer time. As indicated by Jorm,27 the literature is inconsistent as benzodiazepine use has been sustained prophylaxis of the evolving process of the illness.12 and further supports the hypothesis between depressive symptoms and dementia: a community-based prospective study. Arch Gen Psychiatry 1999;56:425–30.28


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**References**


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S
yncope derives from the Greek synkoptein, meaning to strike, cut off, or weary. Hippocrates and biblical texts describe victims of fainting. In the USA syncope accounts for 3% of emergency room visits and 1–6% of all hospital admissions,1 Pierre Adolph Piorry2 (1794–1879) in 1826 reported,

“When a patient faints, symptoms improve when he is laid flat.”

Piorry was a pioneer of percussion and pleximetry, best remembered for his work in chest diseases and for coining the term “uraemia.”

Thomas Addison, when describing postural syncope in adrenal failure, also noted:

“Attacks of giddiness and dinness of sight…would occur always when in the standing posture and were instantly relieved by sitting or lying down.”

Bradbury and Eggleston3 first recorded a recurrent fainting of unknown cause in 1925 in three patients subject to attacks of idiopathic orthostatic hypotension. Many lone and familial cases have subsequently appeared in the literature.4

Orthostatic hy-potension in association with other neurological diseases occurs in tabes, syringomyelia, spinal cord injury, and in polyneuropathies, Parkinsonism, multiple system atrophy (MSA), and a variety of cerebrovascular and tumorous lesions. In 1960 Milton Shy at the NIH and Glen Albert Drager of Houston described comprehensively the clinicopathology of MSA:

“A neurological syndrome associated with orthostatic hypotension—which is of adult onset and consists of orthostatic hypotension, bladder and bowel incontinence, anhidrosis, iris atrophy, amyotrophy, ataxia, rigidity and tremor; intellect is unaffected”.

Schatz later revised5 the nosology for autonomic disorders: (1) primary, autonomic failure (idiopathic orthostatic hypotension or Bradbury-Eggleston syndrome) with no neurologic defects other than autonomic dysfunction; (2) multiple system atrophy, equivalent to Shy-Drager syndrome (a sporadic, progressive, adult onset disorder characterised by autonomic dysfunction, parkinsonism, and ataxia in any combination); and (3) secondary autonomic failure delineates diseases in which the cause is known (for example diabetes, amyloidosis, dopamin beta-hydroxylase deficiency, and drug side effects).

The American Autonomic Society has defined multiple system atrophy as:

“A sporadic, progressive, adult onset disorder characterized by autonomic dysfunction, Parkinsonism, and ataxia (a failure of muscular coordination) in any combination. The main features are: Parkinsonism, cerebellar signs, autonomic failure, striatonigral degeneration, sporadic olivo-ponto-cerebellar atrophy, Shy-Drager Syndrome.”

George Milton Shy (1919–1967) qualified in Oregon in 1943 and after brief war service, in which he was wounded, studied neurology at the Montreal Neurological Institute, and at the National Hospital, Queen Square. Unusually for an American, he became MRCP London by examination, in 1947. He carried out highly regarded research on myopathies and published extensively. His work on the Kearns-Sayre syndrome, oculopharyngeal myopathy (Shy-Gonatas syndrome), and a myopathy of infants (Shy-Magee syndrome) are lasting monuments to his protean interests. In 1953 he became clinical director of the National Institute of Neurological Diseases and Blindness, Bethesda, and in 1962 chairman of neurology in Pennsylvania, before taking the chair and directorship of the New York Neurological Institute, Columbia in 1967. Aged only 47 he died suddenly a few weeks later.

Little is recorded about Drager (1917–1967), a talented physician at Baylor College of Medicine, Houston, who also died young, in the same year as Shy.