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## The Mood Disorder Questionnaire improves recognition of bipolar disorder in psychiatric care

Erkki Isometsä\*<sup>1</sup>, Kirsi Suominen<sup>1,2</sup>, Outi Mantere<sup>2</sup>, Hanna Valtonen<sup>2</sup>, Sami Leppämäki<sup>1,2</sup>, Marita Pippingsköld<sup>2</sup> and Petri Arvilommi<sup>2</sup>

Address: <sup>1</sup>Department of Mental Health and Alcohol Research, National Public Health Institute, Helsinki, Finland and <sup>2</sup>Department of Psychiatry, Jorvi Hospital, Helsinki University Central Hospital, Espoo, Finland

Email: Erkki Isometsä\* - [erkki.isometsa@ktl.fi](mailto:erkki.isometsa@ktl.fi); Kirsi Suominen - [kirsi.suominen@hus.fi](mailto:kirsi.suominen@hus.fi); Outi Mantere - [outi.mantere@hus.fi](mailto:outi.mantere@hus.fi); Hanna Valtonen - [hanna.valtonen@hus.fi](mailto:hanna.valtonen@hus.fi); Sami Leppämäki - [sami.leppamaki@hus.fi](mailto:sami.leppamaki@hus.fi); Marita Pippingsköld - [marita.pippingskold@hus.fi](mailto:marita.pippingskold@hus.fi); Petri Arvilommi - [petri.arvilommi@hus.fi](mailto:petri.arvilommi@hus.fi)

\* Corresponding author

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### Abstract

**Background:** We investigated our translation of The Mood Disorder Questionnaire (MDQ) as a screening instrument for bipolar disorder in a psychiatric setting in Finland.

**Methods:** In a pilot study for the Jorvi Bipolar Study (JoBS), 109 consecutive non-schizophrenic psychiatric out- and inpatients in Espoo, Finland, were screened for bipolar disorder using the Finnish translation of the MDQ, and 38 of them diagnostically interviewed with the SCID.

**Results:** Forty subjects (37%) were positive in the MDQ screen. In the SCID interview, twenty patients were found to suffer from bipolar disorder, of whom seven (70%) of ten patients with bipolar I but only two (20%) of ten with bipolar II disorder had been previously clinically correctly diagnosed. The translated MDQ was found internally consistent (alpha 0.79) and a feasible screening tool.

**Conclusions:** Bipolar disorder, particularly type II, remains commonly unrecognized in psychiatric settings. The Mood Disorder Questionnaire is a feasible screen for bipolar disorder, which could well be integrated into psychiatric routine practice.

### Background

Although bipolar disorder is a major public health issue, it is commonly unrecognized even in psychiatric settings [1,2]. Feasible screening instruments are needed to improve recognition and diagnosis of the various forms of the illness. The Mood Disorder Questionnaire [3] is a recently developed and simple screening method already validated in a US multicenter study. The instrument was found to have relatively good sensitivity (0.73) and very good specificity (0.90) in samples comprising mostly patients with uni- and bipolar mood disorders in aca-

demic centers [3]; in a further general population study the sensitivity turned out to be very low (0.28), but specificity (0.97) remarkably high [4]. The instrument has so far been little investigated by others than its developers.

In the present pilot study, using our Finnish translation of the Mood Disorder Questionnaire among unselected psychiatric patients, we investigated its psychometric and screening properties, and its feasibility in improving recognition of type I and II bipolar disorder.

## Methods

The present investigation was a pilot study for the ongoing Jorvi Bipolar Study (JoBS). The screening for bipolar disorder was conducted at the Department of Psychiatry at Jorvi Hospital, part of the Helsinki University Central Hospital, from 1st–31st October, 2001. The Department has a catchment area of approximately 250 000 inhabitants, but this pilot study was conducted only in two selected community mental health centres (Leppävaara and Kirkkonummi), and in the psychiatric outpatient clinic of Jorvi Hospital and three of its psychiatric wards. The whole JoBS project has been evaluated and accepted in the Ethics Committee of the Helsinki and Uusimaa Hospital District, and it complies with the principles of the Helsinki Declaration.

Three patient groups were screened: (a) all new patients who were referred to treatment in the Department of Psychiatry; (b) all patients who had earlier received treatment in the Department, but now had a new referral, and (c) those already in contact with the facilities, without a clinical diagnosis of ICD-10 schizophrenia, and now showing signs of deteriorating clinical state after at least two months of limited or no symptoms. The aim was to include all incident episodes of bipolar disorder among patients receiving treatment.

The Mood Disorder Questionnaire [3] is a short self-report screening instrument, and was translated into Finnish by the authors. The first question includes 13 items, symptoms or behaviors related to a manic or hypomanic syndrome. The second question asks whether several of the symptoms have been experienced during the same time period, and the third asks about resulting problems, classified as minor, moderate or serious. The screen is regarded positive when seven or more positive symptoms have occurred, several within the same episode, causing moderate to severe problems. If the MDQ was negative, the attending professional was also asked whether bipolar disorder might be present despite a negative screen (clinical suspicion).

All patients with a positive screen or clinically suspected of having bipolar disorder were invited to be interviewed with the Structured Clinical Interview for DSM-IV Axis I Disorders, Clinician Version, SCID-CV [5]. In addition, a systematic sample of every third patient with 4–6 positive items in question 1 of the MDQ was drawn from each setting. All interviewers were either psychiatrists or residents with several years experience in psychiatry and relevant training. The first two authors supervised the diagnostic process.

## Results

### Screening phase

The group of eligible patients comprised 113 subjects, of whom two (2%) refused and two (2%) were excluded because of not speaking Finnish. Altogether 109 (96%) patients were screened. Their mean age was  $37.9 \pm 11.4$  years, and 56 (51%) were female. Forty subjects (37%) were positive in the MDQ. Twenty-nine (27%) subjects had negative MDQ but 4–6 positive items in question 1. This latter group included four of the five subjects (5% of the total) with negative MDQ who were clinically suspected of possibly suffering from bipolar disorder.

The frequency of positive responses to the various symptom items of question 1 ranged from 27% (more social) to 74% (racing thoughts). In the vast majority of those reporting seven or more items, several symptoms had occurred within the same time period (in 87%), and moderate to serious problems were present (in 91%). The internal consistency of the translated instrument was found to be good (Cronbach's alpha 0.79).

### Diagnostic findings

Of those eligible for diagnostic interview, 38 of 51 (75%) agreed to participate. Of those interviewed, 20 (53%) were diagnosed with bipolar disorder: ten (26%, 5 males) with type I and ten (26%, 3 males) with type II (Table 1). The mean ages of the two groups were  $38.2 \pm 8.0$  years and  $39.1 \pm 12.5$  years, respectively. Using the SCID-interview as the gold standard, the MDQ screening produced 9 false positives (5 patients with unipolar MDD; one schizoaffective, brief psychotic, and personality disorders; plus an amphetamine-induced psychosis) and three false negative patients (one type I with < 7 acknowledged items, one type II patient with only minor problems due to hypomania, and one type II with both). Less than half of those diagnosed with bipolar disorder in the SCID interview had had a ICD-10 diagnosis of bipolar disorder, mania or hypomania, or bipolar schizoaffective disorders before the interview. Of the bipolar I patients seven (70%) had a relevant diagnosis before the interview, compared to only two of the bipolar II patients (20%). Despite the small sample size, the difference approached statistical significance (Fisher's exact test,  $p = 0.07$ ).

### The properties of the MDQ as a screen within the diagnostically interviewed sample

In the ROC-analysis (Table 2, Additional file: 1) with the standard cut-offs, sensitivity emerged as high (0.85), but specificity only moderate (0.47). However, the optimal cut-off within this sample was found to be eight symptoms but accepting also minor problems due to episodes (sensitivity 0.90, specificity 0.59). Ignoring severity of problems caused (question three) altogether resulted in very low specificity.

**Table 1: Findings of the Mood Disorder Questionnaire screening vs. diagnostic SCID-interview (N = 37)<sup>a</sup>**

Diagnostic SCID-interview	negative	MDQ-screening positive	total
No bipolar disorder	8	9	17
Bipolar disorder, type I	1	9	10
Bipolar disorder, type II	2	8	10
	11	26	37

<sup>a</sup> One case excluded due to missing response to question 3.

**Discussion**

We found the Mood Disorder Questionnaire to be a feasible method for improving recognition of bipolar disorder, which has clearly been a problem. However, by definition, hypomania involves no marked impairment. Whether necessitating moderate to severe problems to be caused by it is useful in screening should be further investigated. Having minor problems might be enough.

This was a pilot study for the current Jorvi Bipolar Study (JoBS). Several methodological limitations should be noted, some suggesting caution in interpreting the findings. The sample of patients screened with the Mood Disorder Questionnaire was not large (N = 109), albeit probably representative of psychiatric in- and outpatients in the area. The number of patients interviewed with SCID was relatively small. Those interviewed with SCID were not a random sample of all patients screened, but a sample of cases suspected of having bipolar disorder on the basis of the finding in the MDQ, or other clinical factors. The screen thus influenced whether a diagnostic interview was conducted. It is therefore clear that the ROC-analysis overestimated sensitivity and underestimated specificity, and cannot be compared with respective estimates from unselected samples [3]. It only shows that within this sample, the optimal cut-offs would have been different. Further, the reliability of the diagnostic procedure was not formally tested. Finally, the generalizability of our findings within Finland, or to other countries, is not known. Given the relatively good resources and interest in bipolar disorder in the Jorvi psychiatric facilities, we expect recognition to be at least not worse than elsewhere in Finland. Our finding of underrecognition is comparable with those from other countries [6–10].

The Mood Disorder Questionnaire appears to be a feasible method for improving the recognition of bipolar disorder. The internal consistency of the translated instrument was almost as good (Cronbach's alpha 0.79 vs. 0.90) as in the original validation study [3]. Our study population was not a selected sample of patients with mood disorders, but

rather an ordinary sample of secondary care psychiatric patients, schizophrenia excluded. Nevertheless, our prevalences of positive items (27–74% vs. 34–77%) and bipolar disorder (53% vs. 55%) were similar to the original validation study figures, although the proportion of type II disorder was higher (50% vs. 24%) in ours. In the third published study of the MDQ [11] that focused on its factor structure, rates of positive items were convergent, although slightly lower (12–65%) among private practice mood disorder patients.

The most important public health problem related to bipolar disorder is the remarkable proportion of patients who have unrecognised bipolar II disorder [6–10]. The majority of unrecognised patients in our sample, too, had bipolar II disorder. It is therefore vital that the MDQ is also sensitive to bipolar II. Hypomania, as defined in the DSM-IV, must not be related to marked impairment [12]. Thus the last question in the MDQ, necessitating moderate to severe problems due to episodes, appears to have a higher threshold for impairment. In our diagnosed sample, accepting minor impairment too, but necessitating eight symptoms, was found to be the optimal cut-off. Nevertheless, even using the standard cut-offs several patients with previously unrecognised bipolar II disorders were identified, and in fact, ignoring the last question completely resulted in lower specificity. It is to be noted that our findings are based only on ten bipolar II patients and an enriched subsample of patients with bipolar disorder. We suggest that the optimal cut-offs for bipolar II disorders should be further investigated in larger and representative patient samples.

**Conclusions**

The Mood Disorder Questionnaire seems to be a feasible method for improving the recognition of bipolar disorder. Whether screening is actually beneficial is related to the quality of current routine diagnostic procedures. It is useful in psychiatric settings only if recognition without it is a problem, which according to our findings is certainly true. The proportion of previously unrecognised cases was

particularly high in bipolar II disorders, only a few of whom had previously received the correct diagnosis. Our findings support the value and feasibility of screening for bipolar disorder with the MDQ in psychiatric settings.

### Competing interests

No competing interests.

### Authors' contributions

EI supervised the study, analyzed the data and wrote the paper. KS supervised the fieldwork, interviewed patients, and participated in writing the paper. OM, HV, SL, MP and PA interviewed patients and commented on the manuscript. All authors participated in the translation of the Mood Disorder Questionnaire, plus read and approved the final manuscript.

### Additional material

#### Additional file 1

Click here for file  
[<http://www.biomedcentral.com/content/supplementary/1471-244X-3-8-S1.rtf>]

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