



## SCREENING AND RISK ASSESSMENT FOR DEPRESSION IN COMMUNITY PHARMACY-PILOT STUDY

Antonia Kondova<sup>1</sup>, Anna Todorova<sup>1</sup>, Antoaneta Tsvetkova<sup>2</sup>, Mariana Arnaoudova<sup>3</sup>, Kalina Andreevska<sup>4</sup>, Daniela Grekova<sup>4</sup>,

1) Department of Pharmaceutical Sciences, Faculty of Pharmacy, Medical University, Varna, Bulgaria.

2) Medical College, Medical University, Varna, Bulgaria.

3) Department of Psychiatry and Medical Psychology, Faculty of Medicine, Medical University, Varna, Bulgaria.

4) Department of Pharmaceutical Sciences, Faculty of Pharmacy, Medical University, Plovdiv, Bulgaria.

### SUMMARY

**Background:** Depression is the most common mental illness affecting more than 300 million people worldwide and is a significant risk factor for morbidity and mortality. In most cases, it may remain undetected in primary care. Comprehensive screening tools for diagnosing depression might facilitate early detection. As the most accessible health professionals, pharmacists can play an important role in helping to identify individuals at risk.

**Objective:** To differentiate individuals at risk of depression who are seeking a pharmacist consultation and are promptly directed to a psychiatrist.

**Methods:** Depression screening tools The Patient Health Questionnaire (PHQ-2) and (PHQ-9) were applied to 83 individuals with symptoms such as feeling down, tiredness and sleep disturbances for more than 2 weeks, who seek consultation at a pharmacy. Screening with the PHQ-2 was the first step. Patients who screen positive were further evaluated with the PHQ-9.

**Results:** In 70% of the individuals, the PHQ-2 test was positive. After completing PHQ-9, it was found that approximately 55% out of them had indications of mild to moderate depression and were directed to a psychiatrist for further evaluation. Over 50% of suspected depressive individuals had a concomitant chronic disease.

**Conclusion:** Screening for depression should be a routine part of healthcare. Particular attention should be paid to patients with comorbid chronic illnesses, as depression often remains hidden, thus leading to more difficult diagnosis and treatment. Screening would also increase the recognition of depression in patients who have few emotional symptoms but many somatic ones.

**Keywords:** depression, screening, Patient Health Questionnaire, community pharmacy, cardiovascular diseases

### INTRODUCTION:

Depression is the most common mental illness affecting more than 300 million people worldwide and is a significant risk factor for morbidity, disability and mortality. According to data from Eurostat (extracted in January 2017), in 2014 at EU-28 level 3.5 % of the population in Bulgaria reported having chronic depression [1]. Depression has been predicted to be the leading cause of disease burden in 2030 by the World Health Organization (WHO) [2]. It is widespread among people of all ages and social backgrounds, with women being twice as likely as men. The disease usually occurs during the third and fourth decades of life, with an increase in frequency and a reduction in the age of the disease occurring in recent years. In most cases, it may remain undetected in primary care. The high prevalence of depression in patients with physical disorders requires the identification of vulnerable individuals as an important step in the further of these patients.

For that reason, comprehensive screening tools for diagnosing depression might facilitate its early detection [3]. Such screening tools should assess the possibility of depression even in individuals presenting primarily with somatic symptoms and relevantly inform about their general health. Primary care medical specialists should be familiar from one side with the signs and symptoms of depression and on the other -with the popular terms that people use concerning emotional problems. There are some relevant questions that could assist in the identification of depressive symptoms. Although screening self-administrating tools are not diagnostic tools they indicate the need for further evaluation.

The role of pharmacists is not generally accepted and comprehensively defined, but with the growing number of mental disorders and their great impact on patients' everyday life, they will be successfully positioned in the multidisciplinary mental health care teams [4].

**The aim** of our study is to differentiate individuals at risk of depression who have been consulted by a phar-

macist and further referred to a specialist.

**METHODS:**

The Patient Health Questionnaire (PHQ2) and (PHQ9) were applied to 83 individuals (mean age 57,8 years) with some characteristic symptoms as feeling down, tiredness and sleep disturbances for more than 2 weeks, who sought consultation at a pharmacy.

The PHQ-2 is a screening tool that inquires about the frequency of depressed mood and anhedonia over the past 2 weeks, scoring each item as 0 (“not at all”) to 3 (“nearly every day”) with a score ranging from 0 to 6. The authors identified a cut-off score of 3 as the optimal cut point for screening purposes and stated that a cut point of 2 would enhance sensitivity. Patients who screen positive should be further evaluated with the PHQ-9, other diagnostic instruments or direct interview to determine whether they meet criteria for a depressive disorder [5, 6]. A PHQ-2 score of greater than 3 has a sensitivity of 83% and a specificity of 92% for major depression [7]. The PHQ-2 includes the first 2 items of PHQ-9: “Little interest or pleasure in doing things” and “Feeling down, depressed, or hopeless.”

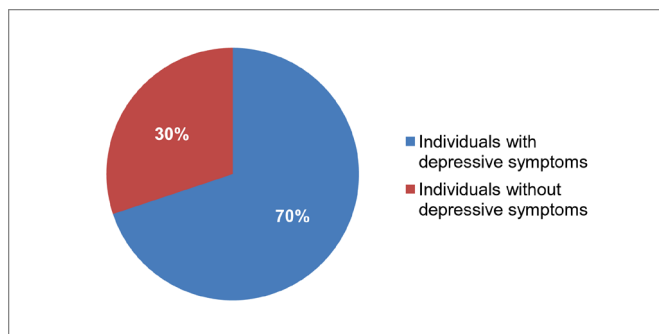
The PHQ-9 is a potentially valuable tool for diagnosis and management of depression because it can generate a diagnosis of major depression, as well a continuous score to monitor treatment. The PHQ-9 has 9 questions with a score ranging from 0 to 3 for each question (maximum score of 27). The PHQ-9 establishes the clinical symptoms of depression. The cut point of the PHQ-9 is equal or greater than 10, which has a sensitivity of 88% and a specificity of 88% for major depression [8]. A threshold score of 15 or more is used in some settings to consider initiating treatment with antidepressants [9]. Once the PHQ-9 is completed, and the test proves to be positive, the patient should be directed to a psychiatrist [10, 11, 12].

Descriptive statistics were used to calculate relative share and average values. A graphic analysis has been made. The processing of the results and the construction of the graphics were done with MS Excel.

**RESULTS AND DISCUSSION:**

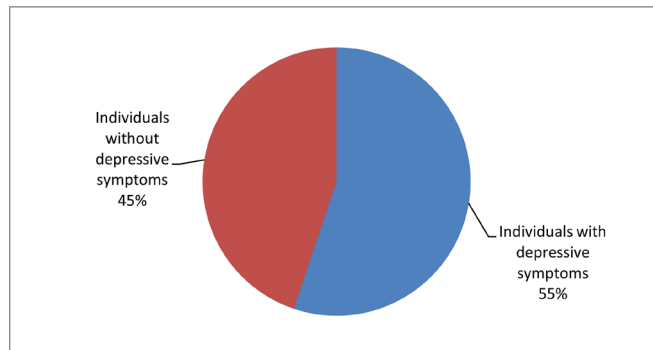
According to the data from PHQ-2, positive for depressive symptoms were 70% of individuals. The distribution of participants with positive PHQ-2 is presented in Fig.1.

**Fig. 1.** Screening with the PHQ-2



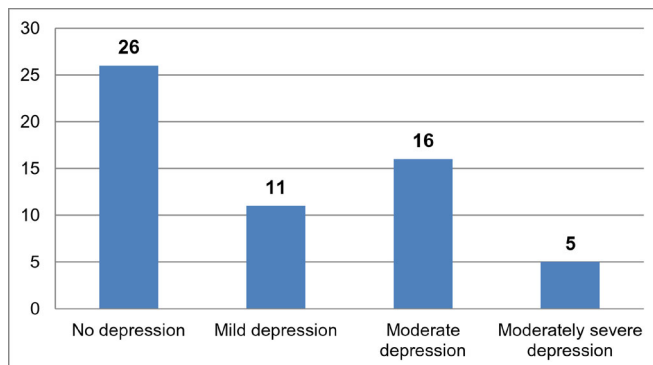
Patients who screen positive by PHQ-2 were further evaluated with the PHQ-9. It was found that approximately 55% of the individuals had indications of mild to moderate depression and were directed to a psychiatrist to confirm the diagnosis (fig.2).

**Fig. 2.** Screening with the PHQ-9



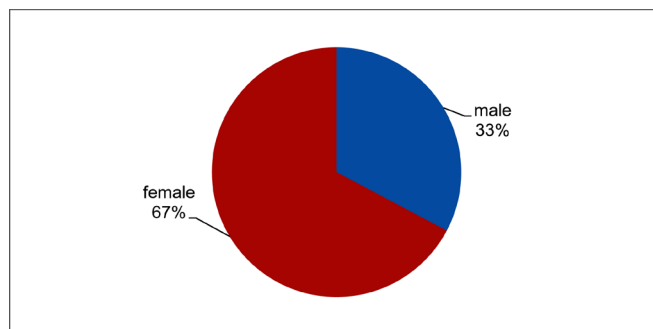
The severity of depression by PHQ-9 is shown in fig. 3. As it is seen, 26 individuals had no symptoms of depression, 11 individuals presented with scores for mild depression and 16 individuals with moderate depression and 5 individuals with moderately severe depression.

**Fig. 3.** Depression severity by PHQ-9



Gender and age are factors influencing depressive symptoms [13]. In men, depression is definitely less common. In our study, females (67%) were twice more likely to develop depression than male individuals- (33%) (fig. 4).

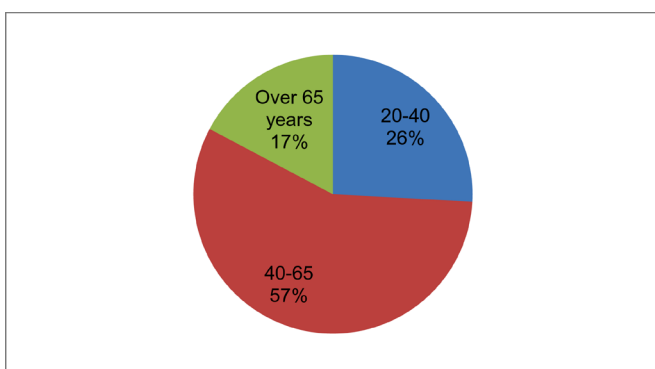
**Fig. 4.** Gender distribution of individuals with depressive symptoms by PHQ-9



For both genders, depression is most common in those who are 25-44 years of age, and least common for those over the age of 65. According to some authors rates of MDD are by and large lower in healthy community-dwelling elderly persons than in younger adult populations, ranging from 1% to 3% [14]. It should be pointed out that these rates depend on factors as different settings, medical and psychiatric comorbidity, social conditions etc. Most common, hereditary factors and personality traits are associated with the early onset, while vascular cognitive impairment is associated with the late onset.

In our study, the proportion of individuals with depressive symptoms in the 40-65 age group was the highest (57%), followed by the 20-40 age group (26%) and over 65 years 17% of participants were with positive for depression PHQ-9 (Fig. 5).

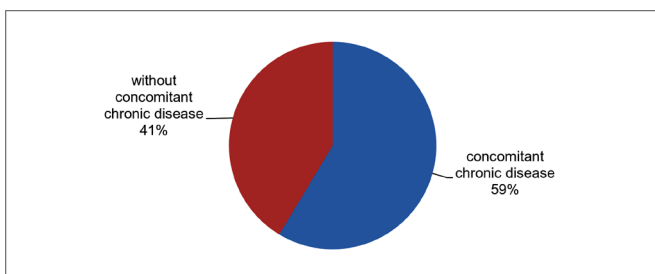
**Fig. 5.** Distribution of age groups



Depressive disorders are highly comorbid with somatic diseases [15]. This comorbidity significantly impacts the clinical presentation of depression, impedes the outcomes both of depression and of the physical disorders, increase mortality and medical costs than when these conditions are present alone. We should bear in mind that depressive somatic symptoms may mask the somatic disease.

Our study found that over 50% of individuals suspected of depression have a concomitant chronic disease (Fig.6).

**Fig. 6.** Individuals with depressive symptoms and concomitant chronic disease



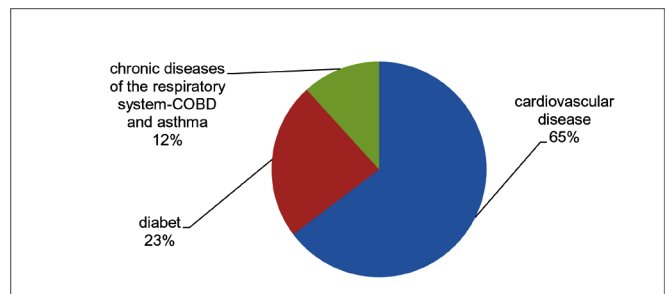
In recent years, there has been much evidence of the link between depression and cardiovascular disease (CVD) [16]. The incidence of clinical depression in patients with coronary artery disease is three times higher than in the gen-

eral population. Approximately every sixth patient with acute myocardial infarction unfolds a major depressive episode, which in one-third of cases does not respond satisfactorily to the usual treatment. Myocardial infarction combined with depression results in much higher mortality within 6 months compared to patients who are not affected by the depressive disorder. According to American Heart Association (AHA) recommendations, patients with cardiovascular disease should be regularly screened for depression [17].

Another important comorbidity is type 2 diabetes [18]. There is evidence that the prevalence of depression is significantly higher in patients with type 2 diabetes than in those without, the relative risk for the occurrence of depression after diabetes being 1.15 and 1.6 for the occurrence of diabetes after depression with the comment that depression is a stronger risk factor for the development of depression than the reverse [19, 20, 21].

The results of our study are consistent with the literature data and show that the largest share is positive for PHQ-9 and concomitant cardiovascular disease— 65%. This includes patients with hypertension, ischemic heart disease. Second come individuals with diabetes 23%, followed by those with chronic respiratory diseases, including chronic obstructive pulmonary disease and asthma- 12%. The distribution of the most common chronic diseases in PHQ-9 depressive symptomatology individuals is presented in Fig.7.

**Fig. 7.** Comorbid chronic physical diseases in depression



## CONCLUSION

Depression screenings should be a routine part of healthcare. Regardless of the limitations of self-reported depression screening tools, it is better to use them to screen for depression than risk missing patients who are suffering from a depressive disorder. Particular attention should be paid to patients with chronic illnesses, as depression often remains hidden and due to significant comorbidity, leading to more difficult diagnosis and treatment. Particular attention should be paid to patients suffering from cardiovascular disorders. These patients should be regularly monitored for depression. Effective treatment of depression aims to improve the emotional, physical, social functioning and quality of life of the patients, to reduce the health care needs and decrease mortality.

## REFERENCES:

1. Eurostat. Mental health and related issues statistics. Data extracted in September 2017. [\[Internet\]](#)
2. World Health Organization. The global burden of disease: 2004 update. Geneva, Switzerland: WHO Press, 2008. [\[Internet\]](#)
3. O'Reilly CL, Wong E, Chen TF. A feasibility study of community pharmacists performing depression screening services. *Res Social Adm Pharm.* 2015 May-Jun;11(3):364-81. [\[PubMed\]](#) [\[CrossRef\]](#)
4. Rubiero-Valera M, Chen TF, O'Reilly CL. New roles for Pharmacists in Community Mental Health Care: A Narrative Review. *Int J Environ Res Public Health.* 2014 Oct;11(10):10967-90. [\[PubMed\]](#) [\[CrossRef\]](#)
5. Spitzer RL, Kroenke K, Williams JB. Validation and utility of a self-report version of PRIME-MD: the PHQ primary care study. Primary Care Evaluation of Mental Disorders. Patient Health Questionnaire. *JAMA.* 1999 Nov;282(18):1737-44. [\[PubMed\]](#) [\[CrossRef\]](#)
6. Kroenke K, Spitzer RL, Williams JB. The Patient Health Questionnaire-2: validity of a two-item depression screener. *Med Care.* 2003 Nov;41(11):1284-92. [\[PubMed\]](#) [\[CrossRef\]](#)
7. Maurer DM. Screening for depression. *Am Fam Physician.* 2012 Jan;85(2):139-44. [\[PubMed\]](#)
8. Kocalevent RD, Hinz A, Brahler E. Standardization of the depression screener patient health questionnaire (PHQ-9) in the general population. *Gen Hosp Psychiatry.* 2013 Sep-Oct;35(5):551-5. [\[PubMed\]](#) [\[CrossRef\]](#)
9. Arroll B, Goodyear-Smith F, Crengle S, Gunn J, Kerse N, Fishman T, et al. Validation of PHQ-2 and PHQ-9 to Screen for Major Depression in the Primary Care Population. *Ann Fam Med.* 2010 Jul-Aug;8(4):348-53. [\[PubMed\]](#) [\[CrossRef\]](#)
10. Kroenke K, Spitzer RL, Williams JB. The PHQ-9: validity of a brief depression severity measure. *J Gen Intern Med.* 2001 Sep;16(9):606-13. [\[PubMed\]](#)
11. Pinto-Meza A, Serrano-Blanco A, Penarrubia MT, Blanco E, Haro JM. Assessing depression in primary care with the PHQ-9: can it be carried out over the telephone? *J Gen Intern Med.* 2005 Aug;20(8):738-42. [\[PubMed\]](#) [\[CrossRef\]](#)
12. Beard C, Hsu KJ, Rifkin LS, Busch AB, Bjorgvinsson T. Validation of the PHQ-9 in a psychiatric sample. *J Affect Disord.* 2016 Mar;193:267-73. [\[PubMed\]](#) [\[CrossRef\]](#)
13. Angst J, Gamma A, Gastpar M, Lepine JP, Mendlewicz J, Tylee A. Gender differences in depression. *Eur Arch Psychiatry Clin Neurosci.* 2002 Oct;252(5):201-9. [\[PubMed\]](#) [\[CrossRef\]](#)
14. Kessler RC, Birnbaum H, Bromet E, Hwang I, Sampson N, Shahly V. Age differences in major depression: results from the National Comorbidity Survey replication (NCS9R). *Psychol Med.* 2010 Feb;40(2):225-37. [\[PubMed\]](#) [\[CrossRef\]](#)
15. Katon W, Lin EH, Kroenke K. The association of depression and anxiety with medical symptom burden in patients with chronic medical illness. *Gen Hosp Psychiatry.* 2007 Mar-Apr;29(2):147-55. [\[PubMed\]](#) [\[CrossRef\]](#)
16. O'Neil A, Williams ED, Stevenson CE, Oldenburg B, Berk M, Sanderson K. Co-morbid cardiovascular disease and depression: sequence of disease onset is linked to mental but not physical self-rated health. Results from a cross-sectional, population-based study. *Soc Psychiatry Psychiatr Epidemiol.* 2012 Jul;47(7):1145-51. [\[PubMed\]](#) [\[CrossRef\]](#)
17. Haddad M, Walters P, Phillips R, Tsakok J, Williams P, Mann A, et al. Detecting depression in patients with coronary heart disease: a diagnostic evaluation of the PHQ-9 and HADS-D in primary care, findings from the UP-BEAT-UK study. *PLoS One.* 2013 Oct;8(10):e78493. [\[PubMed\]](#) [\[CrossRef\]](#)
18. Katon W, Fan MY, Unützer J, Taylor J, Pincus H, Schoenbaum M. Depression and diabetes: a potentially lethal combination. *J Gen Intern Med.* 2008 Oct;23(10):1571-5. [\[PubMed\]](#) [\[CrossRef\]](#)
19. Ali S, Stone MA, Peters JL, Davies MJ, Khunti K. The prevalence of co-morbid depression in adults with type 2 diabetes: a systematic review and meta-analysis. *Diabet Med.* 2006 Nov;23(11):1165-73. [\[PubMed\]](#) [\[CrossRef\]](#)
20. Mezuk B, Eaton WW, Albrecht S, Golden SH. Depression and type 2 diabetes over the lifespan: a meta-analysis. *Diabetes Care.* 2008 Dec;31(12):2383-90. [\[PubMed\]](#) [\[CrossRef\]](#)
21. Schmidt do Prado-Lima PA. Medical comorbidities and functioning in depression: a clinical perspective. *Medicographia.* 2014; 36(4):464-469.

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### Address for correspondence:

Antoniya Kondova  
Department Pharmaceutical Sciences, Faculty of pharmacy, Medical University, Varna,  
84, Tzar Osvoboditel Str., 9000 Varna, Bulgaria.  
e-mail: [antoniakondova@mail.bg](mailto:antoniakondova@mail.bg)