

# Comorbidity of Anxiety and Depression with Hypertension, Diabetes, and Cardiovascular Disease: A Selective Systematic Review from India

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## Abstract

**Context:** Non-communicable diseases (cardiovascular diseases, hypertension, and diabetes) and comorbid common mental disorders are of public health concern because of their high morbidity and mortality rates. The authors undertook a systematic review of studies that reported the prevalence of common mental disorders among non-communicable diseases, specifically in India.

**Evidence acquisition:** Relevant databases (Medline, Google Scholar, EBSCO, and ProQuest) were searched until May 2021. Descriptive and observational studies from the mentioned databases were included.

**Evidence synthesis:** Of the total 6,515 studies, the electronic literature search identified 4,307 studies. Manual cross-referencing identified an additional 2,208 studies. Only 17 studies met the criteria and were included for the review.

**Findings:** Twelve studies focused on the prevalence of anxiety and depression in patients with diabetes, four studies focused on cardiovascular disease, and one on the prevalence of depression in hypertension. The prevalence of anxiety disorder and depression was 3.9–44% and 8–44%, respectively.

**Conclusion:** High prevalence of anxiety and depression is seen in people with diabetes, indicating these are of serious public health concerns in India.

## INTRODUCTION

Non-communicable diseases (NCD) are defined as medical conditions that cannot be transmitted.<sup>1</sup> They are characterised by a chronicity of at least 3 months and are progressive in nature.

The conditions categorised as NCDs are cardiovascular diseases (CVD), respiratory diseases, obesity, gastrointestinal disorders, diabetes, cancer, and endocrine and metabolic disorders.<sup>2</sup> NCDs such as heart diseases, cancer, stroke, and diabetes are the leading

cause of mortality in the world.<sup>3</sup> The combined burden of these diseases is rising fast among lower-income countries.

According to the Global Status Report on NCDs (2010), 80% of CVD and diabetes, and 90% of deaths from chronic obstructive pulmonary disease, occur in low- and middle-income countries.<sup>4</sup> The recent edition (2020) indicates that 70% of deaths worldwide are due to NCDs.<sup>4</sup>

The modifiable risk factors associated with NCDs are tobacco use, physical inactivity, unhealthy diet, and the harmful use of alcohol.<sup>4</sup> Tobacco accounts for 7.2 million deaths every year, more than 3.3 million annual deaths have been attributed to alcohol use from NCDs, including cancer, and 1.6 million deaths can be attributed to insufficient physical activity.<sup>5</sup> These risk factors lead to metabolic and psychological changes in the body, such as raised blood pressure, overweight/obesity, hyperglycaemia, and hyperlipidaemia.<sup>4</sup>

There have been increasing calls to include common mental disorders (CMD) such as depression and anxiety under the umbrella of NCDs. However, the prevention and control of both diseases remain separate and independent. CMDs typically include anxiety, depression, and somatic complaints and cause limitations in daily activities.<sup>6</sup> CMDs are identified as the leading contributors of disability globally, but they are less likely to receive treatment.<sup>7</sup> This reflects stigmatisation, and they are undetected most of the time.<sup>7,8</sup> The aetiological factors for CMDs can be psychological, social, and biological. The causal mechanism of the risk factors of NCDs and CMDs also operate at the individual level (genetic factor), psychological level, and societal level (social determinants). Thus, CMDs and NCDs are interlinked and highly comorbid. The pathways of comorbidity have been described to be bidirectional in nature.<sup>9,10</sup> For example, diabetes elevates the risk of depression and vice versa.<sup>11</sup> These conditions share common diathesis contributing to the onset of other comorbid conditions.<sup>12</sup> The psychological burden of being ill and associated health factors can act as triggering factors in the development of depression and anxiety among patients with diabetes.<sup>13</sup> Similarly, depression leads to unhealthy dietary practices, less physical activity, tobacco use, harmful use of alcohol, and weight gain, which are risk factors

for diabetes. Risk factors for NCDs tend to cluster together.<sup>8</sup>

Depression and anxiety have also been recognised as systemic illnesses that negatively affect physical health.<sup>14,15</sup> People with CMDs may use nicotine and alcohol, which can further affect their physical health. This not only contributes to NCDs but also increases the burden of depression and anxiety. Tobacco and alcohol use are also linked with common and severe mental illness. Along with NCDs, from a societal perspective, tobacco use in mental disorders might be a significant contributor to premature mortality.<sup>16</sup>

The likelihood of depression tends to double for patients with diabetes, when compared with the general population without diabetes.<sup>17,18</sup> Patients with diabetes may have excessive fear and worry about the disease, which can result in anxiety, and anxiety disorders are associated with hypertension.<sup>19,20</sup> Hypertension may trigger an anxiety disorder and, conversely, an anxiety disorder may lead to hypertension.<sup>21</sup> Studies suggest that individuals experiencing depression are at risk of developing hypertension, and are predisposed to stroke and ischaemic heart disease (IHD).<sup>22,23</sup> They are more likely to develop CVD and have a higher mortality rate. Hypertension is also an important risk factor for the development of coronary heart disease.<sup>24</sup> There exists a graded relationship among both: the more severe the depression, the higher the risk of mortality and other cardiovascular events. The economic indicators relating to CVD and depression are high medical cost, increased health service utilisation, and less productivity. These factors contribute to poor quality of life (QoL).<sup>25</sup>

There is also evidence that lifestyle risk factors such as diet, physical inactivity, smoking, and alcohol use, besides non-modifiable risk factors such as age, poverty, and gender, contribute to comorbidity.<sup>12</sup> It is also evident that depression and anxiety are prevalent in people with physical comorbidities, specifically in South Asia.<sup>26</sup> There is an increasing trend in the prevalence of both CMDs and NCDs in developing countries such as India.<sup>26</sup> Major risk factors of NCDs in India are high systolic blood pressure, high fasting plasma glucose, and high BMI.<sup>27</sup> In 2013, there were an estimated 65.1 million diabetes cases; this is expected to increase up to 109.0 million in 2035.<sup>28</sup>

The World Economic Forum (WEF) estimated that countries such as Brazil, China, India, and the Russian Federation lose more than 20 million productive life years annually to NCDs.<sup>3</sup>

Most of the time, depression goes undetected despite its high prevalence in NCDs. Patients do not receive adequate treatment, which further deteriorates their physical health. This suggests addressing the risk factors and integrated management of mental disorders and NCDs. In view of this, it is important to screen patients for NCDs and CMDs. Thus, this study aimed to review the literature from India on the prevalence of CMDs in NCDs and vice versa.

## OBJECTIVES

- To study the prevalence of anxiety and depression in hypertension, diabetes, and CVD in India; and
- To study the prevalence of hypertension, diabetes, and CVD in anxiety and depression in India.

## METHODS

### Evidence

Electronic databases such as PubMed, Medline, EBSCO, CINAHL, ProQuest, and Google Scholar were searched for the time period of 1980 to May 2021. The keywords used for the search strategy were "prevalence," "anxiety," "depression," and "hypertension;" "prevalence," "anxiety," "depression," and "diabetes mellitus;" and "prevalence," "anxiety," "depression," and "cardiovascular diseases." The evidence was screened for inclusion criteria using a three-stage approach: reviewing the title, abstract, and full text. The authors included studies that were published between 1980 and May 2021 in indexed and non-indexed journals, and descriptive and observational studies that allowed an estimation of the prevalence of CMDs and NCDs. For this review, NCDs included hypertension, diabetes, and CVD, whereas CMDs included anxiety and depression. The authors included studies from inpatient, outpatient, and community settings, and whose participants were adults (age >18 years) from India. Studies that did not relate to NCDs and CMDs, and those that were randomised control trials or reviews (systematic or non-

systematic), were excluded. Studies conducted only amongst the elderly or adolescents were also excluded.

### Data Extraction

Microsoft Excel was used to enter the extracted information for the review. The data extraction form included authors' names, publication year, study design, participants' profile, sample size, measurements, and outcomes. One author (Rajan) conducted the screening independently by reviewing articles and abstracts. Full-text articles of the abstracts that met the inclusion criteria were then reviewed by two independent authors (Rajan and Chaturvedi). Data of these studies were extracted by two independent authors (Rajan and Krishna). Discrepancies in article inclusion, data extraction, and bias assessment were resolved by consensus or by referring to the third author (Muliya).

### Risk of Bias

The quality of the included studies was assessed by the National Institutes of Health (NIH) Quality Assessment Tool for Observational Cohort and Cross-Sectional Studies.<sup>29</sup> Three independent authors (Muliya, Krishna, and Rajan) assessed the quality of studies, and disparities were resolved by consensus. The quality rating was based on the total number of "yes" responses and was rated as follows: poor: <50%; fair: 50–75%; and good: ≥75%.

## RESULTS

A total of 17 studies that met the inclusion criteria were reviewed. **Figure 1** illustrates the screening and selection process in the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flow diagram. Of the total 6,515 studies, the electronic literature search identified 4,307 studies. Manual cross-referencing additionally identified 2,208 studies. After removing 100 duplicate studies, 6,415 were screened and 6,397 were excluded as they did not meet the inclusion criteria. The full text was only reviewed in the remaining 25 studies. Further, eight studies were excluded as they did not concur with the inclusion criteria, resulting in 17 studies for the review.

The characteristics of the included studies are provided in **Table 1**.

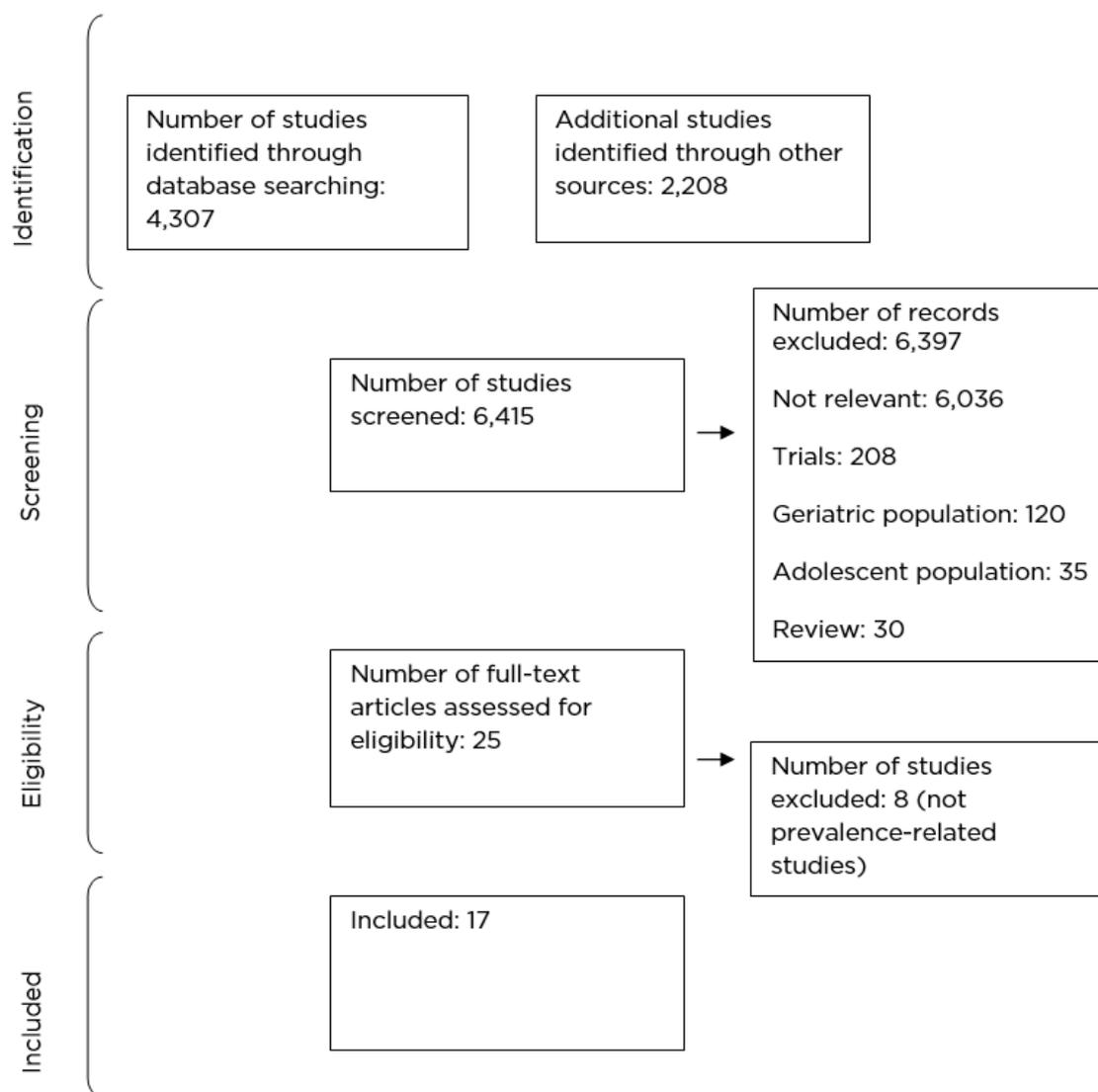


Figure 1: Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flow chart

Table 1: Characteristics of studies included in the review.

Author (year)	Methods			Prevalence	Remarks/socio-demographic correlation
	Design/site of data	Tools and questionnaires used for CMD	Method to diagnose NCD		
Shruthi et al. <sup>30</sup> (2018)	Cross-sectional study N=248 Hospital	MINI and HAM-D	WHO criteria of presence of typical myocardial ischaemic pain, electrocardiographic changes, and cardiac biomarkers	Diabetes: 67.30% Hypertension: 65.33% Depression: 44.00%	Nicotine consumption (smoke form) was found in 66.1% of patients and alcohol use was found in 56.0% of patients

Table 1 continued.

Balhara, Sagar <sup>31</sup> (2011)	Cross-sectional study N=77 Hospital	Brief patient health questionnaire, and HADS	HbA1c levels, fasting blood glucose, and postprandial blood glucose	Depression: 16.9 % Generalised anxiety disorder: 3.9%	HADS anxiety scores were found to be related to HbA1c levels (correlation-coefficient: 0.41; p=0.03) and postprandial blood glucose levels (correlation coefficient: 0.51; p=0.02).
Chaudhary et al. <sup>32</sup> (2017)	Cross-sectional study N=50 Hospital	HAM-D and HAM-A	N/A	Depression among patients with T1D: 38% Depression among patients with T2D: 42% Anxiety among patients with T1D: 44% Anxiety among patients with T2D: 34%	Depressive symptoms were found to be more prominent in patients with T2D compared with patients with T1D
John <sup>33</sup> (2013)	Cross-sectional observational study N=130 Hospital	MMSE or HMSE, SCID-I, and health-related quality of scale (EQ-5D)	ECG and cardiac enzyme studies	Depression: 34.6% Anxiety: 36.9%	Prevalence of dysthymia: 30.0% Panic disorder: 3.1% Generalised anxiety disorder: 3.8% Adjustment disorder: 7.7%
Joseph et al. <sup>34</sup> (2013)	Cross-sectional study N=230 Patients with T2D in hospital	PRIME-MD PHQ-9	Recordings of blood investigation and anthropometric measures	Depression: 45.2%	Categorisation of depression: moderate depression (30.9%) and severe depression (14.3%) Average duration of time since the detection of diabetes was 12.10±7.35 years
Kulkarni et al. <sup>35</sup> (2014)	Cross-sectional study N=282 Hospital	PHQ-SADS	Diagnosed by physician	Anxiety: 19.1% Depression: 29.1%	Prevalence of somatisation: 35.1%
Mathew et al. <sup>36</sup> (2012)	Cross-sectional study N=80 Hospital	MDI, BDI, and the DSM-IV	Diagnosed by physician	Depression: 38.8%	Multiple linear regression models showed that the presence of depression increased HbA1c by an average of 0.94% after adjusting for age and sex

Table 1 continued.

Poongothai et al. <sup>37</sup> (2015)	Cross-sectional study N=1,505 Hospital	PHQ-12	Anthropometric measures and blood investigation	Depression: 16.6%	N/A
Rajput et al. <sup>38</sup> (2016)	Cross-sectional case-control study design N=820 Hospital	HAM-D and HAM-A	Diagnosed by physician and anthropometric measurements were recorded	Depression in patients with diabetes: 26.3% Depression in patients without diabetes: 11.2% Anxiety in patients with diabetes: 27.6% Anxiety in patients without diabetes: 12.7% Comorbid depression and anxiety in patients with diabetes: 21.0%	N/A
Raval et al. <sup>39</sup> (2010)	Cross-sectional study N=80 Hospital	PHQ-9, self-report version of PRIME-MD	Diagnosed by physician and anthropometric measures and blood investigation	Depression: 41%	Depression was strongly associated with age
Singla et al. <sup>40</sup> (2012)	Cross-sectional study N=91 Hospital	MINI Neuropsychiatric Scale	Pre-diagnosed diabetes by physician, routine biochemistry	Depression: 8%	Family history of diabetes was present in 60.43% of patients, 6.59% were active smokers, and 46.15% were obese
Thour et al. <sup>41</sup> (2015)	Cross-sectional study N=73 Hospital	PHQ-9	Pre-diagnosed diabetes by physician	Depression: 41%	Categorisation: severe depression (4%); moderate depression (10%); mild depression (27%) Depression was significantly more prevalent in rural subjects (57%) when compared to urban ones (31%; p=0.049)
Kanwar et al. <sup>42</sup> (2019)	Cross-sectional study N=202 Hospital	Case record form, (Brief IPQ), ICD-10, HAM-D, HAM-A, and MINI	Routine blood investigation	Depression: 41.9% Other psychiatric comorbidity: 58.4%	Depression was higher in female patients and persons aged >50 years
Bhatt et al. <sup>43</sup> (2015)	Observational, single-centre study N=6,867 Hospital	MADRS and SF-36	N/A	Depression: 39.8%	N/A

Table 1 continued.

Weaver, Madhu <sup>44</sup> (2015)	Cross-sectional study N=184 Hospital	HSCL	Finger-stick blood test, pre-diagnosed diabetes by physician	Depression: 19% Anxiety: 26%	N/A
Guruprasad et al. <sup>45</sup> (2012)	Cross-sectional study N=210 Hospital	BDI	GPE, pre-diagnosed diabetes by physician	Depression: 27.6% Comorbid hypertension with depression and diabetes: 51.7% IHD: 25.9% Obesity and diabetes: 15.5% Tobacco use: 65.5%	N/A
Das et al. <sup>46</sup> (2013)	Cross-sectional study N=195 Hospital	HAM-D, Q-LES-Q, and SF	Diagnosed by physician	Depression: 46.15%	Categorisation: mild depression (32.2%); moderate depression (36.7%); severe depression (14.4%); very severe depression (16.7%)

BDI: Beck Depression Inventory; Brief IPQ: Brief Illness Perception Questionnaire; CMD: common mental disorders; DSM-IV: Diagnostic and Statistical Manual of Mental Disorders IV ed. 2000; GPE: General Physical Examination; HADS: Hospital Anxiety and Depression Scale; HAM-A Hamilton Anxiety Rating Scale; HAM-D: Hamilton Depression Rating Scale; HbA1c: glycated haemoglobin; HMSE: Hindi Mental Status Examination; HSCL: Hopkins Symptoms Checklist; ICD-10: International Classification Of Diseases, Tenth Revision; IHD: ischaemic heart disease; MADRS: Montgomery-Åsperg Depression Rating Scale; MDI: Major Depression Inventory; MINI: Mini International Neuropsychiatric Interview PLUS 5.0.0; MMSE: Mini Mental Status Examination; N/A: not applicable; NCD: non-communicable diseases; PHQ-SADS: Patient Health Questionnaire-Somatic, Anxiety, and Depressive Symptoms; PHQ-9: Patient Health Questionnaire-9; PHQ-12: Patient Health Questionnaire-12; PRIME-MD: Primary Care Evaluation of Mental Disorders; Q-LES-Q: Quality of Life Enjoyment and Satisfaction Questionnaire; SCID-I: structured clinical interview for DSM-IV Axis I Diagnosis; SF: Short Form; SF-36: Short Form Health Survey-36; T1D: Type 1 diabetes; T2D: Type 2 diabetes; WHO: World Health Organization.

## REVIEW OF THE INCLUDED STUDIES

All the studies included in this review were conducted in hospital settings and were observational studies. Out of the 17 studies, only one had respondents enrolled from an inpatient setting. Overall, 12 studies assessed the prevalence of depression and/or anxiety in NCDs, particularly in diabetes. Three studies evaluated the prevalence of depression in CVD. Only one study looked into the prevalence of CMD in NCD (diabetes, hypertension, and CVD).

Despite the aetiological differences, many studies have not distinguished between different types of diabetes. Only one study focused on

Type 1 diabetes (T1D) and Type 2 diabetes (T2D) separately;<sup>32</sup> 11 studies have focused only on T2D.

### Studies That Assessed Depression and Anxiety Disorders in Diabetes

The prevalence of depression in patients with diabetes ranged from 8% to 46%. Risk factors of NCDs such as smoking and alcohol use were also reported in three studies. The sample size varied from 50 to 10,450. Depression was identified by diagnostic interview: International Classification of Diseases (ICD) and Diagnostic and Statistical Manual of Mental Disorder (DSM-IV) in two studies; Physical Health Questionnaire-9 (PHQ-9) in five studies; Hamilton Rating Scale for Depression-17

(HAM-D-17) in three studies; Hamilton Depression Rating Scale (HDRS-21) in one study; Beck Depression Inventory (BDI) in one study; Primary Care Evaluation of Mental Disorders (PRIME-D) in one study; and Hopkins Symptom Checklist (HSCL) in one study. Studies were from Northern (10), Southern (six), and Western (two) regions of India. The prevalence of anxiety disorder and depression in any NCD were 3.9–44.0% and 8.0–46.0%, respectively.

Four studies reported an association between the duration of diabetes and the prevalence of depression, as the incidence of diabetic complications increased with illness duration.<sup>31,41,42,45</sup> A higher prevalence of depression was reported in one of the studies among patients having uncontrolled diabetes (glycated haemoglobin test: HbA1c).<sup>45</sup> Two studies concluded that patients diagnosed with T2D and depression had poor QoL compared with those without depression. Health-related QoL was adversely affected by the presence of depression.<sup>46</sup> Negative correlation was reported between HbA1c and QoL in one of the studies. This supports the view that patients with poor control of blood glucose levels have a worse QoL. One of the studies reported that 67% of patients with diabetes were overweight or obese.<sup>45</sup> Two studies indicated that depression was significantly associated with older age,<sup>34,39</sup> female gender,<sup>34</sup> and complications due to T2D or other comorbidities such as hypertension and being overweight.<sup>34,38,39</sup> Conversely, two studies reported that gender was not associated with depression.<sup>38,39</sup> One of the studies indicated that depression was more prevalent in rural populations than urban.<sup>41</sup> One of the studies showed that depression was significantly associated with retinopathy, nephropathy, and IHD.<sup>38</sup> Another study reported that depression was strongly associated with age, central obesity, neuropathy, and nephropathy; however, it also found that depression was not significantly associated with gender.<sup>39</sup>

### Studies That Assess Depression and Anxiety Disorders in Hypertension

Out of 17 studies, only one, conducted in a hospital in Mangalore, India, reported a prevalence of depression in patients with hypertension as 28%.<sup>35</sup>

### Studies That Assess Depression and Anxiety Disorders in Cardiovascular Diseases

Among the four studies that assessed the prevalence of depression in CVD, one focused on IHD in general, two focused on acute coronary artery disease, and one focused on carotid artery intima-media thickness. One study assessed the relationship between IHD and QoL. Poor QoL was reported among patients with IHD and comorbid anxiety and depression.<sup>33</sup> In one of the studies, depressive symptoms were associated with functional and structural markers of atherosclerosis, common carotid intima-media thickness, and augmentation index.<sup>37</sup>

## DISCUSSION

In this review, the authors have presented a comprehensive summary of existing research on the comorbidity of CMDs (anxiety and depression) and NCDs (hypertension, CVD, and diabetes) in India. The data presented in this review reflects the high prevalence of anxiety and depression in NCDs, specifically diabetes. The prevalence of depression in patients with diabetes in the present study was 8–46%. Studies that were based in the USA and the Netherlands, which were included in a systematic review on the epidemiology of diabetes and depression, conducted among patients with diabetes, reported the prevalence rate of depression as 32.1% and 19.3%, respectively. Though these are from high-income countries, they are within the range of the present study.<sup>47</sup> The estimated prevalence of depression in patients with hypertension in one of the meta-analyses was reported to be 21.3%, and the range of point prevalence of 41 studies was observed to be between 0.5% and 73%.<sup>48</sup>

Twelve studies included in the review examined the prevalence of CMDs in diabetes. Diabetes and psychiatric disorders share a bidirectional association, both influencing each other in multiple ways.<sup>49</sup> People with diabetes are two-times more likely to develop depression relative to people without depression, and patients with depression have poorer clinical parameters and outcomes of diabetes because of poor medication compliance and dietary practices.<sup>50</sup> It has also been reported that comorbid anxiety and depression in people with diabetes also

leads to medical complications of diabetes, work disability, and poor QoL.<sup>51</sup> Patients with diabetes are likely to face challenges at physical, emotional, psychological, social, occupational, and interpersonal levels.<sup>52</sup> The complications associated with the management (medication dosing, monitoring blood glucose, eating patterns, and physical activity)<sup>53</sup> of diabetes can lead to negative emotions such as worry, fear, and anger, which result in poor QoL.<sup>50</sup> Among the studies reviewed, 11 have focused on T2D despite the aetiological differences. T1D is an autoimmune condition, whereas T2D is linked with genetic and lifestyle choices.<sup>54</sup> Both types of diabetes lead to health complications. Depressive symptoms seem to be slightly more prevalent in T2D compared with T1D.<sup>11</sup>

The co-occurrence of NCDs and CMDs has been associated with elevated symptom burden, functional impairment, and a decrease in QoL.<sup>9</sup> The authors found that the prevalence of anxiety and depression was higher in females compared with males, similar to the studies done elsewhere.<sup>55</sup> Depression was identified either by diagnostic interviews or diagnosis by a general practitioner (n=2). The tools used in the studies were PHQ-9, HAM-D, and BDI, which are widely used in clinical practice and research. PHQ is a screening tool commonly used in primary care, HAM-D is a widely used clinician-administered depression assessment scale, and BDI is a self-reported inventory to measure the severity of depression. It is reported that the estimate of depression prevalence depends upon the assessment tools (standardised interview or self-report questionnaires, classification of depression, and diabetes type).<sup>11</sup> Depression assessed with self-report questionnaires is found at a rate two-times higher than when assessed with standardised interviews.<sup>17,53</sup> The overall quality of the studies included in the systematic review was good. In the risk of bias assessment, most studies were categorised as good (n=14). All the studies used standardised tools that were reliable and valid for assessment.

CMDs, particularly depression and anxiety, often go undetected, and systematic screening, even in high-income countries, is not satisfactory.<sup>9</sup> Thus, the screening of CMDs in primary care by using a brief instrument should be made mandatory in routine clinical encounters. This integration has also been reported as an evidence-based approach for collaborative care.<sup>16</sup>

Addressing the overlapping risk factors and integrating the management of CMDs and NCDs can improve clinical outcomes.<sup>56</sup> Management of NCDs and CMDs can be made by adopting existing integrated models of care. The inter-related causal mechanism of CMD and NCD argue for an integrated approach to care.<sup>16</sup> The management of T2D and depression in the primary setting has been effective using the educative model, where healthcare professionals are trained to identify, refer, and communicate with the patient about the symptoms of CMD.<sup>57</sup> The efficiencies of integrated care have proven beneficial for both high-income and low- and middle-income countries.<sup>16</sup>

Several policies and programmes have been adopted in India to tackle the rising burden of NCDs and mental illnesses.<sup>58</sup> These include the National Programme for Prevention and Control of Cancer, Diabetes, Cardiovascular Diseases and Stroke (NPCDCS) and District Mental Health Programme (DMHP). It has been reported that the major challenges are in terms of inadequate resources and ineffective implementation of the programmes.<sup>58</sup> To overcome these challenges, health and wellness centres have been envisaged under the Ayushman Bharat programme of the Government of India, encompassing a range of services for NCDs and mental health closer to the community.<sup>59</sup> Health and wellness centres focus on wellness and lifestyle modification through various activities such as yoga, local sports, and other activities. The screening of NCDs is done by accredited social health activists or auxiliary nurse midwives through the Community Based Assessment Checklist (CBAC). Identified individuals are then referred for necessary treatment. According to the recent statistics, approximately 9.1 crore screenings for hypertension, and 7.4 crore screenings for diabetes have been carried out.<sup>60</sup>

In addition, for effective management, social workers, accredited social health activists, auxiliary nurse midwives, and NCD counsellors can be trained for population-based screening of CMDs in NCDs and vice versa, basic counselling skills, brief intervention for lifestyle modification, and behavioural activation.

## STRENGTHS AND LIMITATIONS

The search strategy adopted was extensive and comprehensive in nature. The studies were limited to those conducted in India. This review included only the prevalence of anxiety and depression in NCDs and excluded other CMDs (somatisation) and substance use disorders. The pooled prevalence of anxiety and depression has not been provided because meta-analysis was not conducted. Small sample sizes in some of the studies may have contributed to sampling errors. All the studies were hospital-based and, therefore, the findings are not generalisable to the community. There were insufficient data to draw firm conclusions about the prevalence of anxiety and depression in hypertension and CVD, as only three studies evaluated the prevalence of depression in CVD and only one study examined the prevalence of CMD in hypertension. The broad range of prevalence can be explained by the following four factors: use of different tools to screen and assess anxiety and/or depression;

the studies were conducted in geographic or cultural settings; sample sizes were disparate; and true variation across the different conditions that have been included as NCDs.

## CONCLUSIONS

In summary, the authors found that NCDs and CMDs are underdiagnosed in India. The data related to the prevalence rates of both NCDs and CMDs is limited. Based on studies carried out in India, the prevalence of anxiety disorder and depression in individuals with NCD was 3.9–44.0% and 8.0–46.0%, respectively. The results of the included studies point to disadvantages for people with CMDs and NCDs with respect to risk factors and outcome. Depressive and anxiety symptoms have been strongly associated with poor QoL, health status, and physical function. Thus, future efforts should focus on integrated healthcare, which may help improve screening and identification of NCDs and CMDs; this in turn can improve outcomes.

## References

1. Bloom DE et al. The global economic burden of noncommunicable diseases. Program on the global demography of aging. 2012. Available at: <https://ideas.repec.org/p/gdm/wpaper/8712.html>. Last accessed: 14 February 2022.
2. Appleby L et al. Sudden unexplained death in psychiatric in-patients. *Br J Psychiatry*. 2000;176(5):405-6.
3. World Health Organization (WHO). Noncommunicable diseases country profiles. 2018. Available at: <https://apps.who.int/iris/handle/10665/274512>. Last accessed: 14 February 2022.
4. World Health Organization (WHO). Global summit report on noncommunicable diseases. 2010. Available at: [https://www.who.int/nmh/publications/ncd\\_report\\_full\\_en.pdf](https://www.who.int/nmh/publications/ncd_report_full_en.pdf). Last accessed: 14 February 2022.
5. GBD 2015 Risk Factors Collaborators. Global, regional, and national comparative risk assessment of 79 behavioural, environmental and occupational, and metabolic risks or clusters of risks, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015. *Lancet*. 2016;388(10053):1659-724.
6. Fryers T et al. Social inequalities and the common mental disorders: a systematic review of the evidence. *Soc Psychiatry and Psychiatr Epidemiol*. 2003;38(5):229-37.
7. Görlitz N et al. [Prevalence of complications and comorbidities in Type 2 diabetes: a cross-sectional study of disease management program participants in Bavaria]. *Dtsch Med Wochenschr*. 2008;133(33):1667-72. (In German).
8. Renn BN et al. The bidirectional relationship of depression and diabetes: a systematic review. *Clin Psychol Rev*. 2011;31(8):1239-46.
9. Goodell S et al. Mental disorders and medical comorbidity. 2011. Available at: <https://www.rwjf.org/en/library/research/2011/02/mental-disorders-and-medical-comorbidity.html>. Last accessed: 14 February 2022.
10. Mezuk B et al. Depression and Type 2 diabetes over the lifespan: a meta-analysis. *Diabetes Care*. 2008;31(12):2383-90.
11. Andreoulakis E et al. Depression in diabetes mellitus: a comprehensive review. *Hippokratia*. 2012;16(3):205-14.
12. O'Neil A et al. A shared framework for the common mental disorders and non-communicable disease: key considerations for disease prevention and control. *BMC Psychiatry*. 2015;DOI:10.1186/s12888-015-0394-0.
13. Berge LI, Riise T. Comorbidity between Type 2 diabetes and depression in the adult population: directions of the association and its possible pathophysiological mechanisms. *Int J Endocrinol*. 2015;2015:164760.
14. Goodwin GM. Depression and associated physical diseases and symptoms. *Dialogues Clin Neurosci*. 2006;8(2):259-65.
15. Katon W et al. Diabetes and poor disease control: is comorbid depression associated with poor medication adherence or lack of treatment intensification? *Psychosom Med*. 2009;71(9):965-72.
16. Stein DJ et al. Integrating mental health with other non-communicable diseases. *BMJ*. 2019;364:l295.
17. Anderson RJ et al. The prevalence of comorbid depression in adults with diabetes: a meta-analysis. *Diabetes Care*. 2001;24(6):1069-78.
18. Bădescu S et al. The association between diabetes mellitus and depression. *J Med Life*. 2016;9(2):120-5.
19. Rutledge T, Hogan BE. A quantitative review of prospective evidence linking psychological factors with hypertension development. *Psychosom Med*. 2002;64(5):758-66.

20. Johannessen L et al. Increased risk of hypertension in patients with bipolar disorder and patients with anxiety compared to background population and patients with schizophrenia. *J Affect Disord.* 2006;95(1-3):13-7.
21. Pan Y et al. Association between anxiety and hypertension: a systematic review and meta-analysis of epidemiological studies. *Neuropsychiatr Dis Treat.* 2015;11:1121-30.
22. Rubio-Guerra AF et al. Depression increases the risk for uncontrolled hypertension. *Exp Clin Cardiol.* 2013;18(1):10-2.
23. Meng L et al. Depression increases the risk of hypertension incidence: a meta-analysis of prospective cohort studies. *J Hypertens.* 2012;30(5):842-51.
24. Player MS, Peterson LE. Anxiety disorders, hypertension, and cardiovascular risk: a review. *Int J Psychiatry Med.* 2011;41(4):365-77.
25. Hare DL et al. Depression and cardiovascular disease: a clinical review. *Eur Heart J.* 2014;35(21):1365-72.
26. Uphoff EP et al. A systematic review and meta-analysis of the prevalence of common mental disorders in people with non-communicable diseases in Bangladesh, India, and Pakistan. *J Glob Health.* 2019;9(2):020417.
27. Dandona L et al.; India State-Level Disease Burden Initiative Collaborators. Nations within a nation: variations in epidemiological transition across the states of India, 1990–2016 in the Global Burden of Disease Study. *Lancet.* 2017;390(10111):2437-60.
28. Guariguata L et al. Global estimates of diabetes prevalence for 2013 and projections for 2035. *Diabetes Res Clin Pract.* 2014;103(2):137-49.
29. National Heart, Lung and Blood Institute (NHLBI). National Institutes of Health Quality assessment tool for observational cohort and cross-sectional studies. 2021. Available at: <https://www.nhlbi.nih.gov/health-topics/study-quality-assessment-tools>. Last accessed: 14 February 2022.
30. Shruthi DR et al. Psychiatric comorbidities in acute coronary syndromes: six-month follow-up study. *Indian J Psychiatry.* 2018;60(1):60-4.
31. Balhara YP, Sagar R. Correlates of anxiety and depression among patients with Type 2 diabetes mellitus. *Indian J Endocrinol Metab.* 2011;15(Suppl 1):S50-4.
32. Chaudhary R et al. Comparative study of psychiatric manifestations among Type I and Type II diabetic patients. *Indian J Psychol Med.* 2017;39(3):342-6.
33. John S. Prevalence and pattern of psychiatric morbidity and health related quality of life in patients with ischemic heart disease in a tertiary care hospital. *Indian J Psychiatry.* 2013;55(4):353-9.
34. Joseph N et al. Proportion of depression and its determinants among Type 2 diabetes mellitus patients in various tertiary care hospitals in Mangalore city of South India. *Indian J Endocrinol Metab.* 2013;17(4):681-8.
35. Kulkarni V et al. Psychiatric comorbidities among patients with select non-communicable diseases in a coastal city of South India. *Int J Prev Med.* 2014;5(9):1139-45.
36. Mathew CS et al. Prevalence of depression in consecutive patients with Type 2 diabetes mellitus of 5-year duration and its impact on glycemic control. *Indian J Endocrinol Metab.* 2012;16(5):764-8.
37. Poongothai S et al. Association of depression with common carotid artery intima media thickness and augmentation index in a large Urban South Indian population- The Chennai Urban Rural Epidemiology Study (CURES - 138). *Indian J Endocrinol Metab.* 2015;19(1):136-42.
38. Rajput R et al. Prevalence and predictors of depression and anxiety in patients of diabetes mellitus in a tertiary care center. *Indian J Endocrinol Metab.* 2016;20(6):746-51.
39. Raval A et al. Prevalence and determinants of depression in Type 2 diabetes patients in a tertiary care centre. *Indian J Med Res.* 2010;132:195-200.
40. Singla R et al. Profile of Type 2 diabetes mellitus without overt complications of diabetes mellitus at a tertiary care center. *Indian J Endocrinol Metab.* 2012;16(Suppl 2):S468-70.
41. Thour A et al. Depression among patients with diabetes mellitus in North India evaluated using patient health questionnaire-9. *Indian J Endocrinol Metab.* 2015;19(2):252-5.
42. Kanwar N et al. Prevalence of psychiatric comorbidity among patients of Type 2 diabetes mellitus in a hilly state of North India. *Indian J Endocrinol Metab.* 2019;23(6):602-8.
43. Bhatt P et al. Unique aspects of coronary artery disease in Indian women. *Cardiovasc Drugs Ther.* 2015;29(4):369-76.
44. Weaver LJ, Madhu S. Type 2 diabetes and anxiety symptoms among women in New Delhi, India. *Am J Public Health.* 2015;105(11):2335-40.
45. Guruprasad K et al. A study of association of depressive symptoms among the Type 2 diabetic outpatients presenting to a tertiary care hospital. *Indian J Psychol Med.* 2012;34(1):30-3.
46. Das R et al. Prevalence of depression in patients with Type II diabetes mellitus and its impact on quality of life. *Indian J Psychol Med.* 2013;35(3):284-9.
47. Roy T, Lloyd CE. Epidemiology of depression and diabetes: a systematic review. *J Affect Disord.* 2012;142:S8-21.
48. Li Z et al. Prevalence of depression in patients with hypertension: a systematic review and meta-analysis. *Medicine (Baltimore).* 2015;94(31):e1317.
49. Singh Balhara YP, Kalra S. Psychiatric disorders in diabetes. *J Pak Med Assoc.* 2015;65(10):1137-8.
50. Bhat NA et al. The psychological aspects of diabetes. *EMJ Diabet.* 2020;8(1)90-8.
51. Pouwer F et al. Rates and risks for co-morbid depression in patients with Type 2 diabetes mellitus: results from a community-based study. *Diabetologia.* 2003;46(7):892-8.
52. Kalra S et al. Emotional and psychological needs of people with diabetes. *Indian J Endocrinol Metab.* 2018;22(5):696-704.
53. Fisher L et al. When is diabetes distress clinically meaningful?: establishing cut points for the Diabetes Distress Scale. *Diabetes Care.* 2012;35(2):259-64.
54. Ozougwu J et al. The pathogenesis and pathophysiology of Type 1 and Type 2 diabetes mellitus. *J Physiol Pathophysiol.* 2013;4(4):46-57.
55. Grigsby AB et al. Prevalence of anxiety in adults with diabetes: a systematic review. *J Psychosom Res.* 2002;53(6):1053-60.
56. Das J et al. Mental health patterns and consequences: results from survey data in five developing countries. *World Bank Econ Rev.* 2009;23(1):31-55.
57. Katon W et al. Collaborative management to achieve treatment guidelines: impact on depression in primary care. *JAMA.* 1995;273(13):1026-31.
58. Sivanantham P et al. Profile of risk factors for non-communicable diseases (NCDs) in a highly urbanized district of India: findings from Puducherry district-wide STEPS survey, 2019–20. *PLoS One.* 2021;16(1):e0245254.
59. Bakshi H et al. Ayushman Bharat Initiative (2018): what we stand to gain or lose! *Indian J Community Med.* 2018;43(2):63-6.
60. Department of Health and Family Welfare (MoHFW). 2020-21 annual report. 2021. Available at: <https://main.mohfw.gov.in/sites/default/files/Annual%20Report%202020-21%20English.pdf>. Last accessed: 14 February 2022.