Guidelines for Adolescent Depression in Primary Care (GLAD-PC): I. Identification, Assessment, and Initial Management
Rachel A. Zuckerbrot, Amy H. Cheung, Peter S. Jensen, Ruth E.K. Stein and Danielle Laraque

*Pediatrics* 2007;120;e1299
DOI: 10.1542/peds.2007-1144

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://pediatrics.aappublications.org/content/120/5/e1299.full.html
Guidelines for Adolescent Depression in Primary Care (GLAD-PC): I. Identification, Assessment, and Initial Management

Rachel A. Zuckerbrot, MD, Amy H. Cheung, MD, Peter S. Jensen, MD, Ruth E. K. Stein, MD, Danielle Laraque, MD, and the GLAD-PC Steering Group

ABSTRACT

OBJECTIVES. To develop clinical practice guidelines to assist primary care clinicians in the management of adolescent depression. This first part of the guidelines addresses identification, assessment, and initial management of adolescent depression in primary care settings.

METHODS. By using a combination of evidence- and consensus-based methodologies, guidelines were developed by an expert steering committee in 5 phases, as informed by (1) current scientific evidence (published and unpublished), (2) a series of focus groups, (3) a formal survey, (4) an expert consensus workshop, and (5) draft revision and iteration among members of the steering committee.

RESULTS. Guidelines were developed for youth aged 10 to 21 years and correspond to initial phases of adolescent depression management in primary care, including identification of at-risk youth, assessment and diagnosis, and initial management. The strength of each recommendation and its evidence base are summarized. The identification, assessment, and initial management section of the guidelines includes recommendations for (1) identification of depression in youth at high risk, (2) systematic assessment procedures using reliable depression scales, patient and caregiver interviews, and Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition criteria, (3) patient and family psychoeducation, (4) establishing relevant links in the community, and (5) the establishment of a safety plan.

CONCLUSIONS. This part of the guidelines is intended to assist primary care clinicians in the identification and initial management of depressed adolescents in an era of great clinical need and a shortage of mental health specialists but cannot replace clinical judgment; these guidelines are not meant to be the sole source of guidance for adolescent depression management. Additional research that addresses the identification and initial management of depressed youth in primary care is needed, including empirical testing of these guidelines.
MAJOR DEPRESSION IN adolescents is recognized as a serious psychiatric illness with extensive acute and chronic morbidity and mortality. Research shows that only 50% of adolescents with depression are diagnosed before they reach adulthood. In primary care (PC), as many as 2 in 3 depressed youth are not identified by their PC clinicians and do not receive any kind of care. Even when diagnosed by PC physicians, only half of these patients are treated appropriately. Furthermore, rates of completion of specialty mental health referral for youth with a recognized emotional disorder from general medical settings are quite low.

In view of the shortage of mental health clinicians and barriers to children having access to mental health professionals, the well-documented need for PC clinicians to learn how to manage this condition, the increasing evidence base available to guide clinical practice, increased selective serotonin reuptake inhibitor–prescribing rates in pediatric PC, and new evidence that a multifaceted approach with mental health consultation may improve the management of depression in PC settings, guidelines may be a necessary first step in the identification and management of depression in adolescents in PC. Unfortunately, no depression-management guidelines have been developed for use in the PC setting in the United States or Canada.

Although additional randomized, controlled, clinical trial (RCT) information is urgently needed to guide PC clinicians in optimal management approaches, such studies often take years to complete, and many critical PC adolescent depression-management questions have not been, and will likely never be, addressed in completed or ongoing studies. To address this gap and meet the needs of PC clinicians and families who are on the “front lines” with few mental health resources available, this report and its companion article constitute the first-ever evidence- and expert consensus–derived guidelines for use in the PC setting in the United States or Canada.

In this article, we present the summary result of literature reviews of the available data and the recommendations on the identification and assessment of depression in PC settings; in our accompanying report, we present the results of the reviews and recommendations on treatment (psychotherapy, psychopharmacology, and pediatric counseling) and ongoing management. Although very few studies have addressed adolescent depression identification and management in PC settings, many PC clinicians are already attempting to change their clinical practices; thus, a great need exists to develop and disseminate methods and tools for assisting PC clinicians in managing adolescent depression.

Major depressive disorder (MDD) is a specific diagnosis described in the Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV), which includes symptoms of low mood, anhedonia, and other neurovegetative symptoms (ie, insomnia, decreased concentration, low energy, etc). Other types of depression exist, including dysthymia, subthreshold forms, or those that occur as part of bipolar disorder or other mental illness. Although the evidence for the psychopharmacology recommendations in the accompanying article focus exclusively on MDD, the recommendations around identification, assessment, and initial management can be applied to other forms of depression as well.

Our guidelines also distinguish between mild, moderate, and severe forms of MDD. The DSM-IV depression criteria include 9 specific symptoms that have been shown to cluster together, run in families, and have a genetic basis; a large body of evidence accumulated over time now supports the internal consistency of depressive symptoms and the validity of the major depression construct, based on the validation criteria for all psychiatric diagnoses. According to the DSM-IV, severity of depressive disorders can be based on symptom count. This commonly used method to define depression severity has been used in large population-based studies and may be particularly relevant in PC settings in which less-severe clinical presentations of depression may be more common. Thus, mild depression may be characterized on the basis of lower scores in standardized depression scales with shorter duration of symptoms or meeting minimal criteria for depression. Following the DSM-IV, mild depression might be defined as 5 to 6 symptoms that are mild in severity. Furthermore, the patient might experience only mild impairment in functioning.

In contrast, depression might be deemed to be severe if a patient experiences all of the depressive symptoms listed in the DSM-IV. Depression might also be considered severe if the patient experiences severe impairment in functioning. Moderate depression falls between these 2 categories. In general, however, even if not all 9 DSM-IV–defined symptoms of depression are present, for the purposes of these guidelines, an adolescent who meets at
least 5 criteria for the diagnosis of MDD should be considered to be in the severe category if he or she presents with a specific suicide plan, clear intent, or recent attempt; psychotic symptoms; or severe impairment in functioning (such as being unable to leave home).

These guidelines were developed for PC clinicians who are in a position to identify and assist depressed youth in their practice settings. Although the age range of 10 to 21 years may encompass preteens, adolescents, and young adults in specific instances, this age range was chosen to include those who might be developmentally “adolescent.” Research that supports adult depression guidelines includes adults 18 years and older. Much of the adolescent depression research focuses on children 18 years and younger. However, because adolescent medicine clinicians and school health clinicians often see patients until they are 21 years old, we have included the older adolescents. However, a primary caregiver faced with an adolescent between the ages of 18 and 21 can choose to use either adult or adolescent depression guidelines on the basis of the developmental status of the adolescent and their own comfort and familiarity with each set of guidelines.

METHODS
The following recommendations were developed on the basis of a synthesis of expert consensus- and evidence-based research-review methodologies. We compiled the necessary information to develop these recommendations in 5 phases.

1. To understand the problems and obstacles faced by PC clinicians regarding the management of adolescent depression, we first conducted focus groups with PC clinicians and youth patients and their family members to review issues pertinent to the PC management of depression.

2. Systematic literature reviews were conducted in each of 5 key areas in which recommendations were subsequently developed. Whenever possible, these reviews focused on identifying empirical evidence that was developed within child/adolescent PC settings. When PC studies were unavailable, research from specialty mental health care was reviewed. In all 5 review instances, the GLAD-PC Steering Group first determined the existence of all high-quality, previously published, systematic evidence-based reviews that met the following criteria: (a) clear definition of search terms from Medline, including words and word roots; (b) explicit delineation of years searched; (c) exclusion of non–English-language studies; (d) physical review and reading of search-identified titles and abstracts; and (e) selection, review, and reading of possibly relevant articles before determination of final inclusion. When more than 1 systematic evidence-based review was identified for a given area, all reviews were drawn on to identify relevant articles for potential inclusion. More than 1 systematic evidence-based review was available for the areas of efficacy of psychotherapeutic interventions for youth MDD and efficacy of pharmacologic treatments for youth MDD. For all reviews, when appropriate, we updated the review for any ensuing years transpired since the latest review by using these same 5 methods. When systematic reviews were not available for a given area, the GLAD-PC team conducted a systematic review by using Medline (from inception to 2004/2005) and the criteria described above. Reviews were guided by members of the GLAD-PC Steering Committee, which comprised leading experts in each of these areas.

To address the first key area regarding the identification and assessment of adolescent depression in PC, a systematic evidence review was conducted to identify all available evidence about adolescent depression identification in PC, as well as information regarding current practices. This review has since been published.20 Because of limited information about depression assessment and screening measures in PC specifically, we also reviewed adolescent-screening instruments/tools previously used in psychiatric or community populations. Beginning from 2 previous systematic evidence reviews,21,22 the GLAD-PC team performed an additional systematic review from 1998 to 2004.

To address the second key area regarding the initial management of adolescent depression in PC, a systematic evidence review was conducted to identify all available evidence about interventions for adolescent depression in PC and has since been published as well.23 Other evidence for the initial management of adolescent depression in PC came from systematic evidence reviews that addressed the chronic illness model, systems of care, and safety planning for suicidal patients.

3. On the basis of the questions and issues identified during the focus groups and the literature reviews, we developed a survey to answer questions regarding critical issues in PC management of adolescent depression that have not been answered in the empirical literature. The survey questions were developed and reviewed by clinical and research experts in the area of mental health and PC. Using this survey, research and clinical experts were surveyed on their depression assessment and management recommendations. Depression clinical/research experts (N = 81) from Canada and the United States were asked to complete the 34-item study survey. Of these items, 3 questions dealt with the identification and diagnosis of depression. Subjects were chosen by using 1 of 4 criteria: (a) membership in child and adolescent psychiatric organizations in Canada and the United States including their academies of child and adoles-
cent psychiatry; (b) recipient of federal grants for related research; (c) lead author of at least 2 articles on clinical research in the area from 1999 to 2004 on the basis of Medline citations; or (d) key PC clinical and research leader with expertise in the area of guideline development and/or emotional and behavioral disorders that present in PC settings. Complete survey results (n = 76) will be presented in a subsequent peer-reviewed article and are available from the authors on request.24

4. An expert consensus workshop was held in July 2004 with 81 North American experts on depression, clinical pediatrics, quality improvement, mental health policy, and health economics. Published data from the literature review, unpublished high-quality research currently in process of publication, and the results of the survey were presented to guide the initial discussion and consensus process.

5. Guidelines were developed on the basis of multiple iterations shared among a small group of core writers, guidance of the larger steering committee, and ultimate input of all consensus-conference attendees to obtain full ownership of the final product. The results of this process are presented below.

On the basis of the 5-step method, 2 guidelines were developed to address different areas of adolescent depression management in PC settings: (1) identification, assessment, and initial management and (2) treatment and ongoing management. This part of the guidelines focuses on identification, assessment, and initial management. Each section of both guidelines is composed of individual recommendations followed by a brief rationale that refers to available empirical findings and experts’ consensus opinion on which the recommendations were based. Each recommendation is graded on the basis of the Oxford Centre for Evidence-Based Medicine grade of evidence (A–D) system (see www.cebm.net/levels.of.evidence.asp). In addition, the strength of each recommendation, in terms of the extent to which experts agreed that the recommendation is highly appropriate and a “first-line” practice, was reached for each recommendation. Recommendation strength was rated in 4 categories: very strong (>90% agreement), strong (>70% agreement), fair (>50% agreement), and weak (<50% agreement). The recommendations in the guidelines were developed only in areas of management that had at least “strong agreement” among experts.

RESULTS

Literature Reviews: Identification and Assessment

Twenty-five articles were located that discussed specific identification methods for adolescent depression in pediatric PC. Only 10 of these articles presented psychometric data of any kind, such as sensitivity, specificity, positive predictive value, negative predictive value, and area under the curve (a full table is available in the published review20). One of the 25, plus an additional 5 survey articles, dealt with current identification practices by PC physicians. According to those data, most PC clinicians rely on the use of presenting complaints and family concerns to identify depressed youth.25 Likewise, other surveys confirmed that very few pediatric providers have instituted a systematic assessment or case-finding tool to identify adolescent depression.26,27

Despite clinicians’ principal reliance on adolescent and parental chief complaints and physician interview in current practice, the authors of the review found that the use of these methods alone underidentify adolescent depression.20 Even physicians who are trained in the use of mnemonics to guide interviews underidentify adolescent depression.28 Instead, only by asking patients directly about depression and suicide (versus relying on them to volunteer the information) does one reliably improve case finding and the psychometric quality of diagnostic data.29–32 Using systematic assessment methods with depression-specific questions seems to provide the best identification results.27–29,31

Many steps are involved in implementing systematic depression-identification procedures within a busy practice setting (eg, training office staff in the use of procedures, adjusting other paperwork demands to fit the depression-identification procedures, ensuring that providers review the information, teaching providers how to use the information, and determining whether the procedures actually benefit youth). Although no study has documented the feasibility and outcomes of taking all of these steps within a single study, evidence from the review cited above as well as a recently published pilot study,34 suggests that each of these steps can be implemented in real-world settings (ie, training office staff27,28,30,33–37, ensuring PC providers review any results27,28,32–34,37, and establishing that appropriate counseling and/or needed mental health referrals are made33,39). Finally, and perhaps most importantly, although no study compared outcomes between screened and not-screened groups, 1 study did demonstrate that an identification program in PC when combined with high-quality depression treatment actually yields better outcomes than treatment-as-usual conditions4 (when no high-quality depression treatment is available).

Because of limited information about depression assessment and screening measures in PC specifically, we also reviewed adolescent-screening instruments/tools previously used in psychiatric or community populations. Beginning from 2 previous systematic evidence reviews,21,22 the GLAD-PC team performed an additional systematic review for the years 1998–2004 and found 15 studies with psychometric performance data on depression instruments in English-speaking adolescents. A table that presents these 15 studies, the gold-standard
diagnostic assessment used, sensitivity, specificity, positive predictive value, negative predictive value, prevalence, and study population is available on request. Commonly used adolescent-screening instruments included the Beck Depression Inventory, the Reynolds Adolescent Depression Screen, and the Mood and Feelings Questionnaire. Sensitivities of these more common instruments ranged from .70 to .90, and specificities ranged from .39 to .90. In addition, the Kutcher Adolescent Depression Scale, a newer 6-item instrument, had sensitivity and specificity of .81 and .86, respectively, in a school population with a positive predictive value of .39 and a negative predictive value of .98 given a 10% prevalence.

In summary, no perfect depression screening/assessment tool exists, but a number of adolescent depression assessment instruments do possess adequate psychometric properties to commend their use in depression detection and assessment. Thus, it is reasonable to expect that depression detection in PC can be improved by the use of self- or parent-report checklists. Reliance on adolescent self-report depression checklists alone will lead to substantial numbers of false-positive and false-negative cases. Instead, optimal diagnostic procedures should combine the use of depression-specific screening tools as diagnostic aids buttressed by follow-up clinical interviews in which one obtains information from other informants (eg, parents) and reconciles discrepant information to arrive at an accurate diagnosis and impairment assessment before treatment. For more information about rating scales and cutoff scores, please refer to the GLAD-PC toolkit (available at www.glad- pc.org; see Appendix).

Initial Management of Adolescent Depression

A GLAD-PC review of adolescent depression interventions performed in PC found 4 articles that directly discussed interventions for adolescent depression in PC, only 1 of which reported an intervention by a PC staff member as opposed to a mental health worker. This identified RCT evaluated the effectiveness of a PC-delivered consultation intervention that invited teens from 8 general practices in Britain to discuss their health concerns with a PC nurse who provided individual consultations. In addition to discussing adolescents' general health concerns, the PC nurses offered mental health referrals when they deemed it appropriate. Posthoc analyses indicated that among teens with high Center for Epidemiological Studies Depression Scale for Children (CES-DC) scores, those who were randomly assigned to the PC nurse consultation had lower CES-DC scores on follow-up than adolescents with high CES-DC scores who were not randomly assigned to the consultation, which suggests that a PC-delivered intervention may be useful in addressing depression symptoms and/or similar emotional issues (because these patients were not diagnosed with a depressive disorder per se), perhaps by the consultation itself or by the resulting referral to mental health.

Although the study discussed above was the only one that dealt with an intervention by a PC provider for adolescent depression, the large multisite trial conducted by Asarnow et al showed that improving links between PC and mental health will result in better outcomes for depressed adolescents who present in PC.

In addition, on behalf of the GLAD-PC team, Stein et al reviewed the literature on psychosocial interventions for anticipatory guidance. No RCTs or evidence-based reviews were found. Citing earlier literature reviews in the area of injury prevention and anticipatory guidance, Stein et al found some limited evidence that anticipatory guidance strategies such as education and counseling in the PC setting can be effective.

Another area reviewed by Stein et al involved psychosocial interventions for improved adherence. An evidence review by Lemanek et al on asthma adherence suggested that some educational and behavioral strategies are “probably efficacious” in creating change. In addition, a study that used cognitive behavioral strategies suggested that diabetic adherence can also be improved.

In addition, our team searched the Cochrane database for systematic reviews of all types of interventions implemented in the adherence arena. These reviews suggested that only complex, multifaceted approaches that included convenient care, patient education, reminders, reinforcement, counseling, and additional supervision by a member of the care team were effective in improving adherence in different chronic medical conditions including asthma, hypertension, diabetes, and adult depression. These complex models have also moved beyond the somewhat paternalistic traditional model to a model of shared decision-making between the provider and patient. However, improvements in treatment outcomes remain small even with these complex and resource-intensive interventions. In the pediatric literature, research regarding adherence commonly involved interventions that targeted both patients and their families. Several key components have been identified that may improve compliance/adherence, including patient self-management/monitoring, patient/family education/support, and the setting and documentation of achievement of management goals. The identification and periodic review of short- and long-term goals provides an individualized plan that both the provider and the patient and family can follow over time. However, current research evidence suggests that more complex interventions that include shared decision-making between provider and patient are likely to have the greatest impact on both adherence and treatment outcomes. This kind of coordinated care has been designated the PC medical home and is discussed further in this section.
Factors that pertain to the linkages between/among organizations have been studied as well. For example, the concept of “system of care” was first described in 1986 in a monograph by Stroul and Friedman and used to support the Children and Adolescent Service System Program; their monograph outlined the ideal model for an integrated system of care for children and youth with severe emotional disturbances. Over the last 2 decades, numerous examples of comprehensive, coordinated, community-based systems of care have been implemented and evaluated. Most of them showed no improvement in patient outcomes but significant improvement in other areas such as patient/family satisfaction, decreased wait time, and more appropriate care.

In the chronic care medical literature, PC professionals have been encouraged to provide a medical home for their patients with chronic conditions. In effect, professionals are expected to remain accessible to patients and families through periods of quiescence and medical crises, coordinate care with other health care professionals, advocate with third-party payers, and provide continuity of care to prevent long-term consequences of chronic illnesses. A review of the evidence for the medical home was conducted by Cooley, who found that although no studies of the outcomes of the broad application of the medical home exist, some evidence exists for positive outcomes for different aspects of the concept. Again, no RCTs exist that we could find. In 1 study, children without a medical home were twice as likely to delay or forego needed care and to have unmet healthcare service needs. Two other reviews focused on the impact of coordination of care and continuity of care. As a recent vaccination study demonstrated, just having a medical home available to a patient does not necessarily mean that the patient will make full use of the available services. Thus, as outlined in a recent American Academy of Pediatrics policy statement, the medical home must provide care coordination and help patients to make use of available resources.

Although the “system of care” focuses on the overall coordination of care involving many agencies that are involved in the care of youth (child welfare services, schools, etc), a crucial link in that system and for adolescent depression is that between mental health and PC. Currently, no specific literature addresses the issue of referral to specialty care of adolescent patients with depression (such as which subgroups of patients would benefit from referral to mental health professionals). Studies have been conducted with adults with depression that demonstrate that increased collaborative care between mental health and PC professionals is needed to improve the care of patients with depression in PC. The adult literature shows the importance of a close working relationship between mental health specialists and PC clinicians in the PC setting. Different models of collaboration have been shown to be effective in the adult depression literature, including the use of case managers in PC practices and consultation by on-site mental health clinicians. Other models include shared care and telephone consultation on an ad hoc basis. Although these models suggest improved outcomes for both clinicians and patients, there are significant barriers to the successful implementation of these collaborative models, including funding deficiencies and shortage of mental health providers. Financial and other incentives for both PC and mental health clinicians to develop these models and obtain the training needed to function within these models are essential. Increased development of skills needed for collaborative care and training in mental health may also be addressed at earlier stages of training such as during residency for both PC and mental health clinicians.

Safety Planning
Safety planning with depressed suicidal or potentially suicidal adolescent patients usually consists of instructing the family to remove lethal means, instructing the family to monitor for risk factors for suicide, engaging the potentially suicidal adolescent in his or her treatment, providing adolescents with mutually agreeable and available emergency contacts should they find themselves with increasing suicidality, and establishing clear follow-up. Our review of the literature found no trials that have studied the impact of or how to conduct any of these aspects of safety planning with depressed adolescents. No studies have examined the benefits or risks of a safety contract. Some studies have suggested that limiting access to firearms or other lethal means can decrease suicide by those methods, but the evidence is still unclear as to whether, on a broader population level, restricting access to certain lethal methods results in an overall decrease of suicide rates. In addition, a study by Brent et al found that families of depressed adolescents are frequently noncompliant with recommendations to remove firearms from the house. Yet, a small prospective follow-up of patients seen in an emergency department for mental health concerns found that the majority of families removed or secured lethal means (firearms, alcohol, prescription medications, and over-the-counter medications) after injury-prevention education in the emergency department, whereas no families who did not receive injury-prevention education did so. Some limited evidence suggests that quick and consistent follow-up with a team approach will be most helpful in increasing compliance and engagement among suicidal patients.

GUIDELINES
Identification
Recommendation 1: Patients with depression risk factors (such as history of previous episodes, family history, other psychiatric
disorders, substance abuse, trauma, psychosocial adversity, etc) should be identified (grade of evidence: C; strength of recommendation: very strong) and systematically monitored over time for the development of a depressive disorder (grade of evidence: C; strength of recommendation: very strong).

Although most PC clinicians believe it is their responsibility to identify depression in their adolescent patients, evidence suggests that only a fraction of these youth are identified when they present in PC settings, and only 50% of depressed adolescents are diagnosed before reaching adulthood. As part of overall health care, PC clinicians should routinely monitor the psychosocial functioning of all youth, because problems in psychosocial functioning may be an early indication of a variety of problems, including depression. For those at known increased risk for depression, PC clinicians should use systematic, effective identification strategies. Risk factors that clinicians may use to identify those who are at high risk for depression include a personal history and/or family history of (1) depression, (2) bipolar disorder, (3) suicide-related behaviors, (4) substance abuse, and (5) other psychiatric illness, or (6) significant psychosocial stressors such as family crises, physical and sexual abuse and neglect, and other trauma history. Research evidence shows that patients who present with such risk factors are likely to experience future depressive episodes. Patients who have been treated for depression or suicidality in the past should continue to be monitored. PC clinicians should systematically evaluate adolescents at high risk for depression during health care visits (ie, well-child visits, urgent care visits). This systematic assessment should take place at least once a year, but frequent somatizers may need to be assessed more often.

Identification methods of youth at high risk may involve tools such as standardized written instruments, either generalized (eg, Guidelines for Adolescent Preventive Services and Strength and Difficulties Questionnaire) or specific emotional symptom checklists (eg, Beck Depression Inventory, Kutcher Adolescent Depression Scale). Although mnemonic-based interviews (eg, HEADSS: home, education/employment, activities, drugs, sexuality, suicide/depression) may also be used routinely during visits to guide the direct interview, systematic and scheduled use of psychometrically reliable and practical methods such as brief symptom checklists or validated depression scales are a preferred adjunct.

Assessment/Diagnosis

Recommendation 1: PC clinicians should evaluate for depression in adolescents at high risk as well as those who present with emotional problems as the chief complaint (grade of evidence: B; strength of recommendation: very strong). Clinicians should assess for depressive symptoms on the basis of diagnostic criteria established in the DSM-IV or International Classification of Diseases, 10th Revision (grade of evidence: B; strength of recommendation: very strong) and should use standardized depression tools to aid in the assessment (grade of evidence: A; strength of recommendation: very strong).

PC clinicians should probe for the presence of any of several depressive disorders, including MDD, dysthymia, and depression not otherwise specified by using systematic, rigorous assessment methods. Standardized instruments should be used to help with diagnosis but should not replace direct interview by the clinician. Because adolescents with depression may not be able to clearly identify depressed mood as their presenting complaint, providers need to be aware of common presenting symptoms that may signal MDD. These symptoms may include insomnia, weight loss, decline in academic functioning, family conflict, and other symptoms of depressive disorders. The Diagnostic and Statistical Manual for Primary Care can help PC clinicians distinguish between transient depressive responses and depressive disorders.

Recommendation 2: Assessment for depression should include direct interviews with the patients and families/caregivers (grade of evidence: B; strength of recommendation: very strong) and should include the assessment of functional impairment in different domains (grade of evidence: B; strength of recommendation: very strong) and other existing psychiatric conditions (grade of evidence: B; strength of recommendation: very strong).

Evidence of the core symptoms of depression and functional impairment should be obtained from the youth and from families/caregivers separately. The involvement of the family is critical in all phases of management and should be included in the assessment for depressive disorders. Family relationships also may affect the presentation of depression in adolescents. Cultural background of the patients and their families also must be considered during the assessment, because it can affect the presentation of core symptoms. Collateral information from other sources (such as teachers) may also be obtained to aid in the assessment. Given the high rates of comorbidities, clinicians should assess for the existence of comorbid conditions that may affect the diagnosis and treatment of the depressive disorder. These comorbidities may include 1 or more of the following conditions: substance abuse, anxiety disorder, attention-deficit/hyperactivity disorder, bipolar disorder, physical abuse, sexual abuse, and trauma. Instruments that assess for a range of common comorbid mental health conditions should be considered also. Clinicians should also assess for impairment in key areas of functioning including school, home, and peer settings. Subjective distress should be assessed also. Regardless of the diagnostic impression or any additional treatment plans, a safety assessment must be completed by the clinician (see recommendation 4 in “Initial Management of Depression”).
Initial Management of Depression

Recommendation 1: Clinicians should educate and counsel families and patients about depression and options for the management of the disorder (grade of evidence: C; strength of recommendation: very strong). Clinicians should also discuss limits of confidentiality with the adolescent and family (grade of evidence: D; strength of recommendation: very strong).

Management should be based on a plan developed with the understanding that depression is often a recurring condition. As seen in studies of depression interventions, families and patients need to be educated about the causes and symptoms of depression, impairments associated with it, and the expected outcomes of treatment. Information should be provided at a developmentally appropriate level, in a way that the patient and family can understand the nature of the condition and the management plan. Communication that is developmentally appropriate should facilitate the ability of parents and patients to work with the clinician to develop an effective and achievable treatment plan. To establish a strong therapeutic alliance, the clinician should also take into account cultural factors that may affect the diagnosis and management of this disorder. Clinicians should also be aware of the negative reactions of family members to a possible diagnosis of depression in the teen (ie, sadness, anger, denial). Sample materials are available in the GLAD-PC toolkit and include resources for patients and parents. Because the symptoms of depression can also affect many areas of an adolescent’s life, other ongoing partnerships may need to be established with personnel in schools and other settings (extracurricular activities). Confidentiality must also be discussed with the adolescent and his or her family. Adolescents and their families should be aware of the limits of confidentiality, including the need to involve parents or legal authorities when the risk of harm to the adolescent or others may be imminent. Clinicians should be aware of state laws regarding confidentiality (eg, see www.advocatesforyouth.org/publications/iag/conflhth.htm for additional information).

Recommendation 2: Clinicians should develop a treatment plan with patients and families (grade of evidence: C; strength of recommendation: very strong) and set specific treatment goals in key areas of functioning, including home, peer, and school settings (grade of evidence: D; strength of recommendation: very strong).

From studies of chronic disorders in youth, it is suggested that better adherence to treatment is associated with the identification and tracking of specific treatment goals and outcomes. Written action plans in asthma management have produced evidence for improved outcomes. If a patient presents with moderate-to-severe depression or has persistent depressive symptoms, treatment goals and outcomes should be identified and agreed upon, based on close collaboration with the patient and family at the time of treatment initiation. Treatment goals may include the establishment of a regular exercise routine, adequate nutrition, and regular meetings to resolve issues at home. In the adult depression literature, monitoring seems most effective when implemented through designated case managers who monitor patients’ clinical status and treatment-plan adherence. The benefits of such programs may be enhanced through the use of electronic medical charts and the development of patient registries.

Recommendation 3: The PC clinician should establish relevant links/collaboration with mental health resources in the community (grade of evidence: B; strength of recommendation: very strong), which may include patients and families who have dealt with adolescent depression and are willing to serve as resources to other affected adolescents and their family members (grade of evidence: D; strength of recommendation: very strong).

A major gap in the management of chronic disorders in young people is the lack of linkages between relevant services that make up the system of care for an individual youth. Furthermore, family-based interventions have been shown to help youth with mental illness. Therefore, establishing relevant links/collaboration with mental health resources in the local community, including peer support groups, advocacy groups, and traditional community- or hospital-based mental health services whenever these services are available, is essential to ensure timely and effective access to needed services. Such linkages may include prearranged agreement regarding referral, exchange of clinical information, points of contact, etc. Where appropriate (eg, rural areas), clinicians should also establish links with para-professionals who may provide the bulk of counseling and supportive services in underserved areas.

Recommendation 4: All management must include the establishment of a safety plan, which includes restricting lethal means, engaging a concerned third party, and developing an emergency communication mechanism should the patient deteriorate, become actively suicidal or dangerous to others, or experience an acute crisis associated with psychosocial stressors, especially during the period of initial treatment when safety concerns are highest (grade of evidence: C; strength of recommendation: very strong).

Suicidality, including ideation, behaviors, or attempts, is common among adolescents with depression. In studies of completed suicide, more than 50% of the victims had a diagnosis of depression. Therefore, clinicians who manage this disorder must develop an emergency communication mechanism for handling increased suicidality or acute crises. After assessing a suicidal patient for suicidality, the clinician must obtain information from a third party, assess that adequate adult supervision and support are available, have an adult agree to help remove lethal medications and firearms from the premises, warn the patient of the disint-
hibiting effects of drugs and alcohol, put contingency planning in place, and establish follow-up within a reasonable period of time.\textsuperscript{72,107} This plan should be developed with adolescents (and with their families/caregivers if possible) and should include a list of persons/services for the adolescent to contact in case of acute crisis or increased suicidality. The establishment of this plan is especially important during the period of diagnosis and initial treatment when safety concerns are highest. Clinicians may also work with schools to develop an emergency plan for all students who may experience an acute suicidal crisis. This global approach may prevent, in some instances, having to label a specific child suicidal when providers are merely trying to ensure that safety measures are in place in case the child decompensates. Components of a safety plan may also include a list of persons who are aware of the adolescents’ issues and will be able to assist if contacted during an acute crisis.

**DISCUSSION**

Although not definitive and subject to modification on the basis of ongoing accumulation of additional evidence, this part of the guidelines is intended to address the lack of recommendations regarding the screening, diagnosis, and initial management of depression in adolescents aged 10 to 21 years in PC settings in the United States and Canada. As such, these guidelines are intended to assist clinicians in family medicine, pediatrics, nursing, and internal medicine who may be the first (and sometimes only) clinicians to identify, manage, and possibly treat adolescent depression. These guidelines may also be helpful to allied health professionals who care for adolescents.

Although not all the steps involved in identifying, diagnosing, and initially managing the care for adolescent depression in PC have been (or even can be) subject to rigorous RCTs, there is sound reason to believe that existing tools and management protocols for adolescent depression can be applied in the PC setting. Although more research is needed, our review suggests that these components of the identification and initial management of adolescent depression in PC can be done. The recommendations were developed on the basis of areas that had at least “strong agreement” among experts. However, there were other controversial areas that were not addressed in these recommendations, such as universal screening. New emerging evidence may affect the inclusion of such areas in future iterations of these guidelines and the accompanying toolkit.

**Should These Guidelines Be Universally Deployed?**

One might question whether PC clinicians should identify and diagnose the problem of adolescent depression if the lack of psychiatric services prevents them from referring these youth.\textsuperscript{108} This caution notwithstanding, the increasingly prevailing recommendation is that as a minimum, PC clinicians should be provided the necessary guidance to support their initial management of adolescent depression.\textsuperscript{109,110} Nonetheless, because practitioners and their clinical practice settings vary widely in their degree of “readiness” in identifying and managing adolescent depression, it is likely that a good deal of time and flexibility will be required before these guidelines are adopted systematically or as a universal requirement. It is conceivable that integrated health care systems with electronic medical charts, tracking systems, and access to specialty mental health back-up and consultation will be most “ready” and able to fully implement the guidelines. The second part of the guidelines, the companion article,\textsuperscript{12} addresses the treatment of this disorder. Practices that do identify adolescent depression and have nowhere to refer the patients may benefit from the guidance offered in the next set of recommendations.

**Preparatory Steps**

Because the management of adolescent depression may constitute a new or major challenge for some PC practices, a number of important considerations should be kept in mind when preparing to implement the guidelines, given the findings from studies in the adult literature, input from our focus groups with clinicians, families and patients, and the experience of members of the GLAD-PC Steering Committee. Specifically, PC clinicians who manage adolescent depression should pursue (1) additional education regarding issues such as advances in screening, diagnosis, treatment, and follow-up, liability, consent, confidentiality, and billing, (2) practice and systems changes such as office staff training and “buy-in,” electronic medical charts, and automated tracking systems, whenever available, and (3) establishing linkages with mental health services.

Linkages with community mental health resources are necessary to both meet the learning needs of the PC clinician and facilitate consultation/referral of difficult cases. Practice and systems changes are useful in increasing clinicians’ capacity to ensure monitoring and follow-up of patients with depression. For example, staff training may help prioritize calls from adolescent patients who may not state the nature of their call. Specific tools and/or templates have been developed that offer examples of how to efficiently identify, monitor, track, and refer teens with depression. These materials are available in the GLAD-PC toolkit (available at www.glad-pc.org). The toolkit addresses how each of the recommendations might be accomplished without each practice necessarily having to “reinvent the wheel.”

**CONCLUSIONS**

Review of the evidence suggests that PC clinicians who have appropriate training and are attempting to deliver comprehensive health care should be able to identify and initiate management of adolescent depression. This
will likely require real changes in existing systems of care. As health care models such as the medical home indicate, comprehensive health care must include assessment and coordination of care for both physical and behavioral health. This first part of the guidelines for adolescent depression in PC may enable providers to pull together the current best evidence and attempt to initiate the best available high-quality care, even in instances in which they are not in a position to treat such youth. Mounting evidence suggests that pediatric providers can and should identify and coordinate depression care for their adolescent population.

**APPENDIX: PART I TOOLKIT ITEMS**

- Screening/assessment instruments (ie. Columbia Depression Scale)
- Information sheet on the developmental considerations in the diagnosis of depression
- Assessment algorithm/flow sheet (Fig 1)
- Fact sheet/family educational materials
- Educational materials on suicide prevention/safety planning

**ACKNOWLEDGMENTS**

We thank the Center for Substance Abuse Treatment (Substance Abuse and Mental Health Services Administration), the Josiah Macy, Jr Foundation, the New York State Office of Mental Health, the Lowenstein Foundation, the Center for the Advancement of Children’s Mental Health (Columbia University), Sunnybrook Health Sciences Centre (University of Toronto), the American Academy of Pediatrics (District II, New York chapters 1, 2 and 3), the New York Council on Child and Adolescent Psychiatry, the Children’s Health Forum (New York Academy of Medicine), the Kellogg Foundation, and the Civic Research Institute, Inc for financial support of the GLAD-PC project.

The GLAD-PC Steering Group consists of members of the GLAD-PC project team, steering committee, and official organizational liaisons. The GLAD-PC project team members are Peter S. Jensen, MD (project director, REACH Institute), Amy Cheung, MD (project coordinator, University of Toronto/Columbia University), Rachel A. Zuckerbrot, MD, (project coordinator, Columbia University), Kareem Ghalib, MD (Columbia University), and Anthony Levitt, MD (project consultant, University of Toronto). The steering committee members are (listed alphabetically) Boris Birmaher, MD (Western Psychiatric Institute & Clinic, University of Pittsburgh), John Campo, MD (Ohio State University and Nationwide Children’s Hospital), Greg Clarke, PhD (Center for Health Research, Kaiser Permanente), Dave Davis, MD (University of Toronto), Angela Diaz, MD (Mount Sinai School of Medicine), Allen Dietrich, MD (Dartmouth Hitchcock Medical Center), Graham Emslie, MD (University of Texas Southwestern Medical School), Bernard Ewigman, MD (Department of Family Medicine, University of Chicago), Eric Fombonne, MD (McGill University), Sherry Glied, PhD (Columbia University), Kimberly Eaton Hoagwood, PhD (Office of Mental Health, New York State/Columbia University), Charles Homer, MD (National Initiative for Children’s Healthcare Quality), Danielle Laraque, MD (AAP New York Chapter 3, District II/Mount Sinai School of Medicine), Miriam Kaufman, MD (Hospital for Sick Children, University of Toronto), Kelly J. Kelleher, MD (Ohio State University), Stanley Kutzer, MD (Dalhousie Medical School), Michael Malus, MD (Department of Family Medicine, McGill University), James Perrin, MD (Massachusetts Medical School/Harvard Medical School), Harold Pincus, MD (Columbia University/New York State Psychiatric Institute), Brenda Reiss-Brennan, APRN (Intermountain Health), Diane Sacks, MD (Canadian Paediatric Society), Ruth E. K. Stein, MD (Forum for Child Health, New York Academy of Medicine, Albert Einstein College of Medicine), and Bruce Waslick, MD, Baystate Health Systems, MA). The organizational liaisons are Angela Diaz, MD (American Academy of Pediatrics), Kelly Kelleher, MD (American Academy of Pediatrics), James Perrin, MD (American Academy of Pediatrics), Diane Sacks, MD (American Academy of Pediatrics/Canadian Paediatric Society), Bruce Waslick, MD (American Medical Association), David Fassler, MD (American Academy of Child and Adolescent Psychiatry), Eric Fombonne, MD (Canadian Academy of Child Psychiatry and Canadian Psychiatric Association), James McIntyre, MD (American Psychiatric Association), Judy Garber, PhD (American Psychological Association), Vicky Wolfe, PhD (Canadian Psychological Association), Michael Malus, MD (College of Family Medicine of Canada), Johanne Renaud, MD (Canadian Association for Adolescent Health), Debbie Ebner, PhD (Society for Adolescent Medicine), Stanford Friedman, MD (Society for Developmental and Behavioral Pediatrics), Terry Stancin, PhD (Society for Developmental and Behavioral Pediatrics), Kathryn Salisbury, PhD (Mental Health Association of New York City), Michael Faenza, MSSW (National Mental Health Association), Susan Bergeson (Depression and Bipolar Support Alliance), Darcy Gruttadaro (National Alliance on Mental Illness), Sandra Spencer (Federation of Families for Children’s Mental Health), and Elizabeth Hawkins-Walsh, DNSc, CPNP (National Association for Pediatric Nurse Practitioners).

In addition to members of the GLAD-PC Steering Group, we also acknowledge the GLAD-PC conference attendees: Perry Adler, PhD; Joan Asarnow, PhD; Lorretta Young Au, MD, FAAP; Abraham Bartell, MD; Tamar Bauer; Rachel Bergeson, MD; Blanche Benenson, MD; Linda Theil Cahill, MD; Wayne Cannon, MD; Marie Barone Casalino, MD; Sonia Chehil, MD; Joseph Cra-
Youth presents to clinic for urgent care or health maintenance visit

Early Identification
Systematically identify high-risk youth

(1) Stop assessment
(2) Repeat surveillance as needed

If low risk
If high risk or presenting with emotional issues as chief complaint

Assessment
(1) Assess with systematic depression assessment tool
(2) Interview patient and parent to assess for depression and other psychiatric disorders with DSM-IV or ICD-10 criteria
(3) Assess for safety/suicide risk

Evaluation Negative for Depression, but positive for other MH conditions
(1) Refer to other treatment guidelines; (2) Evaluate for depression at future visits; (3) Book for follow-up visit.

Evaluation Positive for Depression: MILD, MODERATE, SEVERE, or Depression with COMORBIDITIES
(1) Evaluate safety and establish safety plan.
(2) Evaluate severity of depression symptoms.\(^a\)
(3) Patient/Family Education.\(^b\)
(4) Develop treatment plan based on severity-review diagnosis and treatment options with patient/family.

\(^a\)See Guidelines Part I for definition of mild, moderate, and severe depression. Please consult toolkit for methods available to aid clinicians to distinguish between mild, moderate, and severe depression.

\(^b\)Psychoeducation, supportive counseling, facilitate parental and patient self-management, refer for peer support and regular monitoring of depressive symptoms and suicidality.

Refer to Crisis or Emergency Services (may include subsequent referral to inpatient treatment)

Youth or family presents with emotional issues as chief complaint

Preparation for Managing Depression in Primary Care
Preparation through increased training, establishing mental health linkages, and increasing the capacity of practices to monitor and follow-up with patients with depression.

FIGURE 1
Clinical assessment flowchart.
REFERENCES


Available at: www.pediatrics.org/cgi/content/full/120/5/e1313


32. Smith MS, Mitchell J, McCauley EA, Calderon R. Screening


40. Beck AT, Steer RA. *Reynolds Adolescent Depression Scale.* Lutz, FL: Psychological Assessment Resources; 1987

41. Reynolds WM. *Reynolds Adolescent Depression Scale.* Lutz, FL: Psychological Assessment Resources; 1987


56. Stroul BA, Friedman R, eds. *A System of Care for Children and Youth With Severe Emotional Disturbances.* Washington, DC: Georgetown University Child Development Center, CASSP, Technical Assistance Center; 1986


74. Shah S, Hoffman RE, Wake L, Marine WM. Adolescent sui-
103. Stoul BA, Friedman RM. A System of Care for Children and Youth With Severe Emotional Disturbances. Washington, DC: Georgetown University Child Development Center, CASSP Technical Assistance Center
Guidelines for Adolescent Depression in Primary Care (GLAD-PC): I. Identification, Assessment, and Initial Management
Rachel A. Zuckerbrot, Amy H. Cheung, Peter S. Jensen, Ruth E.K. Stein and Danielle Laraque
Pediatrics 2007;120;e1299
DOI: 10.1542/peds.2007-1144

Updated Information & Services
including high resolution figures, can be found at:
http://pediatrics.aappublications.org/content/120/5/e1299.full.html

Supplementary Material
Supplementary material can be found at:
http://pediatrics.aappublications.org/content/suppl/2010/09/03/120.5.e1299.DC1.html

References
This article cites 93 articles, 21 of which can be accessed free at:
http://pediatrics.aappublications.org/content/120/5/e1299.full.html#ref-list-1

Citations
This article has been cited by 15 HighWire-hosted articles:
http://pediatrics.aappublications.org/content/120/5/e1299.full.html#related-urls

Subspecialty Collections
This article, along with others on similar topics, appears in the following collection(s):
Adolescent Health/Medicine
http://pediatrics.aappublications.org/cgi/collection/adolescent_health:medicine_sub

Permissions & Licensing
Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at:
http://pediatrics.aappublications.org/site/misc/Permissions.xhtml

Reprints
Information about ordering reprints can be found online:
http://pediatrics.aappublications.org/site/misc/reprints.xhtml

PEDIATRICS is the official journal of the American Academy of Pediatrics. A monthly publication, it has been published continuously since 1948. PEDIATRICS is owned, published, and trademarked by the American Academy of Pediatrics, 141 Northwest Point Boulevard, Elk Grove Village, Illinois, 60007. Copyright © 2007 by the American Academy of Pediatrics. All rights reserved. Print ISSN: 0031-4005. Online ISSN: 1098-4275.

American Academy of Pediatrics
DEDICATED TO THE HEALTH OF ALL CHILDREN™