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AHA SCIENTIFIC STATEMENT

Psychological Outcomes and Interventions for Individuals With Congenital Heart Disease: A Scientific Statement From the American Heart Association

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ABSTRACT: Although resilience and high quality of life are demonstrated by many individuals with congenital heart disease, a range of significant psychological challenges exists across the life span for this growing patient population. Psychiatric disorders represent the most common comorbidity among people with congenital heart disease. Clinicians are becoming increasingly aware of the magnitude of this problem and its interplay with patients' physical health, and many seek guidance and resources to improve emotional, behavioral, and social outcomes. This American Heart Association scientific statement summarizes the psychological outcomes of patients with congenital heart disease across the life span and reviews age-appropriate mental health interventions, including psychotherapy and pharmacotherapy. Data from studies on psychotherapeutic, educational, and pharmacological interventions for this population are scarce but promising. Models for the integration of mental health professionals within both pediatric and adult congenital heart disease care teams exist and have shown benefit. Despite strong advocacy by patients, families, and health care professionals, however, initiatives have been slow to move forward in the clinical setting. It is the goal of this scientific statement to serve as a catalyst to spur efforts for large-scale research studies examining psychological experiences, outcomes, and interventions tailored to this population and for integrating mental health professionals within congenital heart disease interdisciplinary teams to implement a care model that offers patients the best possible quality of life.

Key Words: AHA Scientific Statements ■ anxiety ■ depression ■ heart defects, congenital ■ psychotherapy ■ psychotropic drugs ■ quality of life

s a result of significant medical advances in recent decades, >90% of newborns with congenital heart disease can now survive to adulthood. There are currently >2.4 million Americans living with congenital heart disease, and adults now outnumber children.¹ Although reductions in mortality and morbidity are appropriately extolled, living with the challenges of congenital heart disease is associated with a higher prevalence of emotional, behavioral, and social difficulties compared with healthy peers. The lifetime prevalence of depression and anxiety in this patient population is as high as 50%²³³ and has been associated with higher medical costs in addition to greater morbidity and mortality.⁴⁵ The importance of this messaging is not limited to physicians and health

care professionals who specialize in congenital heart disease. Awareness of psychological sequelae is necessary for every clinician who takes care of individuals with congenital heart disease, including those who specialize in heart failure, electrophysiology, coronary artery disease, and surgery. Pediatric cardiology and adult congenital heart disease programs have established clear referral and treatment pathways for patients with other clinically significant symptoms or organ dysfunction (eg, referral to pulmonologists, neurologists, or hepatologists), yet the psychological needs of individuals with congenital heart disease have been largely unmet in comparison. The onus on accessing psychological care should not be placed solely on the shoulders of patients and families.

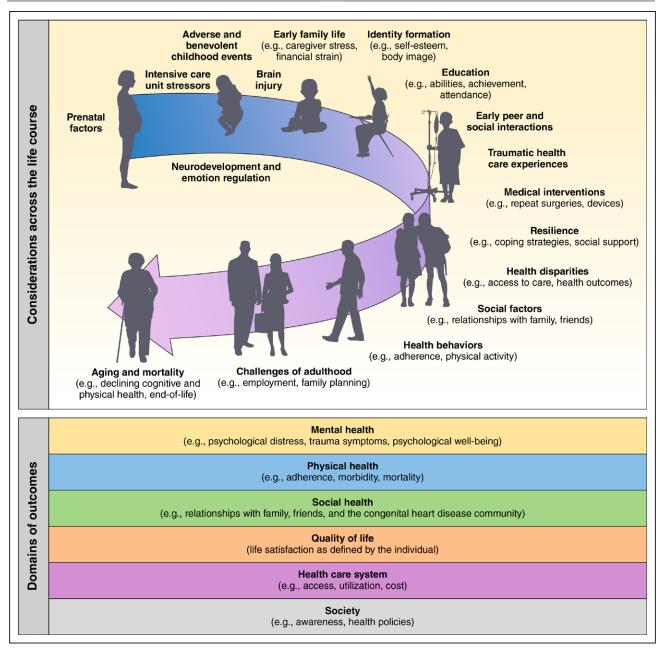


Figure. Diagrammatic representation of contributing factors and domains of outcomes for the psychological well-being of individuals with congenital heart disease.

Such factors and outcomes are multiple, interrelated, and cumulative across the life span with significant individual, family, and societal effects.

Furthermore, it is no longer sufficient to focus primarily on awareness among congenital heart disease clinicians of the challenges that patients and families encounter in maintaining psychological well-being. Collectively, clinicians, their institutions, and health systems are now called on to engage in implementation of strategies to prevent and alleviate significant distress and to support resilience and mental health as part of routine care.

This scientific statement summarizes the psychological outcomes of infants, children, adolescents, and adults with congenital heart disease and reviews mental health interventions, including psychotherapy and

pharmacotherapy. We occasionally use the term psychosocial to highlight ways in which congenital heart disease might affect both psychological and social outcomes and the bidirectional relationship between psychological and social factors. This scientific statement incorporates but does not significantly elaborate on neurodevelopmental outcomes because they have been effectively summarized in a previous scientific statement from the American Heart Association.⁶ Rather, ways in which psychological and neurodevelopmental or neurocognitive outcomes are intertwined across the life span are incorporated as appropriate.⁷

The statement is intended to serve as an urgent call to action for pediatric and adult programs to implement a proactive approach to integrating psychological care and improving emotional, behavioral, and social outcomes for individuals of all ages living with congenital heart disease.

PSYCHOLOGICAL OUTCOMES ACROSS THE LIFE SPAN

Shared Origins of Psychological and Neurodevelopmental Outcomes

Risk factors for adverse psychological and neurodevelopmental outcomes in individuals with congenital heart disease are multiple, interrelated, and cumulative (Figure) and may exert influence in utero and across the life course. Such factors include altered fetal circulation and cerebral hemodynamics, altered placental environment, hypoxia, brain dysmaturation, cardiac arrest, perioperative neurological injury (eg, seizures, stroke), use of mechanical cardiopulmonary support, medications (including anesthetics), pain, prolonged hospitalization, exposure to stress or trauma, genetic and epigenetic factors, and parental mental health.^{6,8} Data suggest that preoperative risk factors (eg, genetics, family socioeconomic hardship, low birth weight, gestational age, brain dysmaturation, neonatal transition), as well as cumulative morbidity and parental mental health, may contribute more strongly to adverse psychological and neurodevelopmental outcomes than intraoperative support techniques such as cardiopulmonary bypass.8 There is also growing recognition of the role of ecological factors acting on genetic predispositions, particularly the potential lifelong consequences of cumulative exposure to adversity and toxic stress in early childhood.9 Furthermore, there is evidence that sociocontextual, economic, and race and ethnicity disparities influence congenital heart disease surgical outcomes, lapses in care, and mortality across the life span¹⁰⁻¹²; it is likely that such disparities may also affect psychological outcomes in congenital heart disease because socioeconomic status and mental health are linked in the general population. 13,14

Overall, the prevalence of neurodevelopmental delay or impairment in congenital heart disease is high, ^{6,8} and the link between psychological and neurodevelopmental outcomes is well established. ^{15–22} The Boston Circulatory Arrest Study, for example, demonstrated that neurodevelopmental difficulties exhibited by school-aged children (eg, attention)²³ and adolescents (eg, executive function, social cognition)²⁴ with dextrotransposition of the great arteries were associated with poorer psychosocial health status, as reported by their parents. Moreover, impaired cognitive functioning in children with dextro-transposition of the great

arteries at 8 years of age was found to predict subsequent psychosocial health at 16 years of age, with attentional difficulties one of the strongest predictors of future lower parent-proxy-reported psychosocial health status in this study.²⁴ Greater clinical severity of attention-deficit/hyperactivity disorder (ADHD) is associated with poorer psychosocial health^{25,26} and quality of life²⁷ in community samples, as well as a greater risk of graduation failure, juvenile justice system involvement, and mortality from suicide. 25,28 These findings highlight the role of attentional difficulties, and neurodevelopmental impairment more broadly, as critical targets for interventions to reduce longer-term psychological suffering and adversity, including physical, mental, and financial health into adulthood. 26,28 Among adolescents and adults, poorer executive functioning, less effective coping skills, and psychological distress are often intertwined.^{29,30}

Challenges and Outcomes During Infancy, Childhood, and Adolescence

Any serious medical illness may profoundly influence a child's developmental trajectory. A child's life can be affected by multiple stressors that may begin before or shortly after birth and continue into childhood, adolescence, and beyond. Infants who undergo cardiac surgery show high levels of stress and distress. Babies with congenital heart disease are often hypersensitive to stimuli and may experience difficulties with early neurobehavioral and regulatory capacities such as feeding, sleeping, settling, and soothing,31 as well as autonomic and motor organization.32 For critically ill infants within the intensive care unit, stressors can originate from the physical environment (eg, bright lights, loud sounds, noxious smells), clinical exposures (eg, painful and invasive procedures, neurotoxic medications),33 and psychosocial environment (eg, high parental emotional distress, limited opportunities for bonding and attachment).34 Early nutritional deficiencies from unrecognized or uncompensated heart failure can lead to failure to thrive, which in general has been associated with poorer cognitive development and educational deficits among children.35 In the context of congenital heart disease, it can be difficult to separate the additional burden of failure to thrive from the confounding effects of the underlying morbidity that can lead to the nutritional deficits in the first place, but it likely poses an additional risk factor among these children. Additional challenges across early childhood can include periods of separation from caregivers and social supports, acute and persistent medical problems, exacerbations of disease, and the burden of care that may disrupt life consistency (eg, school, physical activity, early relational bonds) and place a financial burden on individuals and families.36

As a group, children with more complex congenital heart disease and those with comorbid medical conditions are at greater risk of neurodevelopmental impairment⁶ and emotional and behavioral dysregulation.³⁷ Children with complex congenital heart disease may show early signs of social withdrawal,38 and children and youth have elevated rates of internalizing (eg, anxiety, depression; 25%) and externalizing (eq. aggression, hyperactivity; 15%) difficulties.³⁹ A recent meta-analysis that included parent-proxy reports of psychological distress among children and adolescents with congenital heart disease (all of whom had surgical intervention at <2 years of age) reported an increased prevalence of behavioral difficulties, including aggression, opposition, and defiance.³⁹ For these reasons, it can often be valuable for pediatric congenital heart disease clinicians to collaborate and coordinate efforts with teachers and school counselors to optimize educational, psychological, and social outcomes. The goal should be to support individual patient needs without placing unnecessary limitations on children's activities. 40 Children and adolescents with single-ventricle congenital heart disease have been found to have a 5-fold higher rate of lifetime anxiety disorder diagnosis and an almost 6-fold higher rate of lifetime ADHD diagnosis, which can affect school and social functioning.41 This study reported a lifetime prevalence of any psychiatric disorder of 65% in this patient group.41 A higher prevalence of psychiatric diagnoses, including autism spectrum disorder, compared with the general population has also been reported for other congenital heart disease subgroups.42 A recent meta-analysis of health-related quality of life among individuals with a Fontan circulation synthesized findings from 50 articles reporting 2793 patients and 1437 parents-proxies.⁴³ The authors concluded that poorer emotional functioning was associated with older age at Fontan operation, whereas poorer social functioning was associated with the presence of hypoplastic left-sided heart syndrome. Despite this cumulative evidence, only a small fraction of children with congenital heart disease are offered or participate in formal psychological assessment or treatment.44

A systematic review of 44 qualitative studies that analyzed patterns in the narratives of 995 children and young people with congenital heart disease from 12 countries identified 6 key themes that have implications for psychological outcomes⁴⁵: (1) disruptions to normal life (eg, fluctuations between sickness and health, destabilizing the family dynamic); (2) powerlessness during periods of deteriorating health (eg, vulnerability to complications, thoughts of mortality, exhaustion from medical testing); (3) enduring uncertainty and medical adversity (eg, trauma associated with invasive procedures, uncertainty and disappointment associated with treatment failure, a sense of displacement during transition to adult cardiac services, valuing empathy and continuity in clinical care); (4) warring with the body

(eg, fatigue, physical limitations, distorted body image); (5) hampered goals (eg, feeling disabled, limited goal attainment and maintenance of milestones); and (6) establishing one's own pace (eg, fostering hope and a determination to thrive, embracing the positives, finding personal enrichment and social or spiritual support).

Adolescents with congenital heart disease face the typical challenges of this developmental stage (eg, cultivation of self-identity, independence, autonomy, and social networks) and additional disease-related stressors. Typical adolescent development involves decreasing reliance on parental support and greater investment in peer relationships.²⁹ In the context of congenital heart disease, this process may be altered by multiple factors, including academic challenges, healthrelated school absenteeism, physical limitations that restrict activities, social anxiety, body image concerns (eg, related to surgical scarring), parental overprotection, and ambivalence about disclosing their congenital heart disease diagnosis to peers. 40,46 Among adolescents with congenital heart disease, depressive symptoms and loneliness are closely linked and have been associated with poorer quality of life.47

During adolescence, most individuals with congenital heart disease can ideally begin the transition process and develop knowledge and self-management skills to gradually assume increased responsibility for their health care and to prepare for transfer from pediatric to adult health services. There are unique considerations to ensure a respectfully tailored experience for individuals with neurodevelopmental disabilities. Inherent to this process are incremental shifts in patient-parent and patient-clinician relationships. Best practices for transition emphasize the importance of holistic, psychosocial assessment and early intervention, as well as diseasespecific education and social support, to ensure success in disease self-management.⁴⁸ This transition may result in increased disease-related stress for adolescents with congenital heart disease who vary greatly in their preparedness for tasks such as speaking independently with health care professionals, overseeing medication regimens, and making decisions about treatment. Psychological distress has been linked to poorer treatment adherence among adolescents with other chronic illnesses, 49 and this association presumably also exists in congenital heart disease. Thus, at a time when medical teams typically transition responsibility for disease management from parents to adolescents, individuals with congenital heart disease are also especially vulnerable to psychological distress that might hinder adherence and contribute to lapses in medical care.

Challenges and Outcomes During Adulthood

For adults with congenital heart disease, heightened vulnerability to emotional distress and elevated rates of

mental health diagnoses persist. Disease-related stressors include awareness of the potential for future medical interventions and life-threatening complications, concerns about early mortality, potential restrictions on family planning, and financial challenges.⁵⁰ Medical outcomes (eg, major adverse cardiac events) in adolescents and adults with congenital heart disease are poorer among those with lower socioeconomic resources.⁵¹ With regard to financial stressors, in general, income and rates of employment are lower among adults with congenital heart disease.⁴⁰ Adult patients also might face disability-related discrimination and ableism in the workplace. 40,52,53 Other factors that may contribute to psychological distress include traumatic memories of previous health experiences, vulnerability and disempowerment in hospital encounters (eg, revealing hospital gowns), advanced cardiac therapies (eg, cardiac device implantation, ventricular assist device, transplantation), invasive procedures, and medical emergencies. 52,54 Stress may also arise from challenges accessing specialized adult congenital heart disease care attributable to insurance and financial barriers or the distance of a patient's residence from a qualified adult congenital heart disease clinician. 55 Last, advancing age carries a unique set of experiences that may affect psychological well-being, including sexual dysfunction, cognitive decline (including an increased risk of early-onset dementia), and declining health status and approaching mortality.8,56,57 Older adults with congenital heart disease often face acquired heart disease and heart failure, which have been associated with increased prevalence of depression and anxiety in the general population,⁵⁸⁻⁶⁰ including among older patients.⁶¹

Adults with congenital heart disease also face challenges unrelated or indirectly related to health (eg, relationships, work demands), as well as non-congenital heart disease-related health stressors (eg, other medical diagnoses, COVID-19 [coronavirus disease 2019] pandemic). Racism and racial discrimination are also adversely associated with physical and mental health outcomes. 62,63 Health disparities in congenital heart disease exist, as demonstrated by evidence of lower rates of insurance among Hispanic patients. 64 Congenital heart disease-related challenges, therefore, must be considered within a broader framework of multiple, cumulative, and interactive stressors.

It is therefore unsurprising that, as a group, adults with congenital heart disease are at elevated risk of clinically significant psychological distress. A 2018 review of studies using self-report assessments found that rates of elevated depressive symptoms ranged from 6% to 34%, whereas elevated anxiety symptoms were reported by 34% to 42% of adults with congenital heart disease. Studies from North America, Germany, and France that incorporated structured clinical interviews observed current³ or lifetime²,66 prevalence rates of mood or anxiety disorders of ≈50%. Known risk factors for depression and anxiety among adults with congenital heart disease

include poorer neurocognitive functioning (eg, executive function),²⁹ loneliness,² limited social support,⁶⁷ poorer perceived health status,² and sedentary behavior.⁶⁸

Beyond depression and anxiety, additional mental health concerns warrant attention. For example, there is emerging evidence that adults with congenital heart disease have an increased risk of traumatic stress and posttraumatic stress disorder (PTSD), which is understandable given a lifetime of medical surveillance and often multiple invasive procedures. In a study using the self-report PTSD Checklist-Civilian version to assess symptoms in relation to any trauma, 21% of 134 adults with congenital heart disease met criteria for PTSD symptoms, a rate markedly higher than that reported in the general community.69 Furthermore, in the context of specific genetic conditions such as DiGeorge syndrome (22q11.2 deletion) and 3q29 recurrent deletion, there is an incidence of other psychiatric disorders, including psychotic disorders, autism spectrum disorder, and ADHD.^{70,71} For example, individuals with 22q11.2 deletion syndrome, which is associated with conotruncal defects such as tetralogy of Fallot,72 face an elevated prevalence of psychiatric disorders; ≈25% will develop schizophrenia, highlighting the necessity of interdisciplinary care. 73 Among individuals with Down syndrome, of whom approximately one-third to one-half will have cardiac malformations,74 between one-quarter and onethird demonstrate maladaptive behavior, and the reported prevalence of psychiatric disorders (eg, autism, depression, ADHD) varies significantly, from 6% to >50%.75 We are unaware of published studies of psychological outcomes focused specifically on individuals with both congenital heart disease and Down syndrome.

There is emerging evidence that health care use (eg, more emergency department visits and hospitalizations and costlier hospitalizations) and mortality risk are higher for adults with congenital heart disease and comorbid depression or anxiety.^{5,76} A recent study examining outcomes for 1146 outpatient adults with congenital heart disease found those with a history of major depression to be more likely to have severe complex congenital heart disease, cyanosis, and worse functional class.4 Furthermore, diagnosis of depression was associated with biomarkers indicative of systemic inflammation (eg, high-sensitivity C-reactive protein) and heart failure (eg, NT-proBNP [N-terminal pro-B-type natriuretic peptide]), as well as increased risk of mortality, even after adjustment for important confounders.4 A clinical diagnosis of depression was associated with an increased risk of mortality among both men and women, with no notable differences based on sex.4

Summary

Individuals with congenital heart disease face a range of potential adverse life experiences that may affect psychological, social, educational, vocational, and medical outcomes across the life span. Awareness of patients' psychological needs is a necessary yet insufficient approach to comprehensive care. To use a medical analogy, within our field, it is accepted that one of the advantages of uninterrupted retention in specialized congenital heart disease care is the opportunity to detect cardiac symptoms and diagnoses that warrant prompt medical intervention. The assumption is that the avoidance of unnecessary delays in treatment can improve early and longer-term outcomes. A similar approach to timely mental health intervention is warranted, such that there is clear need for more timely recognition and treatment of significant psychological distress.

MENTAL HEALTH INTERVENTIONS

Given the widely documented challenges to mental health experienced by individuals with congenital heart disease across the life span, investment in psychological intervention is a high priority. Unfortunately, most children, young people, and adults with congenital heart disease who report clinically significant psychological distress in research studies have not received timely mental health treatment (psychotherapy or psychotropic medication) or other appropriate intervention.^{2,42} There are likely multiple reasons for this, including stigma and insurance barriers for many individuals. Social determinants of health must also be considered; a recent study demonstrated that non-Hispanic Black, Hispanic, and Asian American children (compared with non-Hispanic White children) and uninsured children with congenital heart disease are less likely to receive a diagnosis or to have treatment for a mental health condition.⁷⁷ If symptoms of emotional distress impede daily functioning or impair quality of life, treatment referral is clearly warranted. Mental health professional availability, however, varies between institutions and geographic regions.

We strongly advocate for the integration of mental health professionals within congenital heart disease teams or the "medical home." This has become standard in oncology settings; however, it is disappointingly uncommon in congenital heart disease. Embedding psychologists and other mental health professionals within congenital heart disease teams can normalize emotional reactions to health challenges, reduce stigma associated with seeking mental health treatment, and improve timely access to appropriate, specialized mental health care from experienced clinicians. Integration of mental health professionals within congenital heart disease teams allows a more comprehensive interdisciplinary approach to patient and family care. Mental health professionals may also develop or support other opportunities to enhance psychological well-being, including peer support groups and camps for individuals with congenital heart disease, which create a sense of belonging and acceptance and are highly valued by participants. 78,79 Furthermore, they may foster psychologically informed care within the congenital heart disease team, which includes promoting a soothing presence through voice and touch, creating a healing environment, supporting the psychological needs of the whole family, training clinicians to provide emotional support, and empowering patients.^{80,81}

Models for the integration of psychologists and psychiatrists within both pediatric cardiology and adult congenital heart disease care teams exist and have shown benefit.41,81-84 Within these clinical models, the frequency and duration of sessions and the application of specific therapeutic techniques are based on an individual's presenting psychological concerns; current coping abilities; and needs, values, and preferences. A retrospective review of 100 adults with congenital heart disease referred to an experienced psychologist integrated within an adult congenital heart disease team found that cognitive-behavioral therapy was associated with reductions in self-reported psychological distress.83 Integration of psychologists and other mental health professionals within congenital heart disease programs (in contrast to referral to community clinicians) may offer multiple advantages, including the following:

- Timely provision of psychological care, which is grounded in an in-depth understanding of congenital heart disease and tailored to the unique needs of individual patients within a setting with which they are already familiar and engaged;
- Increased awareness among all congenital heart disease team members of consideration of psychological and social factors when making treatment recommendations, creating a culture of psychologically informed care;
- Greater opportunities to provide access to mental health care regardless of socioeconomic or insurance status;
- Knowledge about which forms of mental health treatment, including which approach to psychotherapy, are most appropriate for individual patients;
- Superior interdisciplinary health care coordination and communication (eg, shared electronic medical records, colocation of office and clinic space, attendance at medical rounds and case conferences, mental health care rounds);
- Increased comfort with and confidence in initiating discussions of patient and family psychological wellbeing among congenital heart disease physicians and nurses, knowing that they have a colleague to whom they can refer patients as appropriate;
- Education and skills-based training of congenital heart disease team members and trainees (eg, developmentally appropriate care, psychologically informed and trauma-informed care, delivering difficult news); and
- Interdisciplinary quality improvement and research collaboration.

Table 1. Psychotropic Drugs and Specific Considerations for Individuals With Congenital Heart Disease*

Psychotropic class	Pharmacological effects of cardiovascular significance	Specific considerations for individuals with congenital heart disease				
SSRIs used for mood, anxiety, or obsessive-compulsive disorders	CYP450 inhibition by SSRI/SNRI (fluoxetine, fluvoxamine, and paroxetine have a	Overall, SSRIs are considered safe and typically preferred as first-line pharma- cotherapy for depression and anxiety in pediatric and adult populations.				
(eg, citalopram, escitalopram,† fluoxetine,† fluvoxamine,† ser- traline†) SNRIs used for mood, anxiety, or obsessive-compulsive disorders (eg, duloxetine,† desvenlafaxine, venlafaxine) SNRIs used for ADHD (atom- oxetine‡)	stronger effect than citalopram, escitalo- pram, sertraline, and venlafaxine) ¹⁰⁶ Potassium channel blockade: QTc prolon- gation (variable and overall modest) ¹⁰⁷	SNRIs are also considered safe for use in the management of depression and anx ety in adult patients. Duloxetine is the only SNRI approved for the management of anxiety in pediatric patients. SNRIs cause modest increases in heart rate and blood pressure, and similar considerations apply as with psychostimulants below.				
	Noradrenergic side effects (SNRI): increased heart rate and blood pressure (modest)	Caution is indicated when used with medications that interact with the CYP450 pathway, including anticoagulants (eg, warfarin) and β-blockers (eg, metoprolol). Caution is indicated in patients with hepatic dysfunction/heart failure and in the presence of electrolyte abnormalities.				
		The unknown risk of arrhythmia should be weighed against risk of undertreating anxio-depressive disorders. 108,109				
TCAs (eg, amitriptyline, clomipramine, imipramine)	Sodium channel blockade: type IA antiar- rhythmic effect causing electric conduction delay with prolongation of PR and QRS	TCAs are associated with a mild increase in risk of arrhythmias in the first month of treatment. They should be avoided or carefully monitored in patients with proarrhythmic conditions.				
	Potassium channels blockade: QTc pro- longation (worse than SSRIs) ¹⁰⁷	Patients with postsurgical intracardiac scars or conduction abnormalities (Fontan procedure, tetralogy of Fallot repair, atrial switch, I-looped heart) are more prone to reentrant tachyarrhythmias and higher degree of heart block.				
	$\alpha_{\rm I}/\alpha_{\rm 2}$ -Adrenergic receptor blockade and anticholinergic effects: orthostatic hypotension and sinus tachycardia	Because of the unfavorable side effect profile, TCAs are considered second-line therapy for depression or anxiety in adults ^{106,110} and are not recommended for managing depression or anxiety in children and adolescents with congenital heart disease.				
Mood stabilizers (eg, lithium,§ valproic acid, lamotrigine)	Sodium channel blockade: Lithium can cause dose-dependent inhibition leading to decreased intracellular potassium and related electrocardiographic abnormalities (including increased QTc dispersion but not QTc duration at therapeutic dose), bradycardia, conduction abnormalities, and ventricular tachyarrhythmias. 111,112 Lamictal exhibits class IB antiarrhythmic activity and can slow ventricular conduction.	Lithium can be associated with time-dependent decline in renal function. 113 lt rarely induces bradycardia and should be used with caution with other bradycardia-inducing agents (eg, β-blockers). Thiazides can increase lithium levels and lead to toxicity. Hypokalemia should be avoided or corrected in patients taking lithium. Valproic acid can cause thrombocytopenia, hepatotoxicity, and increased toxicity in hypoalbuminemic states. 113				
		Both valproate and lithium can cause major congenital malformations in the fetus. Lamotrigine should be avoided in patients with cardiac conduction disorders (ie, second- or third-degree heart block). Concomitant use of other sodium channel blockers may increase risk of conduction abnormalities. ¹¹⁴				
Benzodiazepine anxiolytics (eg, alprazolam, clonazepam, lorazepam, oxazepam, temazepam)	No direct pharmacological cardiovascular effects or electrocardiographic changes are seen, but indirect decrease in heart rate and blood pressure can occur as a result of their anxiolytic properties.	Indicated primarily as second-line therapy for acute anxiety management and for short-term use. ¹¹⁰ Can lead to reduction in anxiety-associated tachycardia and myocardial work. Continuous use for ≥4 wk is associated with possible tolerance and withdrawal symptoms. ¹¹³ Metabolized by the liver and can have long-acting metabolites; lorazepam, oxaz-				
		epam, and temazepam undergo conjugation and are less dependent on global liver function. Caution is indicated when used with other medications metabolized by the live and in the presence of hepatic dysfunction attributable to heart failure; lorazepam, oxazepam, and temazepam are safer choices in patients with polypharma				
α_2 -Adrenergic receptor agonists (eg, clonidine,‡ guanfacine,‡ dexmedetomidine)	Central α_2 adrenergic receptor stimulation decreases heart rate and blood pressure (modest). ^{115,116}	or hepatic dysfunction. Patients on antihypertensive or diuretic drugs are at increased risk for hypotension, especially during initiation and after each dose increase.				
Psychostimulants used to reduce ADHD symptoms (eg, amphetamine/dextroamphetamine mixed salts,‡ lisdexamfetamine,‡ methamphetamine,‡ methylphenidate‡)	Catecholaminergic effects: elevation of resting heart rate and increased systolic blood pressure (modest) ¹¹⁷	Caution is needed when used in patients with dynamic subaortic stenosis, dilated aorta, and severe left ventricular dysfunction or those at risk of severe arrhythmias. ^{118,119}				
Antipsychotics (eg, aripiprazole,§ clozapine, haloperidol, olanzapine,§ quetiapine,§ risperidone§)	Potassium channel blockade: prolonged QTc prolongation (variable) 120 α_1 -Adrenergic receptor blockade: orthostatic hypotension	The unknown risk of arrhythmia should be weighed against risk of undertreating psychosis or bipolar disorder. Metabolic syndrome (weight gain, dyslipidemia, and diabetes). 121 Caution when used together with antihypertensive or diuretic drugs because of increased risk for hypotension, especially during initiation and after each dose increase.				

ADHD indicates attention-deficit/hyperactivity disorder; SNRI, serotonin-norepinephrine reuptake inhibitor; SSRI, selective serotonin reuptake inhibitor; and TCA, tricyclic antidepressant.

^{*}This table is not meant to be an exhaustive list of side effects for each medication.

tUS Food and Drug Administration approved for management of major depressive disorder, generalized anxiety disorder, or obsessive-compulsive disorder in pediatric patients. 122

 $[\]pm$ US Food and Drug Administration approved for management of ADHD in pediatric patients. 123

[§]US Food and Drug Administration approved for management of bipolar I disorder, 124 schizophrenia, or irritability associated with autism spectrum disorder in pediatric patients. 125

Coordination of mental health care should not be considered a single event but rather an ongoing collaborative process of communication and shared care among congenital heart disease medical, nursing and allied health professionals; mental health professionals; and patients and families. During referral to psychological services, it is important to consider patient and family needs and values, including preferences for individual, family, or group psychotherapy; pharmacotherapy; or peer support.85,86 Selection of specific therapeutic approaches should be culturally sensitive, developmentally appropriate, and commensurate with the individual patient's cognitive abilities, given the known risks of neurodevelopmental and neurocognitive impairment in congenital heart disease.²⁹ We provide an overview of psychotherapy and pharmacotherapy approaches for the congenital heart disease patient population below.

Psychotherapy

There are many different approaches to psychotherapy (often referred to as talk therapy). Therapists working in medical settings may align mainly with one approach, yet they typically incorporate techniques of others as appropriate. The following is a nonexhaustive list of examples of approaches to psychotherapy that might be offered to individuals with congenital heart disease. Cognitivebehavioral therapy, with a focus on challenging and modifying maladaptive cognitions and behaviors, has a strong evidence base to support its use in the treatment of depression and anxiety with children and adults, including in the primary care setting, and among adults with cardiovascular disease.87-89 It was the chosen framework for a coping skills intervention developed for adults with congenital heart disease.90 Acceptance and commitment therapy is an approach to cognitive-behavioral therapy that emphasizes psychological flexibility, and preliminary research indicates acceptability among adults with heart disease.91 Trauma-focused therapy typically includes exposure techniques and cognitive processing and has proved effective for children and adults with PTSD.92,93 Eye-movement desensitization reprocessing therapy is another treatment that targets trauma and PTSD.94 Mindfulness-based interventions, which typically include meditation and relaxation techniques, have also demonstrated positive psychological outcomes for individuals, including those with cardiovascular disease.95 Attachment-based psychotherapies often take a child-led approach and focus on supporting and strengthening the parent-child attachment relationship.96,97 Psychodynamically oriented therapies focus on patterns in an individual's thoughts, feelings, self-concept, relationship and life experiences using a developmental perspective to understand how early experiences may influence present psychological difficulties.98 We draw on strong empirical support for psychotherapy in the general population and among

individuals with heart disease and on our collective clinical experiences to suggest that psychotherapy is likely beneficial within the congenital heart disease setting.

Published trials of psychological interventions for children, adolescents and parents affected by pediatriconset chronic illness (including but not limited to congenital heart disease) have yielded encouraging results. Demonstrated benefits of these interventions include less procedure-related distress, greater disease-related knowledge, more appropriate health care use and improvements in a range of psychological outcomes. 99,100 Short-term behavioral interventions (mean total duration, 7.5 hours) have been trialed most frequently and appear to have meaningful beneficial effect for up to 12 months after intervention. Differences in intervention effects have been identified for age and sex, with women, older children and adolescents on average reporting greater emotional or behavioral benefits.

Unfortunately, published psychotherapy outcomes in congenital heart disease are scarce. Two recent systematic reviews synthesized evidence from controlled trials of psychological interventions in congenital heart disease. One focused specifically on interventions for parents of infants during intensive care unit admission,34 and the other included interventions for patients of all ages with childhood-onset heart disease or their family members.101 For parents of children with congenital heart disease, interventions focused predominantly on education, parenting and coping skills training, parent-infant interaction and bonding, procedural preparation, child school readiness, or early pediatric palliative care. Tested interventions demonstrated significant reductions in maternal anxiety and improvements in maternal coping, family functioning, child school functioning and absenteeism but yielded mixed evidence of efficacy for other parental (eg, depression) and child (eg, neurodevelopment, feeding) outcomes. For adolescents and adults with congenital heart disease, interventions targeted mindfulness, resilience, cardiac rehabilitation, exercise and peer engagement, and none yielded evidence of changes in psychological outcomes. This finding signals the need to explore other therapeutic approaches and to identify potential barriers and facilitators to successful intervention and support for older patients.

More recently, a pilot study assessing the feasibility of group-based cognitive-behavioral therapy (eight 90-minute sessions) for adults with congenital heart disease identified several barriers to participation, including scheduling and transportation. Intervention content included psychoeducation (eg, information about common psychological aspects of living with congenital heart disease), cognitive-behavioral strategies (eg, cognitive restructuring, relaxation training, behavioral activation, activity pacing, strategies to improve sleep), social interaction and communication skills training. Although the sample was small, a moderate effect size favoring the intervention was found for depressive

symptoms. Given the paucity of interventions for adults with congenital heart disease, extrapolation from acquired heart disease research may be informative. A Cochrane systematic review and meta-analysis of 35 randomized controlled trials of psychological interventions (including at least a 6-month follow-up) for adults with coronary artery disease (N=10703) concluded that psychological interventions improved depressive symptoms, anxiety and stress and reduced cardiovascular mortality. Considerable uncertainty remains, however, about the magnitude of these effects, which approaches are most likely to yield greatest benefit, and for whom.

Across the developmental spectrum, the small number of existing trials in congenital heart disease is limited by the use of nonrandomized designs, substantial variability in the discipline, training and experience of individuals providing intervention, low response and high attrition rates, short duration of follow-up assessment, and limited diversity of study samples (eg, in terms of race, ethnicity, socioeconomic resources, congenital heart disease complexity). Large-scale studies with prespecified trial and intervention procedures, validated measurement techniques and greater participant engagement and diversity are needed to rigorously test the effects of protocolized psychological intervention for individuals with congenital heart disease and their families. Such trials should include well-defined outcomes monitored over at least 6 months (preferably 12 months) after the intervention to assess durability of benefits. Because accessibility is crucial to the success of psychotherapies, innovative delivery mechanisms, including the use of telemedicine and digital platforms, should be considered in future investigations, along with economic and process evaluations to provide evidence for sustainable implementation. 103

We advocate for the rigorous, empirical investigation of protocolized psychological interventions for individuals with congenital heart disease. At the same time, we recognize and value the unique nuances of psychotherapy practiced in the clinical setting, in which treatment goals, foci and therapeutic techniques vary between clinicians and patients. For example, the course of psychotherapy would appropriately differ among a parent-infant dyad to support bonding and attachment during an extended hospital stay, a child experiencing intense separation anxiety, an adolescent struggling with decision-making, a young adult facing a complex surgical intervention, and an adult facing the end of their life. An embedded mental health professional, familiar with common challenges faced by individuals with congenital heart disease, would make an informed decision about the therapeutic approach and techniques appropriate for use in each situation.

Pharmacotherapy

Just as congenital heart disease diagnosis and medical management entail great heterogeneity, so do pathways to psychiatric pharmacotherapy. A first challenge for individuals with congenital heart disease is that symptoms related to their underlying cardiac condition may be misattributed to an emotional or behavioral problem. For example, shortness of breath secondary to cardiac dysfunction may be mislabeled as anxiety, and poor appetite and psychomotor retardation related to heart failure may be misdiagnosed as depression. Psychotropic medications should therefore be initiated only once evidence of a comorbid psychiatric disorder is clearly determined.

In childhood and adolescence, some disorders may be managed without medications through psychotherapy that may enlist support from family members; however, some individuals will benefit from psychopharmacological interventions for improved outcomes and functioning. More common indications for psychotropic medication prescription in the pediatric setting include severe internalizing (ie, anxiety and depression) and neurodevelopmental difficulties (eg, ADHD) associated with significant psychosocial or functional impairment. Among patients with heart transplantation seen at a US pediatric institution and referred to a transplantation psychology/psychiatry service, 23% included a psychopharmacology consultation.⁸⁴

Consistent with pediatric patients, some adults with congenital heart disease and sustained severe psychological distress may also benefit from augmentation with pharmacotherapy, particularly in instances of psychotherapy failure. We are unaware of published trials of psychotropic medication prescription for adults with congenital heart disease. In a retrospective cohort study from a single-center adult congenital heart disease program in the United Kingdom, 3.3% of patients had documented use of antidepressants in their medical record, most often selective serotonin reuptake inhibitors. 104 Patients who had a prescription for antidepressant medication were more likely to be women, older and suffering from more complex congenital heart disease. After propensity score matching for variables potentially affecting risk of death, men but not women who required antidepressants were found to have 44% excess mortality (hazard ratio, 1.44 [95% CI, 1.17-1.84]). Causes of death were generally thought to be unrelated to antidepressant use but may be linked to the severity of the underlying cardiac condition that drove the need to initiate antidepressants. This possibility is further strengthened by the lack of evidence of increased arrhythmogenic or sudden cardiac death in the group receiving antidepressants. It is also possible that antidepressant medications may have been prescribed as part of palliative care, given increasing involvement of psychiatry during end-of-life care. 105 The low rate of antidepressant medication use in this British study (<5%) highlights the difference in prescribing patterns between countries. In a similar study from the United States, the rate of documented antidepressant use in adults with congenital heart disease was 18%.4

Table 2. Clinical Directions for Pediatric Cardiology and Adult Congenital Heart Disease Programs and Clinicians

Identify psychological care and outcomes as essential components of routine congenital heart disease care within one's program

Improve the ability of all clinicians (including physicians and nurses) to recognize psychological distress indicative of the need for clinical intervention

Collaborate with teachers, counselors, and educational systems to enhance neurodevelopmental, academic, and psychosocial outcomes

Address stigma and other barriers that may be associated with reluctance to seek mental health care

Establish guiding principles and metrics for assessing psychological functioning and other patient-reported outcomes, as well as best practices for coordinating mental health care across the life span

Integrate short-term assessment and treatment for health-related psychological challenges within congenital heart disease care teams, and develop pathways for longer-term mental health treatment within the community as appropriate

Expand the care model to address the psychological needs of families, including parents, caregivers, siblings, and spouses/partners

Promote peer mentorship programs for patients and family members

Commit to the provision of psychological care that is culturally sensitive and cognizant of social determinants of both physical and mental health

Ensure that mental health care is individualized to the cognitive abilities of patients, being mindful of the unique needs of those with genetic syndromes that affect neurodevelopmental and psychiatric outcomes

Develop strategies and business models for sustainable integration of mental health professionals within congenital heart disease teams, with particular attention to ensuring equity of access to care regardless of insurance coverage

Engage in advocacy and promote social justice initiatives to ensure inclusion and health equity and to reduce ableism

Table 1 summarizes general and congenital heart disease-specific considerations for each class of psychotropic medications. Among them, the most frequently used in pediatric patients with congenital heart disease are central nervous stimulant and nonstimulant medications targeting ADHD. Although there is no definite contraindication to the use of psychostimulants, antidepressants, mood stabilizers, or antipsychotics in individuals with congenital heart disease as a whole, 126 the safety profile of proposed drugs should be evaluated in the context of the medical history and symptoms of each patient. Despite the reported effects of stimulant and nonstimulant ADHD medications on heart rate and blood pressure, multicenter studies have shown that stimulant use was not associated with increased risk of serious cardiac events, 127,128 even in high-risk groups such as individuals with congenital heart disease,129 who otherwise have no specific arrhythmogenic background. 126 Given the evidence that inadequate treatment of ADHD is associated with significant risks, including increased accidents, academic failure, and substance misuse, 130 a risk-benefit analysis should inform treatment decisions. Pharmacological treatment for ADHD has been shown to improve psychosocial healthrelated quality of life over time, 131 and its use is considered safe in children with congenital heart disease. 132 However, studies are still needed to assess whether diagnosis and treatment of ADHD in school-aged children with congenital heart disease can improve their psychosocial well-being into adulthood.

Before psychotropic treatment is initiated, in addition to a psychological diagnostic evaluation, it is important to conduct a comprehensive medical evaluation, including physical and mental status examinations; history of and current cardiac symptomatology (eg, palpitations or syncope); and review of recent laboratory tests, electrocardiogram, imaging results, and current medications. Information about how long it takes for medications to reach full effectiveness and potential side effects should be shared with patients (and parents/caregivers in the pediatric setting).133 A particular challenge in the use of such medications in children with congenital heart disease is the difficulty in assessing QTc in patients with depolarization/repolarization abnormalities, as is frequently the case among patients with a history of operated congenital heart disease. Careful monitoring and close collaboration between pediatric and adult congenital heart disease cardiologists and psychiatrists have been suggested for best outcomes.¹²⁶ Pharmacists with expertise in cardiovascular medications are also excellent resources.

Summary

Although individuals with congenital heart disease face an elevated prevalence of clinically significant psychological distress, mental health treatment for them is uncommon. There is not a one-size-fits-all approach to mental health treatment; optimal care reflects the individual's needs, cognitive abilities, preferences, and priorities. There are multiple approaches to psychotherapy and options for pharmacotherapy, although empirical investigation in congenital heart disease has been sparse. Mental health professionals with experience and expertise in congenital heart disease could facilitate personalized treatment decisions for each patient. We thus strongly advocate for a culture shift in which mental health professionals are deemed essential members of congenital heart disease teams.

CONCLUSIONS AND FUTURE DIRECTIONS

Infants, children, adolescents, and adults with congenital heart disease experience a range of adversities that can threaten their physical, emotional, neurodevelopmental and social health. Furthermore, as a reflection of the bidirectional relationship between medical and psychological well-being, emerging evidence suggests that sustained, elevated psychological stress or distress may affect congenital heart disease morbidity and mortality. The Figure provides an illustrative summary of some of the antecedents and consequences of psychological distress in congenital heart disease. Much scope remains to deepen our understanding of the nature, prevalence, predictors, and trajectories of mental health outcomes and to increase opportunities for

Table 3. Priority Areas of Future Research Investigation to Understand and Improve Psychological Outcomes of Individuals Living With Congenital Heart Disease

How do fetal and developmental origins, including parent-child bonding and attachment, influence physical and mental health outcomes across the life span?

How do individual and environmental factors affect psychological trajectories, outcomes, and access to treatment?

What are the psychological outcomes among those who also have significant developmental or cognitive differences or disabilities? How can we most effectively improve outcomes?

What factors influence family mental health outcomes (often beginning in the prenatal period), as well as family functioning and well-being?

How do racism and sociocontextual factors affect psychological outcomes and access to treatment, and how can we reduce disparities in access to and outcomes of mental health care?

How do systems of health care influence patient and family mental health outcomes?

What is the best way to identify significant psychological distress in the clinical setting?

Does routine screening for psychological distress improve mental health outcomes?

What is the prevalence of acute and prolonged traumatic stress, and what are the risk and protective factors?

What factors contribute to psychological resilience and well-being?

What is the prevalence of and factors associated with self-harm and substance use disorders, and how can we appropriately target physical and mental health care interventions?

Can we improve medical outcomes (morbidity and mortality) by reducing clinically significant psychological distress?

What is the effectiveness and safety profile of psychotropic medications across age groups?

Which protocolized psychological interventions, tested through large-scale randomized controlled trials with validated assessments, are most effective in the short and long term?

How do we most effectively engage patients across all elements of the research process to ensure that assessment approaches and interventions are acceptable and meaningful?

How can we best offer personalized approaches to mental health interventions, which may include psychotherapy or pharmacotherapy, or both?

Is teletherapy an effective modality for mental health intervention?

What are the benefits and limitations of integration of mental health professionals within congenital heart disease teams?

intervention to improve psychological health and well-being. We recognize that positive psychological adaptation and good quality of life are reported by many individuals with congenital heart disease, and we do not wish to minimize observations of tremendous resilience. Rather, we wish to increase the prevalence of psychological well-being of individuals with congenital heart disease. Table 2 provides clinical suggestions for congenital heart disease clinicians and programs. Table 3 highlights research areas that we consider of highest priority. We advocate for genuine patient engagement in the development of both clinical and research endeavors to ensure that initiatives are acceptable, meaningful, and of high priority to patients and families.

Although psychological well-being is fundamental to living a healthy, productive life, the health care system often falls well short of identifying, addressing, and treating

psychological distress among those living with congenital heart disease. This mental health crisis is certainly not unique to individuals with congenital heart disease. The lifelong effects of psychological disorders and related difficulties come at great cost, not only to individuals' quality of life but also to families and communities, with the global financial impact of mental disorders measured in the trillions of dollars. 134 Within congenital heart disease, depression and anxiety are costly to the health care system and are associated with greater risk of death. Although the field of congenital cardiology is certainly not in a position to improve access to mental health care around the world, it is definitely within reach to improve psychological outcomes within the congenital heart disease community, and there is an urgency to do so. This demands prioritization of psychological outcomes by program, hospital and community leadership.

Despite decades of research investigating psychological outcomes of congenital heart disease and strong advocacy by patients and families, a catalyst is clearly needed to move initiatives forward in the clinical setting. We presume that all stakeholders, including patients, families, those who saved the lives of infants and children born with congenital heart disease, and those who take care of adult patients, are motivated to implement a care model that offers these patients the best possible quality of life. It is the goal of this scientific statement to spur efforts to include dedicated mental health professionals within all pediatric cardiology and adult congenital heart disease interdisciplinary teams. It is time to move from awareness to implementation.

ARTICLE INFORMATION

The American Heart Association makes every effort to avoid any actual or potential conflicts of interest that may arise as a result of an outside relationship or a personal, professional, or business interest of a member of the writing panel. Specifically, all members of the writing group are required to complete and submit a Disclosure Questionnaire showing all such relationships that might be perceived as real or potential conflicts of interest.

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†Significant.

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*Modest.

REFERENCES

- Gilboa SM, Devine OJ, Kucik JE, Oster ME, Riehle-Colarusso T, Nembhard WN, Xu P, Correa A, Jenkins K, Marelli AJ. Congenital heart defects in the United States: estimating the magnitude of the affected population in 2010. *Circulation*. 2016;134:101–109. doi: 10.1161/ CIRCULATIONAHA.115.019307
- Kovacs AH, Saidi AS, Kuhl EA, Sears SF, Silversides C, Harrison JL, Ong L, Colman J, Oechslin E, Nolan RP. Depression and anxiety in adult congenital heart disease: predictors and prevalence. *Int J Cardiol.* 2009;137:158–164. doi: 10.1016/j.ijcard.2008.06.042
- Westhoff-Bleck M, Briest J, Fraccarollo D, Hilfiker-Kleiner D, Winter L, Maske U, Busch MA, Bleich S, Bauersachs J, Kahl KG. Mental disorders in adults with congenital heart disease: unmet needs and impact on quality of life. J Affect Disord. 2016;204:180–186. doi: 10.1016/j.jad. 2016.06.047
- Carazo MR, Kolodziej MS, DeWitt ES, Kasparian NA, Newburger JW, Duarte VE, Singh MN, Opotowsky AR. Prevalence, and prognostic association of a clinical diagnosis of depression in adult congenital heart disease: results of the Boston Adult Congenital Heart Disease Biobank. *J Am Heart Assoc.* 2020;9:e014820. doi: 10.1161/JAHA.119.014820
- Benderly M, Kalter-Leibovici O, Weitzman D, Blieden L, Buber J, Dadashev A, Mazor-Dray E, Lorber A, Nir A, Yalonetsky S, et al; Israeli Congenital Heart Disease Research Group. Depression and anxiety are associated with high health care utilization and mortality among adults with congenital heart disease. *Int J Cardiol.* 2019;276:81–86. doi: 10.1016/j.ijcard.2018.09.005
- Marino BS, Lipkin PH, Newburger JW, Peacock G, Gerdes M, Gaynor JW, Mussatto KA, Uzark K, Goldberg CS, Johnson WH Jr, et al; on behalf of the American Heart Association Congenital Heart Defects Committee, Council on Cardiovascular Disease in the Young, Council on Cardiovascular Nursing, and Stroke Council. Neurodevelopmental outcomes in children with congenital heart disease: evaluation and management: a scientific statement from the American Heart Association. Circulation. 2012;126:1143–1172. doi: 10.1161/CIR.0b013e318265ee8a
- Kovacs AH, Bellinger DC. Neurocognitive and psychosocial outcomes in adult congenital heart disease: a lifespan approach. *Heart*. 2021;107:159– 167. doi: 10.1136/heartjnl-2016-310862
- Marelli A, Miller SP, Marino BS, Jefferson AL, Newburger JW. Brain in congenital heart disease across the lifespan: the cumulative burden of injury. *Circulation*. 2016;133:1951–1962. doi: 10.1161/ CIRCULATIONAHA.115.019881

- Shonkoff JP, Garner AS; Committee on Psychosocial Aspects of Child and Family Health; Committee on Early Childhood, Adoption, and Dependent Care; Section on Developmental and Behavioral Pediatrics. The lifelong effects of early childhood adversity and toxic stress. *Pediatrics*. 2012;129:e232–e246. doi: 10.1542/peds.2011-2663
- Karamlou T, Hawke JL, Zafar F, Kafle M, Tweddell JS, Najm HK, Frebis JR, Bryant RG 3rd. Widening our focus: characterizing socioeconomic and racial disparities in congenital heart disease. *Ann Thorac Surg.* 2022;113:157–165.
- Jackson JL, Morack J, Harris M, DeSalvo J, Daniels CJ, Chisolm DJ. Racial disparities in clinic follow-up early in life among survivors of congenital heart disease. *Congenit Heart Dis.* 2019;14:305–310. doi: 10.1111/chd.12732
- Lopez KN, Morris SA, Sexson Tejtel SK, Espaillat A, Salemi JL. US mortality attributable to congenital heart disease across the lifespan from 1999 through 2017 exposes persistent racial/ethnic disparities. *Circulation*. 2020;142:1132–1147. doi: 10.1161/CIRCULATIONAHA. 120.046822
- Peverill M, Dirks MA, Narvaja T, Herts KL, Comer JS, McLaughlin KA. Socioeconomic status and child psychopathology in the United States: A meta-analysis of population-based studies. Clin Psychol Rev. 2021;83:101933. doi: 10.1016/j.cpr.2020.101933
- Fryers T, Melzer D, Jenkins R. Social inequalities and the common mental disorders: a systematic review of the evidence. Soc Psychiatry Psychiatr Epidemiol. 2003;38:229–237. doi: 10.1007/s00127-003-0627-2
- Drakouli M, Petsios K, Giannakopoulou M, Patiraki E, Voutoufianaki I, Matziou V. Determinants of quality of life in children and adolescents with CHD: a systematic review. *Cardiol Young*. 2015;25:1027–1036. doi: 10.1017/S1047951115000086
- Kahr PC, Radke RM, Orwat S, Baumgartner H, Diller GP. Analysis of associations between congenital heart defect complexity and health-related quality of life using a meta-analytic strategy. *Int J Cardiol.* 2015;199:197–203. doi: 10.1016/j.ijcard.2015.07.045
- Latal B, Helfricht S, Fischer JE, Bauersfeld U, Landolt MA. Psychological adjustment and quality of life in children and adolescents following openheart surgery for congenital heart disease: a systematic review. BMC Pediatr. 2009;9:6. doi: 10.1186/1471-2431-9-6
- McCrindle BW, Williams RV, Mitchell PD, Hsu DT, Paridon SM, Atz AM, Li JS, Newburger JW; Pediatric Heart Network Investigators. Relationship of patient and medical characteristics to health status in children and adolescents after the Fontan procedure. *Circulation*. 2006;113:1123–1129. doi: 10.1161/CIRCULATIONAHA.105.576660

- Neal AE, Stopp C, Wypij D, Bellinger DC, Dunbar-Masterson C, DeMaso DR, Newburger JW. Predictors of health-related quality of life in adolescents with tetralogy of Fallot. J Pediatr. 2015;166:132–138. doi: 10.1016/j.jpeds.2014.09.034
- DeMaso DR, Labella M, Taylor GA, Forbes PW, Stopp C, Bellinger DC, Rivkin MJ, Wypij D, Newburger JW. Psychiatric disorders and function in adolescents with d-transposition of the great arteries. *J Pediatr.* 2014;165:760–766. doi: 10.1016/j.jpeds.2014.06.029
- Wilson WM, Smith-Parrish M, Marino BS, Kovacs AH. Neurodevelopmental and psychosocial outcomes across the congenital heart disease lifespan. Prog Pediatr Cardiol. 2015;39(pt B):113–118.
- Verrall C, Blue G, Loughran-Fowlds A, Kasparian NA, Walker K, Dunwoodie S, Sholler GF, Badawi N, Winlaw DS. "Big issues" in neurodevelopment in congenital heart disease. *Open Heart*. 2019;6:e000998. doi: 10.1136/openhrt-2018-000998
- Dunbar-Masterson C, Wypij D, Bellinger DC, Rappaport LA, Baker AL, Jonas RA, Newburger JW. General health status of children with D-transposition of the great arteries after the arterial switch operation. *Circulation*. 2001;104(suppl 1):1138–1142. doi: 10.1161/hc37t1.094782
- Robson VK, Stopp C, Wypij D, Dunbar-Masterson C, Bellinger DC, DeMaso DR, Rappaport LA, Newburger JW. Longitudinal associations between neurodevelopment and psychosocial health status in patients with repaired D-transposition of the great arteries. *J Pediatr.* 2019;204:38– 45.e1. doi: 10.1016/j.jpeds.2018.08.069
- Matza LS, Rentz AM, Secnik K, Swensen AR, Revicki DA, Michelson D, Spencer T, Newcorn JH, Kratochvil CJ. The link between health-related quality of life and clinical symptoms among children with attention-deficit hyperactivity disorder. J Dev Behav Pediatr. 2004;25:166–174. doi: 10.1097/00004703-200406000-00005
- Brook JS, Brook DW, Zhang C, Seltzer N, Finch SJ. Adolescent ADHD and adult physical and mental health, work performance, and financial stress. *Pediatrics*. 2013;131:5–13. doi: 10.1542/peds.2012-1725
- Bussing R, Mason DM, Bell L, Porter P, Garvan C. Adolescent outcomes of childhood attention-deficit/hyperactivity disorder in a diverse community sample. J Am Acad Child Adolesc Psychiatry. 2010;49:595–605. doi: 10.1016/j.jaac.2010.03.006
- Barbaresi WJ, Colligan RC, Weaver AL, Voigt RG, Killian JM, Katusic SK. Mortality, ADHD, and psychosocial adversity in adults with childhood ADHD: a prospective study. *Pediatrics*. 2013;131:637–644. doi: 10.1542/peds.2012-2354
- Jackson JL, Gerardo GM, Monti JD, Schofield KA, Vannatta K. Executive function and internalizing symptoms in adolescents and young adults with congenital heart disease: the role of coping. *J Pediatr Psychol.* 2018;43:906–915. doi: 10.1093/jpepsy/jsx154
- Monti JD, Jackson JL, Vannatta K. Adolescents and young adults living with congenital heart disease: coping, stress reactivity, and emotional functioning. J Clin Psychol Med Settings. 2018;25:441–451. doi: 10.1007/s10880-018-9554-y
- Massaro AN, Glass P, Brown J, Chang T, Krishnan A, Jonas RA, Donofrio MT. Neurobehavioral abnormalities in newborns with congenital heart disease requiring open-heart surgery. *J Pediatr.* 2011;158:678–681.e2. doi: 10.1016/j.jpeds.2010.11.060
- Limperopoulos C, Majnemer A, Shevell MI, Rosenblatt B, Rohlicek C, Tchervenkov C. Neurodevelopmental status of newborns and infants with congenital heart defects before and after open heart surgery. *J Pediatr*. 2000;137:638-645. doi: 10.1067/mpd.2000.109152
- Carbajal R, Rousset A, Danan C, Coquery S, Nolent P, Ducrocq S, Saizou C, Lapillonne A, Granier M, Durand P, et al. Epidemiology and treatment of painful procedures in neonates in intensive care units. *JAMA*. 2008;300:60–70. doi: 10.1001/jama.300.1.60
- Kasparian NA, Kan JM, Sood E, Wray J, Pincus HA, Newburger JW. Mental health care for parents of babies with congenital heart disease during intensive care unit admission: systematic review and statement of best practice. *Early Hum Dev.* 2019;139:104837. doi: 10.1016/j. earlhumdev.2019.104837
- Corbett SS, Drewett RF. To what extent is failure to thrive in infancy associated with poorer cognitive development? A review and metaanalysis. J Child Psychol Psychiatry. 2004;45:641–654. doi: 10.1111/j. 1469-7610.2004.00253.x
- Pickles DM, Lihn SL, Boat TF, Lannon C. A roadmap to emotional health for children and families with chronic pediatric conditions. *Pediatrics*. 2020;145:e20191324. doi: 10.1542/peds.2019-1324
- 37. Denniss DL, Sholler GF, Costa DSJ, Winlaw DS, Kasparian NA. Need for routine screening of health-related quality of life in families of young

- children with complex congenital heart disease. J Pediatr. 2019;205:21–28.e2. doi: 10.1016/j.jpeds.2018.09.037
- Re JM, Dean S, Mullaert J, Guedeney A, Menahem S. Maternal distress and infant social withdrawal (ADBB) following infant cardiac surgery for congenital heart disease. World J Pediatr Congenit Heart Surg. 2018;9:624– 637. doi: 10.1177/2150135118788788
- Abda A, Bolduc ME, Tsimicalis A, Rennick J, Vatcher D, Brossard-Racine M. Psychosocial outcomes of children and adolescents with severe congenital heart defect: a systematic review and meta-analysis. *J Pediatr Psychol.* 2019;44:463–477. doi: 10.1093/jpepsy/jsy085
- Girouard HS, Kovacs AH. Congenital heart disease: education and employment considerations and outcomes. Int J Cardiol Congenit Heart Dis. 2020;1:100005.
- DeMaso DR, Calderon J, Taylor GA, Holland JE, Stopp C, White MT, Bellinger DC, Rivkin MJ, Wypij D, Newburger JW. Psychiatric disorders in adolescents with single ventricle congenital heart disease. *Pediatrics*. 2017;139:e20162241. doi: 10.1542/peds.2016-2241
- Holland JE, Cassidy AR, Stopp C, White MT, Bellinger DC, Rivkin MJ, Newburger JW, DeMaso DR. Psychiatric disorders and function in adolescents with tetralogy of Fallot. J Pediatr. 2017;187:165–173. doi: 10.1016/j. ipeds.2017.04.048
- Marshall KH, D'Udekem Y, Sholler GF, Opotowsky AR, Costa DSJ, Sharpe L, Celermajer DS, Winlaw DS, Newburger JW, Kasparian NA. Health-related quality of life in children, adolescents, and adults with a Fontan circulation: a meta-analysis. *J Am Heart Assoc.* 2020;9:e014172. doi: 10.1161/JAHA.119.014172
- Loccoh EC, Yu S, Donohue J, Lowery R, Butcher J, Pasquali SK, Goldberg CS, Uzark K. Prevalence and risk factors associated with non-attendance in neurodevelopmental follow-up clinic among infants with CHD. Cardiol Young. 2018;28:554–560. doi: 10.1017/S1047951117002748
- 45. Chong LSH, Fitzgerald DA, Craig JC, Manera KE, Hanson CS, Celermajer D, Ayer J, Kasparian NA, Tong A. Children's experiences of congenital heart disease: a systematic review of qualitative studies. Eur J Pediatr. 2018;177:319–336. doi: 10.1007/s00431-017-3081-y
- Kovacs AH, Utens EM. More than just the heart: transition and psychosocial issues in adult congenital heart disease. *Cardiol Clin.* 2015;33:625–34, ix. doi: 10.1016/j.ccl.2015.07.005
- Luyckx K, Goossens E, Rassart J, Apers S, Vanhalst J, Moons P. Parental support, internalizing symptoms, perceived health status, and quality of life in adolescents with congenital heart disease: influences and reciprocal effects. J Behav Med. 2014;37:145–155. doi: 10.1007/s10865-012-9474-5
- 48. Sable C, Foster E, Uzark K, Bjornsen K, Canobbio MM, Connolly HM, Graham TP, Gurvitz MZ, Kovacs A, Meadows AK, et al; on behalf of the American Heart Association Congenital Heart Defects Committee of the Council on Cardiovascular Disease in the Young, Council on Cardiovascular Nursing, Council on Clinical Cardiology, and Council on Peripheral Vascular Disease. Best practices in managing transition to adulthood for adolescents with congenital heart disease: the transition process and medical and psychosocial issues: a scientific statement from the American Heart Association. Circulation. 2011;123:1454–1485. doi: 10.1161/CIR.0b013e3182107c56
- Kongkaew C, Jampachaisri K, Chaturongkul CA, Scholfield CN. Depression and adherence to treatment in diabetic children and adolescents: a systematic review and meta-analysis of observational studies. *Eur J Pediatr*. 2014;173:203–212. doi: 10.1007/s00431-013-2128-y
- Jackson JL, Gerardo GM, Daniels CJ, Vannatta K. Perceptions of disease-related stress: a key to better understanding patient-reported outcomes among survivors of congenital heart disease. *J Cardiovasc Nurs*. 2017;32:587–593. doi: 10.1097/JCN.0000000000000371
- Tillman AR, Colborn KL, Scott KA, Davidson AJ, Khanna A, Kao D, McKenzie L, Ong T, Rausch CM, Duca LM, et al. Associations between socioeconomic context and congenital heart disease related outcomes in adolescents and adults. *Am J Cardiol.* 2021;139:105–115. doi: 10.1016/j.amjcard.2020.10.040
- Roseman A, Morton L, Kovacs AH. Health anxiety among adults with congenital heart disease. Curr Opin Cardiol. 2021;36:98–104. doi: 10.1097/HC0.0000000000000811
- Krnjacki L, Priest N, Aitken Z, Emerson E, Llewellyn G, King T, Kavanagh A. Disability-based discrimination and health: findings from an Australian-based population study. Aust NZ J Public Health. 2018;42:172– 174. doi: 10.1111/1753-6405.12735
- Morton L, Cogan N, Kornfält S, Porter Z, Georgiadis E. Baring all: the impact of the hospital gown on patient well-being. Br J Health Psychol. 2020;25:452–473. doi: 10.1111/bjhp.12416
- 55. Gurvitz M, Valente AM, Broberg C, Cook S, Stout K, Kay J, Ting J, Kuehl K, Earing M, Webb G, et al; Alliance for Adult Research in Congenital Cardiology

- (AARCC) and Adult Congenital Heart Association. Prevalence and predictors of gaps in care among adult congenital heart disease patients: HEART-ACHD (the Health, Education, and Access Research Trial). *J Am Coll Cardiol.* 2013;61:2180–2184. doi: 10.1016/j.jacc.2013.02.048
- 56. Bhatt AB, Foster E, Kuehl K, Alpert J, Brabeck S, Crumb S, Davidson WR Jr, Earing MG, Ghoshhajra BB, Karamlou T, et al; on behalf of the American Heart Association Council on Clinical Cardiology. Congenital heart disease in the older adult: a scientific statement from the American Heart Association [published correction appears in Circulation. 2015;131:e510]. Circulation. 2015;131:1884–1931. doi: 10.1161/CIR.00000000000000204
- Bagge CN, Henderson VW, Laursen HB, Adelborg K, Olsen M, Madsen NL. Risk of dementia in adults with congenital heart disease: populationbased cohort study. *Circulation*. 2018;137:1912–1920. doi: 10.1161/ CIRCULATIONAHA.117.029686
- Jha MK, Qamar A, Vaduganathan M, Charney DS, Murrough JW. Screening and management of depression in patients with cardiovascular disease: *JACC* state-of-the-art review. *J Am Coll Cardiol*. 2019;73:1827–1845. doi: 10.1016/j.jacc.2019.01.041
- Celano CM, Millstein RA, Bedoya CA, Healy BC, Roest AM, Huffman JC. Association between anxiety and mortality in patients with coronary artery disease: a meta-analysis. *Am Heart J.* 2015;170:1105–1115. doi:10.1016/j. ahj.2015.09.013
- Celano CM, Villegas AC, Albanese AM, Gaggin HK, Huffman JC. Depression and anxiety in heart failure: a review. Harv Rev Psychiatry. 2018;26:175– 184. doi: 10.1097/HRP.000000000000162
- Liguori I, Russo G, Curcio F, Sasso G, Della-Morte D, Gargiulo G, Pirozzi F, Cacciatore F, Bonaduce D, Abete P, et al. Depression and chronic heart failure in the elderly: an intriguing relationship. *J Geriatr Cardiol*. 2018;15:451– 459. doi: 10.11909/j.issn.1671-5411.2018.06.014
- Williams DR, Mohammed SA. Racism and health I: pathways and scientific evidence. Am Behav Sci. 2013;57:10.1177/0002764213487340. doi: 10.1177/0002764213487340
- Cobbinah SS, Lewis J. Racism & health: a public health perspective on racial discrimination. J Eval Clin Pract. 2018;24:995–998. doi: 10.1111/jep.12894
- Salciccioli KB, Oluyomi A, Lupo PJ, Ermis PR, Lopez KN. A model for geographic and sociodemographic access to care disparities for adults with congenital heart disease. *Congenit Heart Dis.* 2019;14:752–759. doi: 10.1111/chd.12819
- 65. Jackson JL, Leslie CE, Hondorp SN. Depressive and anxiety symptoms in adult congenital heart disease: prevalence, health impact and treatment. *Prog Cardiovasc Dis.* 2018;61:294–299. doi: 10.1016/j.pcad.2018.07.015
- Kasmi L, Calderon J, Montreuil M, Geronikola N, Lambert V, Belli E, Bonnet D, Kalfa D. Neurocognitive and psychological outcomes in adults with dextro-transposition of the great arteries corrected by the arterial switch operation. *Ann Thorac Surg.* 2018;105:830–836. doi: 10.1016/j.athoracsur.2017.06.055
- Kim MY, Johnson JL, Sawatzky R. Relationship between types of social support, coping strategies, and psychological distress in individuals living with congenital heart disease. *J Cardiovasc Nurs*. 2019;34:76–84. doi: 10.1097/JCN.00000000000000531
- 68. Ko JM, White KS, Kovacs AH, Tecson KM, Apers S, Luyckx K, Thomet C, Budts W, Enomoto J, Sluman MA, et al; APPROACH-IS consortium and International Society for Adult Congenital Heart Disease (ISACHD). Differential impact of physical activity type on depression in adults with congenital heart disease: a multi-center international study. J Psychosom Res. 2019;124:109762. doi: 10.1016/j.jpsychores.2019.109762
- Deng LX, Khan AM, Drajpuch D, Fuller S, Ludmir J, Mascio CE, Partington SL, Qadeer A, Tobin L, Kovacs AH, et al. Prevalence and correlates of post-traumatic stress disorder in adults with congenital heart disease. *Am J Cardiol.* 2016;117:853–857. doi: 10.1016/j.amjcard.2015.11.065
- Sagar A, Bishop JR, Tessman DC, Guter S, Martin CL, Cook EH. Cooccurrence of autism, childhood psychosis, and intellectual disability associated with a de novo 3q29 microdeletion. Am J Med Genet A. 2013;161A:845–849. doi: 10.1002/ajmg.a.35754
- 71. Yi JJ, Tang SX, McDonald-McGinn DM, Calkins ME, Whinna DA, Souders MC, Zackai EH, Goldmuntz E, Gaynor JW, Gur RC, et al. Contribution of congenital heart disease to neuropsychiatric outcome in school-age children with 22q11.2 deletion syndrome. Am J Med Genet B Neuropsychiatr Genet. 2014;165B:137-147. doi: 10.1002/ajmg.b.32215
- Goldmuntz E, Clark BJ, Mitchell LE, Jawad AF, Cuneo BF, Reed L, McDonald-McGinn D, Chien P, Feuer J, Zackai EH, et al. Frequency of 22q11 deletions in patients with conotruncal defects. *J Am Coll Cardiol*. 1998;32:492–498. doi: 10.1016/s0735-1097(98)00259-9

- Fung WL, Butcher NJ, Costain G, Andrade DM, Boot E, Chow EW, Chung B, Cytrynbaum C, Faghfoury H, Fishman L, et al. Practical guidelines for managing adults with 22q11.2 deletion syndrome. *Genet Med.* 2015;17:599–609. doi: 10.1038/gim.2014.175
- Santoro SL, Steffensen EH. Congenital heart disease in Down syndrome: a review of temporal changes. *J Congenit Cardiol.* 2021;5:1. doi: 10.1186/s40949-020-00055-7
- Windsperger K, Hoehl S. Development of Down syndrome research over the last decades: what healthcare and education professionals need to know. Front Psychiatry. 2021;12:749046. doi: 10.3389/fpsyt.2021.749046
- Kim YY, Gauvreau K, Bacha EA, Landzberg MJ, Benavidez OJ. Resource use among adult congenital heart surgery admissions in pediatric hospitals: risk factors for high resource utilization and association with inpatient death. Circ Cardiovasc Qual Outcomes. 2011;4:634–639. doi: 10.1161/CIRCOUTCOMES.111.963223
- Gonzalez VJ, Kimbro RT, Cutitta KE, Shabosky JC, Bilal MF, Penny DJ, Lopez KN. Mental health disorders in children with congenital heart disease. *Pediatrics*. 2021;147:e20201693. doi: 10.1542/peds.2020-1693
- Callus E, Pravettoni G. The role of clinical psychology and peer to peer support in the management of chronic medical conditions: a practical example with adults with congenital heart disease. Front Psychol. 2018;9:731. doi: 10.3389/fpsyg.2018.00731
- Desai PP, Sutton LJ, Staley MD, Hannon DW. A qualitative study exploring the psychosocial value of weekend camping experiences for children and adolescents with complex heart defects. *Child Care Health Dev.* 2014;40:553–561. doi: 10.1111/cch.12056
- Morton L. Using psychologically informed care to improve mental health and wellbeing for people living with a heart condition from birth: a statement paper. J Health Psychol. 2020;25:197–206. doi: 10.1177/1359105319826354
- Kasparian NA, Winlaw DS, Sholler GF. "Congenital heart health": how psychological care can make a difference. *Med J Aust.* 2016;205:104–107. doi: 10.5694/mja16.00392
- Brosig C, Yang K, Hoffmann RG, Dasgupta M, Mussatto K. The role of psychology in a pediatric outpatient cardiology setting: preliminary results from a new clinical program. *J Clin Psychol Med Settings*. 2014;21:337–346. doi: 10.1007/s10880-014-9404-5
- Ferguson M, Kovacs AH. An integrated adult congenital heart disease psychology service. Congenit Heart Dis. 2016;11:444-451. doi: 10.1111/chd.12331
- Thomson K, McKenna K, Bedard-Thomas K, Oliva M, Ibeziako P. Behavioral health care in solid organ transplantation in a pediatric setting. *Pediatr Transplant*. 2018;22:e13217. doi: 10.1111/petr.13217
- Pagé MG, Kovacs AH, Irvine J. How do psychosocial challenges associated with living with congenital heart disease translate into treatment interests and preferences? A qualitative approach. *Psychol Health*. 2012;27:1260– 1270. doi: 10.1080/08870446.2012.667099
- Kovacs AH, Bendell KL, Colman J, Harrison JL, Oechslin E, Silversides C. Adults with congenital heart disease: psychological needs and treatment preferences. *Congenit Heart Dis.* 2009;4:139–146. doi: 10.1111/j. 1747-0803.2009.00280.x
- Crowe K, McKay D. Efficacy of cognitive-behavioral therapy for child-hood anxiety and depression. *J Anxiety Disord*. 2017;49:76–87. doi: 10.1016/j.janxdis.2017.04.001
- Zhang A, Borhneimer LA, Weaver A, Franklin C, Hai AH, Guz S, Shen L. Cognitive behavioral therapy for primary care depression and anxiety: a secondary meta-analytic review using robust variance estimation in meta-regression. *J Behav Med.* 2019;42:1117–1141. doi: 10.1007/s10865-019-00046-z
- Reavell J, Hopkinson M, Clarkesmith D, Lane DA. Effectiveness of cognitive behavioral therapy for depression and anxiety in patients with cardiovascular disease: a systematic review and meta-analysis. *Psychosom Med.* 2018;80:742–753. doi: 10.1097/PSY.0000000000000626
- Kovacs AH, Grace SL, Kentner AC, Nolan RP, Silversides CK, Irvine MJ. Feasibility and outcomes in a pilot randomized controlled trial of a psychosocial intervention for adults with congenital heart disease. *Can J Cardiol.* 2018;34:766–773. doi: 10.1016/j.cjca.2018.02.023
- Rashidi A, Whitehead L, Newson L, Astin F, Gill P, Lane DA, Lip GYH, Neubeck L, Ski CF, Thompson DR, et al. The role of acceptance and commitment therapy in cardiovascular and diabetes healthcare: a scoping review. *Int J Environ Res Public Health*. 2021;18:8126. doi: 10.3390/ijerph18158126
- Charney ME, Hellberg SN, Bui E, Simon NM. Evidenced-based treatment of posttraumatic stress disorder: an updated review of validated psychotherapeutic and pharmacological approaches. *Harv Rev Psychiatry*. 2018;26:99– 115. doi: 10.1097/HRP.000000000000186

- 93. Mavranezouli I, Megnin-Viggars O, Daly C, Dias S, Stockton S, Meiser-Stedman R, Trickey D, Pilling S. Research review: psychological and psychosocial treatments for children and young people with post-traumatic stress disorder: a network meta-analysis. *J Child Psychol Psychiatry*. 2020;61:18–29. doi: 10.1111/jcpp.13094
- 94. Wilson G, Farrell D, Barron I, Hutchins J, Whybrow D, Kiernan MD. The use of eye-movement desensitization reprocessing (EMDR) therapy in treating post-traumatic stress disorder: a systematic narrative review. Front Psychol. 2018;9:923. doi: 10.3389/fpsyg.2018.00923
- Scott-Sheldon LAJ, Gathright EC, Donahue ML, Balletto B, Feulner MM, DeCosta J, Cruess DG, Wing RR, Carey MP, Salmoirago-Blotcher E. Mindfulness-based interventions for adults with cardiovascular disease: a systematic review and meta-analysis. *Ann Behav Med.* 2020;54:67–73. doi: 10.1093/abm/kaz020
- Zilberstein K. The use and limitations of attachment theory in child psychotherapy. Psychotherapy (Chic). 2014;51:93–103. doi: 10.1037/a0030930
- Tesson S, Butow PN, Marshall K, Fonagy P, Kasparian NA. Parent-child bonding and attachment during pregnancy and early childhood following congenital heart disease diagnosis [published online June 8, 2021]. Health Psychol Rev. doi: 10.1080/17437199.2021.1927136. https://www.tandfonline.com/doi/abs/10.1080/17437199. 2021.1927136
- Steinert C, Munder T, Rabung S, Hoyer J, Leichsenring F. Psychodynamic therapy: as efficacious as other empirically supported treatments? A Metaanalysis testing equivalence of outcomes. *Am J Psychiatry*. 2017;174:943– 953. doi: 10.1176/appi.ajp.2017.17010057
- Kibby MY, Tyc VL, Mulhern RK. Effectiveness of psychological intervention for children and adolescents with chronic medical illness: a meta-analysis. Clin Psychol Rev. 1998;18:103–117. doi:10.1016/s0272-7358(97)00049-4
- Law E, Fisher E, Eccleston C, Palermo TM. Psychological interventions for parents of children and adolescents with chronic illness. *Cochrane Database* Syst Rev. 2019;3:CD009660. doi: 10.1002/14651858.CD009660.pub4
- 101. Tesson S, Butow PN, Sholler GF, Sharpe L, Kovacs AH, Kasparian NA. Psychological interventions for people affected by childhood-onset heart disease: a systematic review. *Health Psychol*. 2019;38:151–161. doi: 10.1037/hea0000704
- 102. Richards SH, Anderson L, Jenkinson CE, Whalley B, Rees K, Davies P, Bennett P, Liu Z, West R, Thompson DR, et al. Psychological interventions for coronary heart disease: Cochrane systematic review and meta-analysis. Eur J Prev Cardiol. 2018;25:247–259. doi: 10.1177/2047487317739978
- Lamb T, Pachana NA, Dissanayaka N. Update of recent literature on remotely delivered psychotherapy interventions for anxiety and depression. *Telemed J E Health*. 2019;25:671–677. doi: 10.1089/tmj.2018.0079
- 104. Diller GP, Bräutigam A, Kempny A, Uebing A, Alonso-Gonzalez R, Swan L, Babu-Narayan SV, Baumgartner H, Dimopoulos K, Gatzoulis MA. Depression requiring anti-depressant drug therapy in adult congenital heart disease: prevalence, risk factors, and prognostic value. Eur Heart J. 2016;37:771–782. doi: 10.1093/eurheartj/ehv386
- 105. Samsel C, Tapsak S, Thomson K, McKenna K, McGregor K, Forbes P, Ibeziako P. Psychotropic medication use trends in a large pediatric and young adult solid organ transplant population. *Pediatr Transplant*. 2019;23:e13380. doi: 10.1111/petr.13380
- 106. Kennedy SH, Lam RW, McIntyre RS, Tourjman SV, Bhat V, Blier P, Hasnain M, Jollant F, Levitt AJ, MacQueen GM, et al; CANMAT Depression Work Group. Canadian Network for Mood and Anxiety Treatments (CANMAT) 2016 clinical guidelines for the management of adults with major depressive disorder: section 3: pharmacological treatments. Can J Psychiatry. 2016;61:540–560. doi: 10.1177/0706743716659417
- 107. Beach SR, Kostis WJ, Celano CM, Januzzi JL, Ruskin JN, Noseworthy PA, Huffman JC. Meta-analysis of selective serotonin reuptake inhibitor-associated QTc prolongation. J Clin Psychiatry. 2014;75:e441-e449. doi: 10.4088/JCP.13r08672
- Brouillette J, Nattel S. A practical approach to avoiding cardiovascular adverse effects of psychoactive medications. Can J Cardiol. 2017;33:1577–1586. doi: 10.1016/j.cjca.2017.09.001
- 109. Fanoe S, Kristensen D, Fink-Jensen A, Jensen HK, Toft E, Nielsen J, Videbech P, Pehrson S, Bundgaard H. Risk of arrhythmia induced by psychotropic medications: a proposal for clinical management. *Eur Heart J.* 2014;35:1306–1315. doi: 10.1093/eurheartj/ehu100
- 110. Katzman MA, Bleau P, Blier P, Chokka P, Kjernisted K, Van Ameringen M, Antony MM, Bouchard S, Brunet A, Flament M, et al; Canadian Anxiety Guidelines Initiative Group on behalf of the Anxiety Disorders Association

- of Canada/Association Canadienne des troubles anxieux and McGill University. Canadian clinical practice guidelines for the management of anxiety, posttraumatic stress and obsessive-compulsive disorders. *BMC Psychiatry*. 2014;14(suppl 1):S1. doi: 10.1186/1471-244X-14-S1-S1
- 111. Reilly JG, Ayis SA, Ferrier IN, Jones SJ, Thomas SH. QTc-interval abnormalities and psychotropic drug therapy in psychiatric patients. *Lancet*. 2000;355:1048–1052. doi: 10.1016/s0140-6736(00)02035-3
- 112. Mehta N, Vannozzi R. Lithium-induced electrocardiographic changes: a complete review. *Clin Cardiol.* 2017;40:1363–1367. doi: 10.1002/clc.22822
- Puzantian TC, Carlat DJ. Medication Fact Book for Psychiatric Practice. Carlat Publishing; 2016:170.
- 114. French JA, Perucca E, Sander JW, Bergfeldt L, Baulac M, Auerbach DS, Keezer M, Thijs RD, Devinsky O, Vossler DG, et al. FDA safety warning on the cardiac effects of lamotrigine: an advisory from the Ad Hoc ILAE/AES Task Force. *Epilepsy Curr.* 2021;21:1535759721996344. doi: 10.1177/1535759721996344
- 115. Biederman J, Melmed RD, Patel A, McBurnett K, Konow J, Lyne A, Scherer N; SPD503 Study Group. A randomized, double-blind, place-bo-controlled study of guanfacine extended release in children and adolescents with attention-deficit/hyperactivity disorder. *Pediatrics*. 2008;121:e73-e84. doi: 10.1542/peds.2006-3695
- 116. Kollins SH, Jain R, Brams M, Segal S, Findling RL, Wigal SB, Khayrallah M. Clonidine extended-release tablets as add-on therapy to psychostimulants in children and adolescents with ADHD. *Pediatrics*. 2011;127:e1406–e1413. doi: 10.1542/peds.2010-1260
- 117. Mick E, McManus DD, Goldberg RJ. Meta-analysis of increased heart rate and blood pressure associated with CNS stimulant treatment of ADHD in adults. *Eur Neuropsychopharmacol*. 2013;23:534–541. doi: 10.1016/j.euroneuro.2012.06.011
- 118. Priori SG, Blomström-Lundqvist C, Mazzanti A, Blom N, Borggrefe M, Camm J, Elliott PM, Fitzsimons D, Hatala R, Hindricks G, et al; ESC Scientific Document Group. 2015 ESC guidelines for the management of patients with ventricular arrhythmias and the prevention of sudden cardiac death: the Task Force for the Management of Patients with Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death of the European Society of Cardiology (ESC). Eur Heart J. 2015;36:2793–2867. doi: 10.1093/eurhearti/ehv316
- 119. Zhang C, Kutyifa V, Moss AJ, McNitt S, Zareba W, Kaufman ES. Long-QT syndrome and therapy for attention deficit/hyperactivity disorder. J Cardiovasc Electrophysiol. 2015;26:1039–1044. doi: 10.1111/jce.12739
- 120. Goodnick PJ, Jerry J, Parra F. Psychotropic drugs and the ECG: focus on the QTc interval. Expert Opin Pharmacother. 2002;3:479–498. doi: 10.1517/14656566.3.5.479
- McIntyre RS, Jerrell JM. Metabolic and cardiovascular adverse events associated with antipsychotic treatment in children and adolescents. *Arch Pediatr Adolesc Med.* 2008;162:929–935. doi: 10.1001/archpedi.162.10.929
- 122. Centers for Medicare and Medicaid Services. Antidepressant medications: U.S. Food and Drug Administration-approved indications and dosages for use in pediatric patients. Accessed July 7, 2020. https://www.cms.gov/ Medicare-Medicaid-Coordination/Fraud-Prevention/Medicaid-Integrity-Education/Pharmacy-Education-Materials/Downloads/ad-pediatric-dosingchart11-14.pdf
- 123. Centers for Medicare and Medicaid Services. Stimulant and related medications: use in pediatric patients. Accessed July 7, 2020. https://www.cms.gov/Medicare-Medicaid-Coordination/Fraud-Prevention/Medicaid-Integrity-Education/Pharmacy-Education-Materials/Downloads/stim-pediatric-factsheet11-14.pdf
- 124. Lorberg B, Robb A, Pavuluri M, Chen DT, Wilens T. Pediatric psychopharmacology: food and drug administration approval through the evidence lens. J Am Acad Child Adolesc Psychiatry. 2014;53:716-719. doi:10.1016/j. jaac.2014.04.015
- 125. Centers for Medicare and Medicaid Services. Atypical antipsychotic medications: use in pediatric patients. Accessed July 7, 2020. https://www.cms.gov/Medicare-Medicaid-Coordination/Fraud-Prevention/Medicaid-Integrity-Education/Pharmacy-Education-Materials/Downloads/atyp-antipsych-pediatric-factsheet11-14.pdf
- 126. Vetter VL, Elia J, Erickson C, Berger S, Blum N, Uzark K, Webb CL. Cardiovascular monitoring of children and adolescents with heart disease receiving medications for attention deficit/hyperactivity disorder: a scientific statement from the American Heart Association Council on Cardiovascular Disease in the Young Congenital Cardiac Defects Committee and the Council on Cardiovascular Nursing [published correction appears in Circulation. 2009;120:e55–e59]. Circulation. 2008;117:2407–2423. doi: 10.1161/CIRCULATIONAHA.107.189473

- 127. Cooper WO, Habel LA, Sox CM, Chan KA, Arbogast PG, Cheetham TC, Murray KT, Quinn VP, Stein CM, Callahan ST, et al. ADHD drugs and serious cardiovascular events in children and young adults. N Engl J Med. 2011;365:1896-1904. doi: 10.1056/NEJMoa1110212
- 128. Habel LA, Cooper WO, Sox CM, Chan KA, Fireman BH, Arbogast PG, Cheetham TC, Quinn VP, Dublin S, Boudreau DM, et al. ADHD medications and risk of serious cardiovascular events in young and middle-aged adults. JAMA. 2011;306:2673–2683. doi: 10.1001/jama.2011.1830
- 129. Winterstein AG, Gerhard T, Kubilis P, Saidi A, Linden S, Crystal S, Zito J, Shuster JJ, Olfson M. Cardiovascular safety of central nervous system stimulants in children and adolescents: population based cohort study. BMJ. 2012;345:e4627. doi: 10.1136/bmj.e4627
- Able SL, Johnston JA, Adler LA, Swindle RW. Functional and psychosocial impairment in adults with undiagnosed ADHD. *Psychol Med.* 2007;37:97– 107. doi: 10.1017/S0033291706008713
- 131. Newcorn JH, Spencer TJ, Biederman J, Milton DR, Michelson D. Atomoxetine treatment in children and adolescents with attention-deficit/ hyperactivity disorder and comorbid oppositional defiant disorder. *J Am Acad Child Adolesc Psychiatry*. 2005;44:240–248. doi: 10.1097/00004583-200503000-00008
- 132. Batra AS, Alexander ME, Silka MJ. Attention-deficit/hyperactivity disorder, stimulant therapy, and the patient with congenital heart disease: evidence and reason. *Pediatr Cardiol.* 2012;33:394–401. doi: 10.1007/s00246-012-0162-6
- 133. Miller L, Campo JV. Depression in adolescents. *N Engl J Med.* 2021;385:445–449. doi: 10.1056/NEJMra2033475
- 134. Patel V, Saxena S, Lund C, Thornicroft G, Baingana F, Bolton P, Chisholm D, Collins PY, Cooper JL, Eaton J, et al. The Lancet Commission on global mental health and sustainable development. *Lancet*. 2018;392:1553–1598. doi: 10.1016/S0140-6736(18)31612-X