Depression in children and adolescents


Definition
Compared with adult depression, depression in children (6–12 years) and adolescents (13–18 years) may have a more insidious onset, may be characterised more by irritability than sadness, and occurs more often in association with other conditions such as anxiety, conduct disorder, hyperkinesis, and learning problems.

Incidence/prevalence
Estimates of prevalence of depression among children and adolescents in the community range from 2–6%.[4,5] Prevalence tends to increase with age, with a sharp rise around onset of puberty. Pre-adolescent boys and girls are affected equally by the condition, but depression is seen more frequently among adolescent girls than boys.[6]

Aetiology/risk factors
The aetiology is uncertain, but may include genetic vulnerability, childhood events and current psychosocial adversity.[7]

Prognosis
In children and adolescents, the recurrence rate following a first depressive episode in childhood or adolescence is 70% by 5 years, which is similar to the recurrence rate in adults. It is not clear if this is related to severity of depression.[8] Young people experiencing a moderate to severe depressive episode may be more likely than adults to have a manic episode within the next few years.[9,10] Trials of treatment for child and adolescent depression have found high rates of spontaneous remission (as much as two thirds of people in some inpatient studies).

What are the effects of treatments?

BENEFICIAL

Cognitive therapy (in mild to moderate depression)
One systematic review in children and adolescents with mild to moderate depression has found that cognitive behavioural therapy significantly improves symptoms compared with non-specific support.

Interpersonal therapy in adolescents (in mild to moderate depression)
Two RCTs found that interpersonal therapy versus clinical monitoring or waiting list control significantly increased recovery rate over 12 weeks in adolescents with mild to moderate depression.

TRADE OFF BETWEEN BENEFITS AND HARMs

Selective serotonin reuptake inhibitors
One RCT found no significant difference, one RCT found equivocal results (statistically significant differences on some depression measures but not others), and one RCT found a significant improvement in depressive symptoms with fluoxetine versus placebo after 8–9 weeks. One RCT found that, in adolescents with major depression, paroxetine versus placebo significantly improved remission after 8 weeks. We found no

RCTs on other selective serotonin reuptake inhibitors. Selective serotonin reuptake inhibitors are frequently associated with dizziness, lightheadedness, drowsiness, poor concentration, nausea, headache, and fatigue if treatment is reduced or stopped.

UNKNOWN EFFECTIVENESS

Cognitive therapy (depressed adolescents with depressed parent)
One RCT in depressed adolescents with depressed parents found no significant difference in recovery from depression with cognitive behavioural therapy plus usual care versus usual care alone over 2 years.

Electroconvulsive therapy
We found no RCTs on electroconvulsive therapy in children and adolescents with depression.

Monoamine oxidase inhibitors
One RCT found insufficient evidence to compare moclobemide versus placebo in improvement of depressive symptoms in children aged 9–15 years with major depression. We found no RCTs on non-reversible monoamine oxidase inhibitors in children or adolescents.

St John’s Wort
We found no RCTs on St John’s Wort (Hypericum perforatum) in children or adolescents with depression.

Venlafaxine
One RCT found no significant difference with venlafaxine versus placebo in global assessment or depression scores after 6 weeks in children with depression and family history of bipolar affective disorder. Lithium was associated with adverse effects.

Monoamine oxidase inhibitors
One RCT found no significant difference with lithium versus placebo in improvement of depressive symptoms in children aged 9–15 years with major depression. We found no RCTs on non-reversible monoamine oxidase inhibitors in children or adolescents.

St John’s Wort
We found no RCTs on St John’s Wort (Hypericum perforatum) in children or adolescents with depression.

Electroconvulsive therapy
We found no RCTs on electroconvulsive therapy in children and adolescents with depression.

Long term effects of treatments
We found no systematic review or RCTs examining long term outcomes of interventions for depression in children and adolescents.

Mood stabilisers
One RCT found no significant difference with lithium versus placebo in improvement of depressive symptoms in children with depression and family history of bipolar affective disorder. Lithium was associated with adverse effects.

Long term effects of treatments
We found no systematic review or RCTs examining long term outcomes of interventions for depression in children and adolescents.

Monoamine oxidase inhibitors
One RCT found insufficient evidence to compare moclobemide versus placebo in improvement of depressive symptoms in children and adolescents with major depression after 6 weeks.

Family therapy; group treatments other than cognitive behavioural therapy
We found insufficient evidence in children and adolescents about the effects of these interventions.

UNLIKELY TO BE BENEFICIAL

Oral tricyclic antidepressants (adolescents)
One systematic review found no significant difference with oral tricyclic antidepressants (amitriptyline, desipramine, imipramine, nortriptyline) versus placebo in depression scores in adolescents and children with depression. Subgroup analyses...
found that oral tricyclic antidepressants versus placebo significantly reduced symptoms in adolescents but not in children. The review also found that oral tricyclic antidepressants were associated with adverse effects.

LIKELY TO BE INEFFECTIVE OR HARMFUL

Oral tricyclic antidepressants (children)

Subgroup analyses in one systematic review found no significant difference with oral tricyclic antidepressants (amitriptyline, desipramine, imipramine, nortriptyline) versus placebo in children with depression. The review also found that oral tricyclic antidepressants were associated with adverse effects.

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