

# ACIP Evidence to Recommendations Framework

**Question:** Should hepatitis A catch-up vaccination be recommended for children age 2–18 years?

**Population:** Children aged 2–18 years

**Intervention:** Hepatitis A vaccination (2-dose schedule)

**Comparison(s):** Hepatitis A catch-up vaccination based on individual clinical decision-making

**Outcome:**

- Hepatitis A infection

**Background:** Despite successful implementation of the United States' routine hepatitis A vaccination recommendation for children aged 12–23 months, there remains a small gap to close among adolescents who have not received hepatitis A vaccine and are thus missing long-term and potential lifetime protection from hepatitis A.

Decreased hepatitis A incidence in the United States and reduced exposure to hepatitis A virus (HAV) have resulted in decreased anti-HAV seroprevalence among adults, and an increased proportion of susceptible adults, including younger adults (i.e., aged 20–29 years). Vaccinating adolescents will lead to increased protection among adults—particularly young adults—more quickly than waiting for the routinely vaccinated cohort of children to reach adulthood. It also will protect those with undisclosed risk factors (e.g., drug use, men who have sex with men [MSM]) more consistently and earlier in life.

In 2006, the Advisory Committee on Immunization Practices (ACIP) recommended routine hepatitis A vaccination for all children aged 12–23 months. Children who are not vaccinated by age 2 years can be vaccinated at subsequent health care visits, and catch-up vaccination of unvaccinated children aged 2–18 years can be considered based on shared clinical decision-making. Catch-up vaccination ensures that the percentage of children/adolescents who miss vaccination as scheduled or who were born outside of the routinely vaccinated cohort (i.e., born prior to 2006) are protected and is a way to increase herd immunity.

In 2017, national hepatitis A vaccination coverage among adolescents aged 13–17 years was 77.2% for 1 dose and 68.4% for  $\geq 2$  doses, compared to 36.2% and 25.3% for 1 and  $\geq 2$  doses, respectively, in 2008. This indicates substantial catch-up implementation and acceptance despite a recommendation based on clinical decision-making (1, 2).

The vaccines containing HAV antigen that are currently licensed in the United States for children are the inactivated, single-antigen vaccines HAVRIX® (manufactured by GlaxoSmithKline, Rixensart, Belgium) and VAQTA® (manufactured by Merck & Co., Inc., Whitehouse Station, New Jersey).

*Additional background information supporting the ACIP recommendations on the use of hepatitis A vaccines can be found in the relevant publication of the recommendations referenced on the ACIP website. <https://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hepa.html>*



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			<p>Among adolescents aged 13–17 years living in states where routine vaccination at 12–23 months was first recommended in 2006, coverage is only slightly lower than the total cohort, at 71% for 1 dose and 61% for <math>\geq 2</math> doses.</p>	
<b>BENEFITS &amp; HARMS</b>	<p>How substantial are the desirable anticipated effects?</p>	<p><i>Minimal</i>   <i>Small</i>   <i>Moderate</i>   <i>Large</i>   <i>Don't know</i>   <i>Varies</i></p> <p><input type="checkbox"/>   <input type="checkbox"/>   <input type="checkbox"/>   <input checked="" type="checkbox"/>   <input type="checkbox"/>   <input type="checkbox"/></p>	<p>HAVRIX and VAQTA are highly immunogenic when administered to children and adolescents according to multiple schedules.</p> <ul style="list-style-type: none"> <li>97%–100% of persons aged 2–18 years had protective levels of antibody 1 month after receiving the first dose, and 100% had protective levels 1 month after the second dose, with high geometric mean concentrations (6–14).</li> </ul> <p>Due to demonstrated long-term protection from the vaccines, there is little concern that childhood vaccination will result in risk later in life due to waning immunity.</p> <ul style="list-style-type: none"> <li>Antibody to HAV (anti-HAV) has been shown to persist for at least 22 years in adults administered inactivated vaccine as children (aged 3–6 years) (15, 16).</li> <li>Mathematical modelling based on persons vaccinated as adults predicted that seropositive anti-HAV levels would persist in <math>\geq 95\%</math> of vaccinees at year 30 and <math>\geq 90\%</math> at year 40 (17).</li> <li>Vaccine-induced cellular immunity has been shown to promote HAV-specific</li> </ul>	<p>Catch-up vaccination ensures children and adolescents who miss vaccination as scheduled or who were born outside of the routinely vaccinated cohort are protected and increases herd immunity.</p> <p>In addition, since implementation of risk-based vaccination in adults has been poor, catch-up vaccination will more rapidly increase the proportion of adults with risk factors who are protected.</p>

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		cellular immunity similar to that induced by natural infection (18).	
CRITERIA	WORK GROUP JUDGMENTS	RESEARCH EVIDENCE	ADDITIONAL INFORMATION
How substantial are the undesirable anticipated effects?	<p><i>Minimal Small Moderate Large Don't know Vari</i></p> <p><input checked="" type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/></p>	<p>More than 20 years of safety monitoring have shown no safety concerns.</p> <p>No unusual or unexpected safety patterns were observed in the Vaccine Adverse Event Reporting System for any hepatitis A vaccines (1).</p> <p>Rates of adverse events following TWINRIX® vaccination were similar to those seen with separately administered hepatitis A and hepatitis B vaccines (19). (TWINRIX® is approved for use in persons 18 years of age or older.)</p>	
Do the desirable effects outweigh the undesirable effects?	<p><i>Favors Favors Favors Favors Unclear</i> <i>intervention comparison both neither</i></p> <p><input checked="" type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/></p>	<p>Hepatitis A vaccination affords long-term protection against HAV infection.</p> <ul style="list-style-type: none"> <li>• Disease severity increases as persons age.</li> <li>• More than 20 years of safety monitoring have shown no safety concerns.</li> <li>• Earlier hepatitis A vaccination among healthy children/adolescents provides protection against HAV infection before persons develop increased risk for HAV infection or HAV-associated complications.</li> </ul>	
What is the overall certainty of this evidence for the critical outcomes?	<p>Effectiveness of the intervention</p> <p><i>No included studies</i> <input checked="" type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/></p> <p>4 <i>Very low</i> 3 <i>Low</i> 2 <i>Moderate</i> 1 <i>High</i></p> <p>Safety of the intervention</p> <p><i>No included studies</i> <input checked="" type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/></p> <p>4 <i>Very low</i> 3 <i>Low</i> 2 <i>Moderate</i> 1 <i>High</i></p>	<p>GRADE was not used to evaluate the evidence.</p> <ul style="list-style-type: none"> <li>• Hepatitis A vaccine has been recommended for administration to children since 1996.</li> <li>• Hepatitis A vaccine has been recommended for catch-up vaccination based on clinical decision-making since 2006.</li> <li>• The efficacy and safety of Hepatitis A vaccines has been evaluated and well documented.</li> </ul>	

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			<ul style="list-style-type: none"> <li>There are no known safety concerns with hepatitis A vaccines.</li> </ul>	
VALUES	<p>Does the target population feel that the desirable effects are large relative to undesirable effects?</p>	<p>No    Probably no    Uncertain    Probably yes    Yes    Varies</p> <p><input type="checkbox"/>    <input type="checkbox"/>    <input type="checkbox"/>    <input checked="" type="checkbox"/>    <input type="checkbox"/>    <input type="checkbox"/></p>	<p>In 2017, national hepatitis A vaccination coverage among adolescents aged 13–17 years was 77.2% for 1 dose and 68.4% for ≥2 doses compared to 36.2% and 25.3% for 1 and ≥2 doses, respectively, in 2008 (1, 2).</p> <ul style="list-style-type: none"> <li>This indicates substantial catch-up vaccination implementation and acceptance despite a recommendation based on clinical decision-making.</li> <li>Specific studies of demand among older children for hepatitis A vaccine have not been done. However, almost 70% of teens have already initiated the vaccine series even though catch-up vaccination is not routinely recommended, suggesting parental acceptance or demand for hepatitis A vaccination.</li> </ul>	<p>Compared to other vaccines with a routine catch-up schedule, hepatitis A adolescent 2-dose coverage is comparable (e.g., quadrivalent meningococcal conjugate vaccine) or greater (e.g., human papillomavirus vaccine), providing evidence of acceptability by the target population (1).</p>
	<b>CRITERIA</b>	<b>WORK GROUP JUDGMENTS</b>	<b>RESEARCH EVIDENCE</b>	<b>ADDITIONAL INFORMATION</b>
	<p>Is there important uncertainty about or variability in how much people value the main outcomes?</p>	<p><i>Important uncertainty or variability</i>    <input type="checkbox"/></p> <p><i>Possibly important uncertainty or variability</i>    <input type="checkbox"/></p> <p><i>Probably no important uncertainty or variability</i>    <input checked="" type="checkbox"/></p> <p><i>No important uncertainty or variability</i>    <input type="checkbox"/></p> <p><i>No known undesirable outcomes</i>    <input type="checkbox"/></p>	<p>The high coverage rate, despite this recommendation being based on clinical decision-making, provides strong and consistent evidence that most parents believe that it is important.</p>	

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ACCEPTABILITY	<p>Is the intervention acceptable to key stakeholders?</p>	<p>No <input type="checkbox"/>    Probably no <input type="checkbox"/>    Uncertain <input type="checkbox"/>    Probably yes <input checked="" type="checkbox"/>    Yes <input type="checkbox"/>    .....    Varies <input type="checkbox"/></p>	<p>CDC currently does not have a routine catch-up recommendation for children aged 2–18 years, yet 21 states have introduced mandates for daycare, daycare plus school, or school alone. This represents an increase in states with mandates from 28% in 2011 to 40% in 2018 (20).</p> <ul style="list-style-type: none"> <li>As of October 2018, 12 states have a daycare and school mandate; 8 states plus part of Arizona have a daycare-only mandate; 1 state has a school-only mandate (Indiana).</li> </ul>	<p>Children who were 1 year of age when the routine recommendation was first published in 2006 were ~12–13 years old in 2017 when these data were collected; nearly all of the cohort of children 13–17 assessed in 2017 were not subject to the routine recommendation for children 12–23 months of age. High overall coverage in this age group demonstrates that catch-up vaccination is occurring.</p> <p>As noted above, the hepatitis A adolescent 2-dose coverage is comparable or greater than other vaccines with a routine catch-up schedule, which provides evidence of acceptability by key stakeholder populations.</p>
RESOURCE USE	<p>Is the intervention a reasonable and efficient allocation of resources?</p>	<p>No <input type="checkbox"/>    Probably no <input type="checkbox"/>    Uncertain <input type="checkbox"/>    Probably yes <input checked="" type="checkbox"/>    Yes <input type="checkbox"/></p>	<p><b>Cost analysis:</b> After universal childhood recommendation implementation, a cost-effectiveness model used to assess nationwide routine hepatitis A vaccination was adapted to assess the cost-effectiveness of catch-up hepatitis A vaccination among unvaccinated and partially vaccinated children compared with unvaccinated children (21).</p> <ul style="list-style-type: none"> <li>Over the cohort’s lifetime, catch-up vaccination would reduce the total number of infections relative to baseline by 741.</li> <li>Catch-up vaccination would increase net cost by \$2.38 per person.</li> </ul>	<p><b>Other studies:</b> To assess the population-level impact and cost-effectiveness of US hepatitis vaccination programs, an age-structured population model of hepatitis A transmission dynamics was developed to evaluate two policies of administering a 2-dose hepatitis A vaccine to children aged 12 to 18 months (22).</p> <ul style="list-style-type: none"> <li>The model predicted universal childhood</li> </ul>

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			<ul style="list-style-type: none"> <li>• Incremental cost of hepatitis A vaccine catch-up intervention at age 10 years, the midpoint of the ages modeled, was \$452,239 per QALY gained.</li> <li>• Across age cohorts, cost effectiveness of catch-up vaccination is most favorable at age 12 years, resulting in an incremental cost-effectiveness ratio (ICER) of \$189,000 per QALY gained.             <ul style="list-style-type: none"> <li>○ The impact of vaccination on the ICER was most sensitive to the discount rate, followed by the rate of adult vaccination.</li> </ul> </li> <li>• The model assumed administration costs of hepatitis A vaccination were split with other vaccines routinely administered at age 12 years, thus lowering the cost of vaccination.</li> <li>• Catch-up vaccination in adolescence is more effective when it is assumed to replace later vaccination as an adult.             <ul style="list-style-type: none"> <li>○ Catch-up became more cost effective when targeting children in late adolescence, due to higher probability of symptomatic disease among older children, less discounting of future costs of disease, and less delay in averting adult vaccination costs.</li> </ul> </li> <li>• Limitations: model output is based on hepatitis A incidence from 2008 to 2012 and the cost-effectiveness conclusions are strongly tied to factors such as vaccine uptake and disease transmission patterns which may change over time, altering future cost.</li> <li>• The model also excluded herd immunity effects from vaccination.</li> </ul> <p>Incremental costs of catch up now, given current rates of coverage among 13–17-year-</p>	<p>hepatitis A vaccination would lead to significant reduction in hepatitis A mortality and morbidity.</p> <ul style="list-style-type: none"> <li>• Universal vaccination was cost saving compared with a regional vaccination policy.</li> <li>• The model predicts US incidence will fall to 27/100,000 by 2020, compared to a CDC-reported incidence (adjusted for underreporting) of just 1.71/100,000.</li> <li>• This model overestimates incidence which limits applicability of inferences from these findings to catch-up policies.</li> <li>• While this model provides a favorable cost-effectiveness estimate, limitations exist.</li> </ul> <p><b>Additional Considerations:</b> Cost of outbreak response and hospitalization is substantial and would offset some of additional costs of catch-up vaccination.</p> <p>Based on 2017 hepatitis A vaccination coverage rates:</p>
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			<p>olds, would be more favorable, because hepatitis A vaccination coverage rates are higher among vaccinated children who are aging into the adolescent cohort:</p> <ul style="list-style-type: none"> <li>• 2008, 1 dose: 36.2%, ≥2-dose: 25.3%, versus 2017, 1 dose: 77.2%, 2-dose 68.5%. (1,2).</li> </ul> <p>Achieving 80%–90% coverage among teens would require a much smaller number of additional vaccines given. In addition, HAV incidence overall is higher due to the ongoing multistate outbreaks (0.5 cases/100,000 population in 2012 versus 1.0 case/100,000 population in 2017) (1, 22).</p>	<ul style="list-style-type: none"> <li>• To achieve 80% hepatitis A vaccine series completion among adolescents, 2,205,491 additional persons would need to be vaccinated with one dose and 701,747 persons would need to be vaccinated with 2 doses, at a cost of \$59.3M (private) or \$37M (CDC).</li> <li>• To achieve 90% hepatitis A vaccine series initiation among adolescents, 3,207,987 persons would need to be vaccinated with 1 dose, at a cost of \$52.8M (private), \$32.9M (CDC).</li> </ul> <p>Assumptions:            1) 2017 coverage rates apply to all children; however, in actuality, younger adolescents have higher coverage.            2) 100% of 1-dose recipients complete series when calculating cost of 80% completion.            3) Private sector covers 50% of these adolescents and CDC covers 50%.</p>
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FEASIBILITY	<p>Is the intervention feasible to implement?</p>	<p> <i>No</i>    <i>Probably no</i>    <i>Uncertain</i>    <i>Probably yes</i>    <i>Yes</i>    <i>Varies</i>  <input type="checkbox"/>    <input type="checkbox"/>    <input type="checkbox"/>    <input type="checkbox"/>    <input checked="" type="checkbox"/>    <input type="checkbox"/> </p>	<p>Of the 30 registries American Immunization Registry Association (AIRA) were able to query to test forecasting algorithms, 27 already routinely forecast hepatitis A vaccine for an 18-year-old who has never been vaccinated. All 30 algorithms forecast the second dose in any 18-year-old who has had one dose. Essentially, these algorithms are implemented as routine catch-up. Therefore, 27 of 30 registries would not have to change to implement routine catch-up.</p> <p>Findings from a 2014 survey (23) indicate:</p> <ul style="list-style-type: none"> <li>• If ACIP made a recommendation for catch-up hepatitis A vaccination at health maintenance visits for all children 2 to 18 years of age, 96% of pediatricians and 79% of family physicians reported it would be very feasible to routinely assess hepatitis A vaccination status and vaccinate children and adolescents who were not fully vaccinated.             <ul style="list-style-type: none"> <li>○ An additional 4% and 19%, respectively, indicated it would be moderately feasible.</li> </ul> </li> <li>• The most common barriers to implementing a hepatitis A vaccine catchup recommendation included infrequent visits by adolescent patients (65%); parents' lack of knowledge on the seriousness of HAV disease (39%); hepatitis A vaccination not being required for childcare or school entry (39%); difficulty obtaining immunization records to determine a patient's hepatitis A vaccination status (35%); and parental concerns about giving too many vaccines at 1 visit (33%).</li> </ul>	<p>Routine hepatitis A catch-up already exists in states that have school mandates (21 states have introduced mandates for daycare, daycare plus school, or school alone.). In New York City, all children and adolescents not previously vaccinated should receive the two-dose hepatitis A vaccine series by their 19th birthday for long-term protection (24).</p> <p>There are opportunities to administer hepatitis A vaccine to adolescents concurrently with vaccines protecting against other infections, such as human papillomavirus and meningococcal disease.</p>

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			<ul style="list-style-type: none"> <li>This study was performed 6 years ago and there is currently more education and awareness among providers and the public due to ongoing outbreaks, likely decreasing barriers to vaccination.</li> </ul>			
Balance of consequences	Undesirable consequences <i>clearly outweigh</i> desirable consequences in most settings <input type="checkbox"/>	Undesirable consequences <i>probably outweigh</i> desirable consequences in most settings <input type="checkbox"/>	The balance between desirable and undesirable consequences <i>is closely balanced or uncertain</i> <input type="checkbox"/>	Desirable consequences <i>probably outweigh</i> undesirable consequences in most settings <input checked="" type="checkbox"/>	Desirable consequences <i>clearly outweigh</i> undesirable consequences in most settings <input type="checkbox"/>	There is insufficient evidence to determine the balance of consequences <input type="checkbox"/>
Is there sufficient information to move forward with a recommendation? Yes <input checked="" type="checkbox"/> No <input type="checkbox"/>						
Policy Options for ACIP Consideration	ACIP does not recommend the intervention <input type="checkbox"/>	ACIP recommends the intervention for individuals based on clinical decision-making <input type="checkbox"/>			ACIP recommends the intervention <input checked="" type="checkbox"/>	
Recommendation (text)	Recommended for all unvaccinated children and adolescents aged 2–18 years. Children and adolescents who have not previously received hepatitis A vaccine should be vaccinated routinely at any age (i.e., children and adolescents are recommended for catch-up vaccination).					
Additional considerations (optional)						

**Final deliberation and decision by the ACIP**

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Final ACIP recommendation	ACIP does not recommend the intervention <input type="checkbox"/>	ACIP recommends the intervention for individuals based on shared clinical decision-making <input type="checkbox"/>	ACIP recommends the intervention <input checked="" type="checkbox"/>
ACIP considerations			

i This Evidence to Recommendation table is based on the GRADE Evidence to Decision framework developed through the *DECIDE* project. Further information is available at <http://www.decide-collaboration.eu/evidence-decision-etc-framework>

### References:

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