

REPORT FROM CLINICAL PANEL

TITLE **D09X04**
Auditory brainstem implant in patients with congenital abnormalities of the auditory nerves or cochleae

CRG: Specialised Ear Services

NPOC: Trauma

Lead: Fiona People

Date: 18 November 2015

Question	Conclusion of the panel	If there is a difference between the evidence review and the policy please give a commentary
<p><u>The population</u></p> <p>1. What are the eligible and ineligible populations defined in the policy and are these consistent with populations for which evidence of effectiveness is presented in the evidence review?</p>	<p>The eligible population(s) defined in the policy are the same or similar to the population(s) for which there is evidence of effectiveness considered in the evidence review</p>	
<p><u>Population subgroups</u></p> <p>2. Are any population subgroups defined in the policy and if so do they match the subgroups for which there is evidence presented in the evidence review?</p>	<p>The population subgroups defined in the policy are the same or similar as those for which there is evidence in the evidence review</p>	

<p><u>Outcomes - benefits</u></p> <p>3. Are the clinical benefits demonstrated in the evidence review consistent with the eligible population and/or subgroups presented in the policy?</p>	<p>The clinical benefits demonstrated in the evidence review support the eligible population and/or subgroups presented in the policy</p>	
<p><u>Outcomes – harms</u></p> <p>4. Are the clinical harms demonstrated in the evidence review reflected in the eligible population and/or subgroups presented in the policy?</p>	<p>The clinical harms demonstrated in the evidence review are reflected in the eligible population and/or subgroups presented in the policy</p>	<p>The procedure has a risk profile that seems similar to other skull base procedures such as microvascular decompression surgery. This includes a 1-2% risk of intracranial bleeding, stroke or meningitis, a 1-2% risk to surrounding cranial nerves (particularly the facial, glossopharyngeal and vagal) and a 10-15% risk of CSF leak of whom half may require reoperation).</p>
<p><u>The intervention</u></p> <p>5. Is the intervention described in the policy the same or similar as the intervention for which evidence is presented in the evidence review?</p>	<p>The intervention described in the policy the same or similar as in the evidence review</p>	
<p><u>The comparator</u></p> <p>6. Is the comparator in the policy the same as that in the evidence review?</p> <p>7. Are the comparators in the evidence review the most plausible comparators</p>	<p>The comparator in the policy is the same as that in the evidence review</p> <p>The comparators in the evidence review include</p>	<p>Profound Deafness</p>

<p>for patients in the English NHS and are they suitable for informing policy development.</p>	<p>plausible comparators for patients in the English NHS and are suitable for informing policy development.</p>	
<p><u>Advice</u></p> <p>The Panel should provide advice on matters relating to the evidence base and policy development and prioritisation. Advice may cover:</p> <ul style="list-style-type: none"> • Uncertainty in the evidence base • Challenges in the clinical interpretation and applicability of policy in clinical practice • Challenges in ensuring policy is applied appropriately • Issues with regard to value for money • Likely changes in the pathway of care and therapeutic advances that may result in the need for policy review. 		<p>No further advice (all advice from the July Clinical Panel noted / reviewed).</p>

Overall conclusions of the panel

The policy reflects the findings of the clinical evidence review and should progress.

Report approved by:

James Palmer
 Chair Clinical Panel

FOR PUBLIC CONSULTATION ONLY