

**SPECIALISED COMMISSIONING - CLINICAL EVIDENCE EVALUATION
CRITERIA FOR A PROPOSITION FOR A CLINICAL COMMISSIONING POLICY
FOR NON-ROUTINE COMMISSIONING**

URN: B01X09

TITLE: Proton beam therapy for prostate cancer

CRG: Radiotherapy

NPOC: Cancer

Lead: Nicola McCulloch

Date: 17th February 2016

The panel were presented a policy proposal for non-routine commissioning

Question	Conclusion of the panel	<i>If there is a difference between the evidence review and the policy please give a commentary</i>
<p><u>The population</u></p> <p>1. Are the eligible and ineligible populations defined in the policy consistent with the evidence of effectiveness, and evidence of lack of effectiveness; and where evidence is not available for the populations considered in the evidence review?</p>	<p>The ineligible population(s) defined in the policy are the same or similar to the population(s) for which there is evidence of lack of effectiveness or inadequate evidence of effectiveness demonstrated 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 considered by the evidence review?</p>	<p>The population subgroups defined in the policy are the same or similar as those considered by the evidence review.</p>	
<p><u>Outcomes - benefits</u></p> <p>3. Are the clinical benefits demonstrated in the evidence review</p>	<p>The lack of benefit or absence of evidence of benefit demonstrated in</p>	

<p>consistent with the eligible population and/or subgroups presented in the policy?</p>	<p>the evidence review is consistent with the ineligible 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 and / or ineligible population and/or subgroups presented in the policy?</p>	<p>The clinical harms demonstrated in the evidence review are reflected in the eligible and / or ineligible population and/or subgroups presented in the policy.</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 is 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 for patients in the English NHS and are they suitable for informing policy development.</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 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</p>		<p>The Panel agreed the policy proposition. There was a question about a more recent paper but</p>

<p>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>reading of the abstract of this during the meeting did not seem to indicate that the outcome of the paper would change the policy proposition. The paper would be reviewed through the Public Health Network. The view of the Panel was that stakeholder engagement should continue but that there should be a specific question asked around whether or not the evidence review had captured all the relevant and eligible papers.</p>
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Overall conclusions of the panel

The policy should proceed as a non-routine commissioning policy.

Report approved by:
James Palmer
Clinical panel Chair
17/2/16

Post meeting note:

A specific question relating to whether or not the evidence review had captured all the relevant and eligible papers was included for consultation.