SPECIALISED COMMISSIONING - CLINICAL EVIDENCE EVALUATION CRITERIA FOR CLINICAL COMMISSIONING POLICY PROPOSITION

URN: 1842

TITLE: Proton Beam Therapy for Hepatocellular Carcinoma

CRG: Radiotherapy NPOC: Cancer Date: 16/01/19

	T		
This policy is being	For routine	Not for routine	X
considered for:	commissioning	commissioning	
Is the population	Yes, patients with inoperable hepatocellular carcinoma for		
described in the policy	whom radiotherapy is a treatment option.		
similar to that in the			
evidence reviewed,			
including subgroups?			
Is the intervention	Yes.		
described in the policy			
similar to the			
intervention for which			
evidence is presented in			
the evidence review?			
Are the comparators in	The studies are weak	The comparators are conventi	ional
the evidence reviewed	radiotherapy and stereotactic ablative radiotherapy (SABR).		
plausible clinical	1 7	17 (,
alternatives within the			
NHS and are they			
suitable for informing			
policy development?			
Are the clinical benefits	The clinical benefits of	this treatment are not well	
described in the		consistent with the policy state	ement
		utine commissioning position.	JIIIOIIL
evidence review likely to	procenting a net for re	aurie cerminosiermig pecinem	
apply to the eligible			
population and/or			
subgroups in the policy?	Daniel a diversidados de	hat than an an actual all also at an	.1.1
Are the clinical harms	_	hat there are potential short an	-
described in the		ng the exposure of sensitive no	
evidence review likely to		rgeted forms of radiotherapy th	
apply to the eligible and		surrounding local normal tissu ical advantages. Proton bean	
or ineligible population		potential to result in less radiat	
and/or subgroups in the		ng tissue and Stereotactic Abla	
policy?	<u> </u>	is another form of radiotherapy	
		n to the tumour. In patients wit	
		ects are less likely to be clinical	
		e not expected to be cured as	•
	•	ave a shortened life expectant	
		nediate damage to surrounding	-
		nt but the degree to which this	
		its place in the pathway relative	
		ABR itself is not routinely	
	commissioned and is	subject to an evaluation progra	mme.

The Panel should provide advice on matters relating to the evidence base and policy development and prioritisation. Advice may cover: Balance between benefits and harms Quality and uncertainty in the evidence base Challenges in the clinical interpretation and applicability of policy in clinical practice Challenges in ensuring policy is applied appropriately Likely changes in the pathway of care and therapeutic advances that may result in the need for policy review.	the PPP stated 'toxicity data studies, and as a result it is recompare acute and late treat data'. There is some suggestion the complications form PBT may radiotherapy modalities. How are inadequate to determine certainty. The Panel supported the not policy statement but noted the research and evaluation in dethis treatment in the pathway. Panel recommended that the for public consultation outside policy statement development.	at adverse events and tend to be lower than for other wever, the quality of the studies this with any degree of for routine commissioning are may be a place for further etermining the potential place of the standard process for ant.
Overall conclusion	This is a proposition for routine commissioning and	Should proceed for routine commissioning Should be reversed and proceed as not for routine commissioning Should X
		proceed for

This is a proposition for not routine	not routine commissioning
commissioning and	Should be reconsidered
	by the PWG

Overall conclusions of the panel Report approved by:

Report approved by: David Black Clinical Panel Chair 25/1/19

Post meeting note:

On review of stakeholder feedback, which was supportive of the policy statement, the Programme of Care Board agreed that public consultation was not required.