SPECIALISED COMMISSIONING - CLINICAL EVIDENCE EVALUATION CRITERIA FOR A PROPOSITION FOR A CLINICAL COMMISSIONING POLICY NOT FOR ROUTINE COMMISSIONING

URN: 1779

TITLE: Hyperbaric oxygen therapy for diabetic limb ulceration (diabetic foot ulcer)

NPOC: Trauma

Lead: TBC / Jacquie Kemp

Date: 20 June 2018

This policy is being	For routine		Not for routine	Х	
considered for:	commissioning		commissioning		
Is the population	Yes.				
described in the policy					
the same as that in the					
evidence review					
including subgroups?					
Is the intervention	Yes, 'dose' of hyperbaric oxygen treatment varies (no				
described in the policy	standardisation).				
the same or similar as					
the intervention for which					
evidence is presented in					
the evidence review?					
Is the comparator in the	No. The studies are variable and best available wound				
policy the same as that	care is the appropriate comparator but at best evidence				
in the evidence	of effectiveness in comparison is limited.				
review? 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?					
development:					
Are the clinical benefits	Not demonstrated.				
demonstrated in the					
evidence review					
consistent with the					
eligible population and/or					
subgroups presented in					
the policy?					
Are the clinical harms	Recognised hazards				
demonstrated in the					
evidence review					
reflected in the eligible					
and /or ineligible					
population and/or					

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Supported the not for routine commissioning position based on the lack of evidence of efficacy.
The PWG has expressed concerns about how these studies have been compared and conclusions reached about the robustness and applicability of the studies. The PWG have stated 'In our opinion, the decision to consider the trial by Londahl at al less reliable than Fedorko et al is unfounded and the endorsement of a speculative mechanism for the non-existent difference in outcomes misleads further'.
Clinical Panel recognised the careful consideration and hard work of the PWG to date. Panel also recognised the process that is used to develop the evidence reviews to inform policy and also understood that process needs to be followed in order that there is consistency in policy production. Then PWG concerns have been considered by the evidence reviewers and some amendments have been made. Panel understands that these do not address all the concerns of the PWG.
Panel determined that the policy should progress to the stakeholder and consultation and that the consultation process has the potential to generate comments on the policy and evidence base that informs it. If received, these will be considered through the policy development process. Panel noted that the PWG were not advising that the differences between the PWG interpretation of the evidence and that contained in the independent evidence review were sufficient to result in a change in the proposed policy position.
Proceed as NRC.

 ensuring policy is applied appropriately Likely changes in the pathway of care and therapeutic advances that may result in the need for policy review. 			
Overall conclusion	This is a proposition for routine commissioning and	Should proceed for routine commissioning Should reversed and proceed as not for routine commissioning	
	This is a proposition for not routine commissioning and	Should proceed for not routine commissioning Should be reconsidered by the PWG	X

Report approved by: Dr David Black Deputy Medical Director, Specialised Services 13 July 2018