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## Medicines Management Dept

Patient Group Direction for the Supply/administration of ....

## Patient Group Direction for the Supply/administration of Hepatitis B vaccination in those adults considered to be at high risk (excluding those presenting with a needlestick injury) by qualified and suitably trained Community Pharmacists

Define situation/condition	Immunisation against Hepatitis B		
	UK National guideline on the management of viral hepatitides A, B and C, 2008 http://www.bashh.org/guidelines/2005/hepatitis_abc_final_0905.pdf		
Criteria for inclusion	<ul> <li>Homosexual males</li> <li>Bisexual males</li> <li>Female sexual partners of bisexual males</li> <li>Intravenous drug users</li> <li>Alcohol Detox patients</li> <li>Sexual partners of intravenous drug users</li> <li>Commercial sex workers</li> <li>Regular users of commercial sex workers</li> <li>Those travelling abroad frequently to endemic regions</li> </ul>		
Criteria for exclusion	All patients outside the target group Under 18 years of age Acute severe febrile illness Known hepatitis B positive individuals Known HIV positive patients or immunosuppressed individuals. A confirmed anaphylactic reaction to the vaccine or any component of the vaccine Pregnant Women Post exposure Prophylaxis No valid consent.		
Action if excluded	Refer to GP or Sexual Health Service		
Contraindications	Engerix B should not be administered to subjects with known hypersensitivity to any component of the vaccine, or to subjects having shown signs of hypersensitivity after previous Engerix B		

	administration. As with other vaccines, the administration of Engerix B should be postponed in subjects suffering from acute severe febrile illness. The presence of a minor infection,
	however, is not a contra-indication for immunisation. Pregnancy and Breastfeeding mothers
Action if contraindications	Document and Refer to GP or Sexual health service
Action if patient declines	Document and Refer to GP or Sexual Health Services

<ul> <li>Practising Pharmacists registered with the RPSGB who have completed the PCT approved training to deliver this service</li> <li>Pharmacy actively engaged with Needle Exchange and supervised consumption of methadone services. Completion of two day vaccination training programme provided by M and K updates; Basic life support and anaphylaxis-provided by trust and full understanding of PGD.</li> </ul>		
Annual update in Blood Borne Virus Training.		
Annual update in BLS and anaphylaxis.		
The pharmacist should be aware of any change to the recommendations for the medicine listed.		
Continued professional development is the responsibility of the pharmacist. He/ She should keep up to date with developments in areas relevant to this PGD.		

3. Description of Treatment			
Name of Medicine	Hepatitis B recombinant vaccine adsorbed (Engerix B prefilled syringes)		
Legal status of medicine	РОМ		
Dose	Adults over 18 years- ultra rapid course: One dose(1ml prefilled syringe-20mcg).		
Route	IM into right or left deltoid muscle		
Frequency	One dose (1ml prefilled syringe-20mcg) given at 0,7, and 21 days. A booster dose is recommended at 12 months to provide long term protection.		

Drug	Contraindications/ Cautions	Common Adverse Effects	Interactio ns	Notes
Hepatitis B recombinant vaccine adsorbed (Engerix B prefilled syringe 20mcg/1ml suspension)	<ul> <li>Pregnancy</li> <li>Breastfeeding</li> <li>Known allergy and/or</li> <li>hypersensitivity to Engerix B.</li> <li>Thrombocytopenia.</li> <li>Bleeding disorders</li> <li>Febrile Illness</li> </ul>	Very rare – see SPC • Injection site pain • Injection site Erythema • Injection Site Induration	Non detailed in SPC	See SPC Must not be administered into the buttock or intradermally at any site.

Follow up treatment	
Written/verbal advice for patient	<ul> <li>Check client's ID.</li> <li>Explain common side effects of vaccination.</li> <li>Obtain verbal consent to give vaccination.</li> <li>Give appointment for next dose prior to client leaving the service. Agree sms contact if appropriate</li> <li>Emphasis the importance of completing the vaccine course.</li> <li>Offer safer sex advice and condoms.</li> </ul>
Specify method of recording supply and /or administration	<ul> <li>Completion of IOWPCT treatment form, including additional record entry in patient PMR.</li> <li>The pharmacist must keep a record of the consultation for at least two years. The following should be noted in the pharmacist's records: <ul> <li>Assessment of client need in relation to the intervention.</li> <li>Date and time of supply and administration.</li> <li>Dose given</li> <li>Record of dose number as per schedule</li> <li>Batch number and expiry date.</li> <li>Advice given and leaflets supplied.</li> <li>Signature of client.</li> </ul> </li> </ul>
Procedure for reporting ADRs to Medical Practitioner	Whilst rare, all serious ADRs should be reported, even if the effectis well recognised. (See British National Formulary (BNF) forsupporting information.)ADRs should be reported to:The patient's GPThe Committee on Safety of Medicines, using the Yellow ADR cardsystem. Cards are available: in the BNF; from the MedicinesManagement Teams; and electronic versions atwww.yellowcard.gov.uk.

Management of Group Directions: Group direction developed by:	Mr Kevin Noble – Community Pharmacy Lead IW PCT Mr Paul Jerram – Head of Medicines Management IW PCT Dr John Partridge- Clinical Governance Lead	
Authorizing Doctor/s:	Signature.	
Date applicable:	Dr John Partridge Date signed off	
Review date:	2 years or as appropriate	
Senior Pharmacist	Signature.	
Clinical Directorate Pharmacist	Mr Kevin Noble- Community Pharmacy Lead Signature.	
	Mr Paul Jerram- Head of Medicines Management	
Approved by Pharmacy Group	Signature	
Assured by Oliviaal Otenderde Oregun	Mr Paul Jerram- Head of Medicines Management	
Approved by Clinical Standards Group	Signature	
	Signed by chair of committee (making the Trust liable for the supply and administration of medicines under the PGD, subject to its proper application by authorised and competent personnel.	

The group direction is to be read, agreed to, and signed by all staff it applies to. One copy is to be given to the health professional, another kept in the department.

## I have read the group direction and agreed to use it in accordance with the criteria described.

## All professionals who will be using the PGD need to read it and sign. Their review date should ideally be linked to appraisals or other personal review processes to ensure that they are still competent to be approved practitioners under the PGD