Facility: COM HKH MQE MVH RNS RYD

REMOTE HEPATOLOGY CONSULT:

	This PDF will	expire on 1	1 August 2024
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FAMILY NAME			MRN		
GIVEN NAME			MALE	FEMALE	
D.O.B. DD / MM / YYYY M.O.					
ADDRESS					
			PH		
M/C		FIN			
LOCATION (MADD			ADM DD (/M + VVVV	

HCV TRE	VTN/ENIT		LOCAT	TION / WARD ADM DD / MM / YYYYY			ſΥ
		COMPLETE ALL DETAILS OR AFFIX PATIENT LABEL HERE					
Note: This form is no	ot a referral for a	patient appointment					
For the Attention	of the Hepatol	ogist					
	gastroenterologis	nedical practitioner e st, hepatologist or info					
Patient First Name				Patient Surname .			<u>-</u>
Date of Birth: $__/$	/			Gender:	M F		
Address						Postcode	
Hepatitis C History Date of HCV Diagno				Non-pregnant (fen Discussion re cont		Yes h	No
Known Cirrhosis*?		Yes	No	male & female pat	ients?	Yes	No
Prior Antiviral Trea	tment			Current Medication			
Has patient previous antiviral treatment? Has prior treatment	,	Yes	No			riptions, herbal, over the cou ttach a summary page of	
Boceprevir/Telaprev		Yes	No				
I have checked for p	_	_					
interactions with cu	rrent medications	s.‡ Yes	No				
Intercurrent Condit	tions						
Diabetes		Yes	No				
Obesity		Yes	No				
Hepatitis B HIV		Yes	No				
Alcohol > 40 g/day		Yes Yes	No No				
	ania hanatia daga	ompensation or HBV/		nfaction aboutd be re	oforrad to a appa	iolist	
	•	nt and attach a PDF f			-		
Laboratory Resul	ts (or attached cor	by of results)			'		
Test	Date	Result		Test	Date	Result	
HCV genotype	/ /			Creatinine	/ /		
HCV RNA level	/ /			eGFR	/ /		
ALT	/ /			Haemoglobin	/ /		
AST	/ /			Platelet Count	/ /		
Bilirubin				INR			
Albumin	/_/_			Weight (kg)	//		
Liver Fibrosis Ass	sessment**						
Test	Date	Result					
FibroScan or	/ /						
Other (e.g. APRI)	/ /						
APRI, AST to platel	et ratio index: wv	 ww.hepatitisc.uw.edu	ı/page/d	clinical-calculators/a	apri		
						ıld be referred to a specia	alist.



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FAMILY NAME		MRN
GIVEN NAME		☐ MALE ☐ FEMALE
D.O.B. DD / MM / YYYY	M.O.	
ADDRESS		
		PH
M/C	FIN	
LOCATION / WARD		ADM DD / MM / YYYY

Treatment Choice (Select one prescribing choice)						
Regimen	Genotype	Status	Duration	Select		
Sofosbuvir plus Ledipasvir	1	Treatment naive – no cirrhosis HCV RNA < 6 x 10 ⁶ IU/mL	8 weeks			
	1	No cirrhosis	12 weeks			
Sofosbuvir plus Daclatasvir	1	No cirrhosis treatment – naive	12 weeks			
		No cirrhosis treatment – experienced	12 weeks or 24 weeks			
Sofosbuvir plus Daclatasvir	3	No cirrhosis	12 weeks			
Sofosbuvir plus Ribavirin	2	No cirrhosis	12 weeks			
Paritaprevir/Ritonavir plus Ombitasvir plus Dasabuvir	1b	No cirrhosis	12 weeks			
Paritaprevir/Ritonavir plus Ombitasvir plus Dasabuvir Ribavirin	1a	No cirrhosis	12 weeks			
Grazoprevir-Elbasvir	1 or 4	Naive - no cirrhosis	12 weeks			
	1b	Treatment experienced - no cirrhosis	12 weeks			
Grazoprevir-Elbasvir Ribavirin	1a or 4	Treatment experienced - no cirrhosis	16 weeks			
Sofosbubir-Velpatasvir	1, 2, 3, 4,	Treatment naive - no cirrhosis	12 weeks			
	5 or 6	Treatment experienced - no cirrhosis	12 weeks			

Virus Infection: A Consensus Statement (www.gesa.org.au). Information is also available at: www.pbs.gov.au/info/healthpro/explanatory-notes/general-statement-hep-c

Patients must be tested for HCV RNA at least 12 weeks after completing treatment to determine outcome. Please notify the specialist below of the Week 12 Post Treatment result.

Declaration by General Practitioner/Medical Officer				
I declare all of the information provided above is true and correct:				
GP/MO Name				
Address Postcode				
Phone	. Fax			
Mobile Phone	Email			
Signature	Date:/			
Once completed please return both pages and the drug interactions and other attachments by email:				

NSLHD-RNS-HepatologyService@health.nsw.gov.au

Approval by Specialist Experienced in the Treatment o	f HCV	(Office Use On	lν
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I agree with your decision to treat this person based on the information provided above

This patient would be more suited to HCV treatment under specialist supervision. Please forward a formal referral for this patient and an appointment will be scheduled.

Name	Signature
Designation	Dotos / /