RS1827 - Pegylated interferon alfa-2a

Chronic Hepatitis C - genotype 1 infection treatment more than 4 years prior - INITIATION Chronic hepatitis C - genotype 1 infection - CONTINUATION	
Chronic hepatitis C - genotype 1, 4, 5 or 6 infection or co-infection with HIV or genotype 2 or 3 post liver trans - INITIATION	
Chronic hepatitis C - genotype 2 or 3 infection without co-infection with HIV - INITIATION Hepatitis B - INITIATION	
Myeloproliferative disorder or cutaneous T cell lymphoma - INITIATION Myeloproliferative disorder or cutaneous T cell lymphoma - CONTINUATION	3
Ocular surface squamous neoplasia - INITIATION Ocular surface squamous neoplasia - CONTINUATION	4
Post-allogenic bone marrow transplant - INITIATION Post-allogenic bone marrow transplant - CONTINUATION	4
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HOSPITAL MEDICINES LIST RESTRICTIONS CHECKLIST

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to Section H of the Pharmaceutical Schedule. For community funding, see the Special Authority Criteria.

PRESCRIBER	PATIENT:
Name:	Name:
Ward:	NHI:

Pegylated interferon alfa-2a

INITIATION – Chronic hepatitis C - genotype 1, 4, 5 or 6 infection or co-infection with HIV or genotype 2 or 3 post liver transplant Re-assessment required after 48 weeks			
Prerequisites (tick boxes where appropriate)			
Patient has chronic hepatitis C, genotype 1, 4, 5 or 6 infection or O Patient has chronic hepatitis C and is co-infected with HIV or O Patient has chronic hepatitis C genotype 2 or 3 and has received a liver transplant			
Note: Consider stopping treatment if there is absence of a virological response (defined as at least a 2-log reduction in viral load) following 12 weeks of treatment since this is predictive of treatment failure. Consider reducing treatment to 24 weeks if serum HCV RNA level at Week 4 is undetectable by sensitive PCR assay (less than 50IU/ml) AND Baseline serum HCV RNA is less than 400,000IU/ml.			
CONTINUATION – Chronic hepatitis C - genotype 1 infection Re-assessment required after 48 weeks Prerequisites (tick boxes where appropriate)			
 Prescribed by, or recommended by a gastroenterologist, infectious disease specialist or general physician, or in accordance with a protocol or guideline that has been endorsed by the Health NZ Hospital. and 			
Patient has chronic hepatitis C, genotype 1 and			
O Patient has had previous treatment with pegylated interferon and ribavirin and			
or O Patient has responder relapsed O Patient was a partial responder			
Patient is to be treated in combination with boceprevir			
INITIATION – Chronic Hepatitis C - genotype 1 infection treatment more than 4 years prior Re-assessment required after 48 weeks Prerequisites (tick boxes where appropriate)			
O Prescribed by, or recommended by a gastroenterologist, infectious disease specialist or general physician, or in accordance with a protocol or guideline that has been endorsed by the Health NZ Hospital.			
O Patient has chronic hepatitis C, genotype 1 and			
O Patient has had previous treatment with pegylated interferon and ribavirin			
O Patient has responder relapsed			
O Patient was a partial responder			
O Patient received interferon treatment prior to 2004			
O Patient is to be treated in combination with boceprevir			

HOSPITAL MEDICINES LIST RESTRICTIONS CHECKLIST

Use this checklist to determine if a patient meets the restrictions for funding in the hospital setting	J. For more details, refer to Section H of the Pharmaceutical
Schedule. For community funding, see the Special Authority Criteria.	

PRESCRIBER	PATIENT:	
Name:	Name:	
Ward:	NHI:	
Pegylated interferon alfa-2a - continued		
INITIATION – Chronic hepatitis C - genotype 2 or 3 infection without co-it Re-assessment required after 6 months Prerequisites (tick box where appropriate) O Patient has chronic hepatitis C, genotype 2 or 3 infection	nfection with HIV	
INITIATION – Hepatitis B Re-assessment required after 48 weeks Prerequisites (tick boxes where appropriate) O Prescribed by, or recommended by a gastroenterologist, infectious of guideline that has been endorsed by the Health NZ Hospital. and	lisease specialist or general physician, or in accordance with a protocol or	
Patient has confirmed Hepatitis B infection (HBsAg positive for and Patient is Hepatitis B treatment-naive and ALT > 2 times Upper Limit of Normal and HBV DNA < 10 log10 IU/ml and O HBeAg positive or O Serum HBV DNA greater than or equal to 2,000 units/m moderate fibrosis)	r more than 6 months) I and significant fibrosis (greater than or equal to Metavir Stage F2 or	
and Compensated liver disease and No continuing alcohol abuse or intravenous drug use and Not co-infected with HCV, HIV or HDV and Neither ALT nor AST > 10 times upper limit of normal and No history of hypersensitivity or contraindications to pegylated	l interferon	
INITIATION – myeloproliferative disorder or cutaneous T cell lymphoma Re-assessment required after 12 months Prerequisites (tick boxes where appropriate)		
or Patient has a cutaneous T cell lymphoma* Patient has a myeloproliferative disorder* and Patient is intolerant of hydroxyurea and Treatment with anagrelide and busulfan is not clinically	appropriate	
or O Patient has a myeloproliferative disorder and O Patient is pregnant, planning pregnancy or lactating		

HOSPITAL MEDICINES LIST RESTRICTIONS CHECKLIST

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PRESCRIBER	PATIENT:
Name:	Name:
Ward:	NHI:
Pegylated interferon alfa-2a - continued	
CONTINUATION – myeloproliferative disorder or cutaneous T cell lymph Re-assessment required after 12 months Prerequisites (tick boxes where appropriate) No evidence of disease progression and O The treatment remains appropriate and patient is benefitting fr and O Patient has a cutaneous T cell lymphoma* or O Patient has a myeloproliferative disorder* and	rom treatment
Note: Indications marked with * are unapproved indications	
Re-assessment required after 12 months Prerequisites (tick box where appropriate) O Prescribed by, or recommended by an ophthalmologist, or in accord Hospital. and O Patient has ocular surface squamous neoplasia*	ance with a protocol or guideline that has been endorsed by the Health NZ
CONTINUATION – ocular surface squamous neoplasia Re-assessment required after 12 months Prerequisites (tick box where appropriate) O Prescribed by, or recommended by an ophthalmologist, or in accord Hospital. and O The treatment remains appropriate and patient is benefitting from tre Note: Indications marked with * are unapproved indications	ance with a protocol or guideline that has been endorsed by the Health NZ eatment
INITIATION – post-allogenic bone marrow transplant Re-assessment required after 3 months Prerequisites (tick box where appropriate) O Patient has received an allogeneic bone marrow transplant* and has CONTINUATION – post-allogenic bone marrow transplant Re-assessment required after 3 months	s evidence of disease relapse
Prerequisites (tick box where appropriate) O Patient is responding and ongoing treatment remains appropriate Note: Indications marked with * are unapproved indications	

Signed: Date: