

NC Medicaid Pharmacy Prior Approval Request for Viekira

Beneficiary Information

		5. Beneficiary Gender:
Prescriber Information		
6. Prescribing Provider NPI #:		
7. Requester Contact Information - N		
Drug Information		
8. Drug Name:	9. Strength:	10. Quantity Per 30 Days: <u>112</u>
11. Length of Therapy (in days):	☐ 12 weeks ☐ 24 Weeks	
Clinical Information		
Total Length of Therapy (Check ONE):		
\square 12 weeks = Genotype 1a, without cirrho	osis, or genotype 1b, with cirrhosis	
☐ 24 weeks = Genotype 1a, with compens		
1. Is the beneficiary is 18 years of age or olde		
	npensated cirrhosis or confirmed genotype 1	la without cirrnosis or with
compensated cirrhosis in combination wit	n ribavirin? 🗆 Yes 🗆 No	
Genotype is:	. 1 h (ihla ah ainnh anin)ill hunadhna anh in alh	د ما المان ما المان
2. For all treatment courses except genotype	16 (Without cirrnosis), will treatment include	e the use of ribavirin?
☐ Yes ☐ No 3. As the provider, are you reasonably certain	a that treatment will improve the handician	's averall health status?
☐ Yes ☐No	i that treatment will improve the beneficiary	s over all fleatth status:
4. Has the provider assessed for laboratory a	nd clinical evidence of henatic decompensat	ion? 🗆 Ves 🗆 No
5. Does the beneficiary have cirrhosis? \square Y		
	es in No II allower is yes, please allower the r or clinical signs and symptoms of hepatic deco	
encephalopathy, variceal hemorrh		, p
		evels at baseline and during the first four weeks of
starting treatment and as clinically	indicated? ☐ Yes ☐ No	
6. Is Viekira Pak being used in combination	with other protease inhibitors used to treat	CHC (i.e. boceprevir, simeprevir, or telaprevir) or
in combination with another nucleotide	e NS5B polymerase inhibitor such as Sovaldi	* (sofosbuvir)?
☐ Yes ☐ No		
7. Is the beneficiary using Viekira Pak in co	mbination with another NS5A inhibitor? \Box Y	es □ No
		eve a SVR (defined as a lower limit HCV RNA of 25 ment regimen consisting of Sofosbuvir? \(\text{Yes} \subseteq \text{No}\)
9. Is the beneficiary requesting the regimer	n for re-treatment and either failed to achiev	ve a SVR (defined as a lower limit HCV RNA of 25
		tment regimen consisting of Ledipasvir? \Box Yes \Box No
		assification score of Child Class B or C (VIEKIRA
· · · · · · · · · · · · · · · · · · ·	s with moderate to severe hepatic impairme	•

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 12. Does the beneficiary have decompensated liver disease as defined by Child-Pugh classification score of Child Class B or C (VIEKIRA PAK™ is contraindicated in beneficiaries with moderate to severe hepatic impairment (Child-Pugh B and C)? Yes □ No 13. Has the beneficiary attempted a previous course of therapy with Viekira Pak? □ Yes □ No 14. Does the beneficiary have any FDA labeled contraindications to Viekira Pak? □ Yes □ No 15. Has the beneficiary tried and failed 2 preferred medications in this class or does the beneficiary have a reason or contraindication to the preferred medications in the class? □ Yes □ No Please list t/f medications and/or any contraindications to the preferred medications:
 13. Has the beneficiary attempted a previous course of therapy with Viekira Pak? No 14. Does the beneficiary have any FDA labeled contraindications to Viekira Pak? No 15. Has the beneficiary tried and failed 2 preferred medications in this class or does the beneficiary have a reason or contraindication to the
 14. Does the beneficiary have any FDA labeled contraindications to Viekira Pak? See No 15. Has the beneficiary tried and failed 2 preferred medications in this class or does the beneficiary have a reason or contraindication to the
15. Has the beneficiary tried and failed 2 preferred medications in this class or does the beneficiary have a reason or contraindication to the
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Signature of Prescriber: Date:
(Prescriber Signature Mandatory)

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

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