



PRIOR AUTHORIZATION REQUEST FORM
DESCOVY®

For authorization, please answer each question and fax this form PLUS chart notes back to Real Rx at 385-425-4052.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for assistance 385-425-5094.

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:	ID#:
DOB:	Gender:	Physician:
Office Phone:	Office Fax:	Office Contact:

Height/Weight:

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.

Product being requested: Descovy® (emtricitabine and tenofovir alafenamide)

Dosing/Frequency: _____

If the request is for reauthorization, proceed to reauthorization section.

Questions	Yes	No	Comments/Notes
1. Is this request for an expedited review? By checking the “ Yes ” box to request an expedited review (24 hours), you are certifying that applying the standard review time frame (72 hours) may place the member's life, health, or ability to regain maximum function in serious jeopardy.	<input type="checkbox"/>	<input type="checkbox"/>	
2. Does the member have documentation of renal dysfunction or worsening of renal dysfunction after starting a tenofovir disoproxil fumarate regimen?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Is the member taking any medications that are considered medically necessary and likely to cause or exacerbate renal dysfunction?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
4. Does the member have an intolerance or contraindication to emtricitabine and tenofovir disoproxil fumarate (generic Truvada®)?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
5. Does the member have documentation of osteoporosis confirmed by DEXA Scan OR do serial DEXA scans show osteopenia with progression of bone loss?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
6. For treatment of HIV infection, will Descovy® be used as part of an antiretroviral treatment (ART) regimen?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
7. For PrEP, is the request for an at-risk adult or adolescent (≥ 35 kg) to reduce the risk of sexually acquired HIV-1 infection?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
8. For PrEP, is the member confirmed to be HIV-negative within 30 days prior to initiation of therapy?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation

REAUTHORIZATION			
1. Is the request for reauthorization of therapy?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Has Descovy shown to be tolerable and effective?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Does the member have a continued medical need for therapy?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
4. For PrEP, does the member have a documented negative HIV-1 tests every 3 months?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.			
Additional information:			
Physician Signature:			

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Policy: PHARM-111
 Origination Date: 10/29/2020
 Reviewed/Revised Date: 01/17/2024
 Next Review Date: 01/17/2025
 Current Effective Date: 02/01/2024

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