

US Family Health Plan
 Prior Authorization Request Form for
Enasidenib (Idhifa) and Ivosidenib (Tibsovo)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be faxed to 855-273-5735

OR

The patient may attach the completed form to the prescription and **mail** it to:

Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? Call 1-877-880-7007

Prior authorization for Idhifa will expire in 1 year. Prior authorization for Tibsovo is indefinite.

Step 1 Please complete patient and physician information (please print):

Patient Name: _____ Address: _____ Sponsor ID #: _____ Date of Birth: _____	Physician Name: _____ Address: _____ Phone #: _____ Secure Fax #: _____
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Step 2 Please complete the clinical assessment:

1. Is the patient GREATER THAN or EQUAL TO 18 years of age?	<input type="checkbox"/> Yes Proceed to question 2	<input type="checkbox"/> No STOP Coverage not approved
2. Is the requested medication being prescribed by or in consultation with hematologist or oncologist?	<input type="checkbox"/> Yes Proceed to question 3	<input type="checkbox"/> No STOP Coverage not approved
3. For which medication is coverage being requested?	<input type="checkbox"/> Idhifa Proceed to question 4	<input type="checkbox"/> Tibsovo Proceed to question 11
4. Does the patient have a diagnosis of relapsed or refractory acute myelogenous leukemia (AML)?	<input type="checkbox"/> Yes Proceed to question 5	<input type="checkbox"/> No Proceed to question 9
5. Does the patient exhibit the IDH2 mutation as determined by an FDA approved test?	<input type="checkbox"/> Yes Proceed to question 6	<input type="checkbox"/> No Proceed to question 9
6. Has the patient received this medication under the TRICARE benefit in the last 6 months? Please choose "No" if the patient did not previously have a TRICARE approved PA for Idhifa.	<input type="checkbox"/> Yes Proceed to question 7	<input type="checkbox"/> No Proceed to question 8
7. Has the patient experienced disease progression?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Sign and date on next page

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<p>8. Will the requested medication be used in combination with standard chemotherapy protocols?</p>	<p><input type="checkbox"/> Yes Sign and date below</p>	<p><input type="checkbox"/> No STOP Coverage not approved</p>
<p>9. Please provide the diagnosis.</p>	<hr/> <p>Proceed to question 10 Please provide the diagnosis.</p>	
<p>10. Is the diagnosis cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B</p>	<p><input type="checkbox"/> Yes Sign and date below</p>	<p><input type="checkbox"/> No STOP coverage not approved</p>
<p>11. Does the patient have a diagnosis of relapsed/refractory acute myeloid leukemia (AML) with a susceptible isocitrate dehydrogenase-1 (IDH1) mutation as detected by a FDA-approved test?</p>	<p><input type="checkbox"/> Yes Proceed to question 19</p>	<p><input type="checkbox"/> No Proceed to question 12</p>
<p>12. Has the patient been newly diagnosed with acute myelogenous leukemia (AML)?</p>	<p><input type="checkbox"/> Yes Proceed to question 13</p>	<p><input type="checkbox"/> No Proceed to question 16</p>
<p>13. Is the patient using Tibsovo as monotherapy OR in combination with azacitidine (Vidaza)?</p>	<p><input type="checkbox"/> Yes Proceed to question 14</p>	<p><input type="checkbox"/> No Proceed to question 16</p>
<p>14. Is the patient GREATER THAN or EQUAL TO 75 years of age?</p>	<p><input type="checkbox"/> Yes Proceed to question 19</p>	<p><input type="checkbox"/> No Proceed to question 15</p>
<p>15. Does the patient have comorbidities that preclude use of intensive induction chemotherapy with a susceptible IDH1 mutation as detected by a FDA-approved test?</p>	<p><input type="checkbox"/> Yes Proceed to question 19</p>	<p><input type="checkbox"/> No Proceed to question 16</p>
<p>16. Does the patient have previously treated, locally advanced, or metastatic cholangiocarcinoma with an IDH1 mutation as detected by a FDA-approved test?</p>	<p><input type="checkbox"/> Yes Proceed to question 19</p>	<p><input type="checkbox"/> No Proceed to question 17</p>
<p>17. Please provide the diagnosis.</p>	<hr/> <p>Proceed to question 18</p>	
<p>18. Is the diagnosis cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B</p>	<p><input type="checkbox"/> Yes Proceed to question 19</p>	<p><input type="checkbox"/> No STOP Coverage not approved</p>
<p>19. Will the patient be monitored for differentiation syndrome?</p>	<p><input type="checkbox"/> Yes Proceed to question 20</p>	<p><input type="checkbox"/> No STOP Coverage not approved</p>

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20. Will the patient be monitored for Guillain-Barre Syndrome?

Yes
Sign and date below

No
STOP
Coverage not approved

Step 3 I certify the above is true to the best of my knowledge. Please sign and date:

Prescriber Signature

Date

[05 April 2023]