ForwardHealth requires certain information to enable the programs to authorize and pay for medical services provided to eligible members.

Members of ForwardHealth programs are required to give providers full, correct, and truthful information for the submission of correct and complete claims for reimbursement. Per Wis. Admin. Code § DHS 104.02(4), this information should include, but is not limited to, information concerning enrollment status, accurate name, address, and member ID number.

Under Wis. Stat. § 49.45(4), personally identifiable information about program applicants and members is confidential and is only used for purposes directly related to ForwardHealth administration, such as determining eligibility of the applicant, processing prior authorization (PA) requests, or processing provider claims for reimbursement. Failure to supply the information requested by the form may result in denial of PA or payment for the services.

The use of this form is mandatory when requesting PA for certain drugs. Refer the applicable service-specific publications for service restrictions and additional documentation requirements. Provide enough information for ForwardHealth to make a determination about the request. Prescribers and pharmacy providers are required to retain a completed copy of the form.

INSTRUCTIONS
Prescribers are required to complete and sign the Prior Authorization/Preferred Drug List (PA/PDL) for Cytokine and Cell Adhesion Molecule (CAM) Antagonist Drugs for Hidradenitis Suppurativa, F-01674. Pharmacy providers are required to use the PA/PDL for Cytokine and CAM Antagonist Drugs for Hidradenitis Suppurativa form to request PA using the Specialized Transmission Approval Technology-Prior Authorization (STAT-PA) system or by submitting a PA request on the ForwardHealth Portal, by fax, or by mail. Prescribers and pharmacy providers are required to retain a completed copy of the form.

Pharmacy providers may submit PA requests on a PA/PDL form in one of the following ways:

1) For STAT-PA requests, pharmacy providers should call 800-947-1197.

2) For requests submitted on the ForwardHealth Portal, pharmacy providers may access www.forwardhealth.wi.gov/.

3) For PA requests submitted by fax, pharmacy providers should submit a Prior Authorization Request Form (PA/RF), F-11018, and the appropriate PA/PDL form to ForwardHealth at 608-221-8616.

4) For PA requests submitted by mail, pharmacy providers should submit a PA/RF and the appropriate PA/PDL form to the following address:

   ForwardHealth
   Prior Authorization
   Ste 88
   313 Blettner Blvd
   Madison WI  53784

The provision of services that are greater than or significantly different from those authorized may result in nonpayment of the billing claim(s).

SECTION I – MEMBER INFORMATION

Element 1 – Name – Member
Enter the member’s last name, first name, and middle initial. Use Wisconsin’s Enrollment Verification System (EVS) to obtain the correct spelling of the member’s name. If the name or spelling of the name on the ForwardHealth identification card and the EVS do not match, use the spelling from the EVS.

Element 2 – Member Identification Number
Enter the member ID. Do not enter any other numbers or letters. Use the ForwardHealth card or the EVS to obtain the correct member ID.

Element 3 – Date of Birth – Member
Enter the member’s date of birth in MM/DD/CCYY format.
SECTION II – PRESCRIPTION INFORMATION

Element 4 – Drug Name
Enter the drug name.

Element 5 – Drug Strength
Enter the strength of the drug listed in Element 4.

Element 6 – Date Prescription Written
Enter the date the prescription was written.

Element 7 – Directions for Use
Enter the directions for use of the drug.

Element 8 – Name – Prescriber
Enter the name of the prescriber.

Element 9 – National Provider Identifier (NPI) – Prescriber
Enter the prescribing provider’s National Provider Identifier (NPI).

Element 10 – Address – Prescriber
Enter the address (street, city, state, ZIP+4 code) of the prescribing provider.

Element 11 – Telephone Number – Prescriber
Enter the telephone number, including the area code, of the prescribing provider.

SECTION III – CLINICAL INFORMATION FOR HIDRADENITIS SUPPURATIVA

Element 12 – Diagnosis Code and Description
Enter the appropriate and most-specific International Classification of Diseases (ICD) diagnosis code and description most relevant to the drug requested. The ICD diagnosis code must correspond with the ICD description.

Element 13
Check the appropriate box to indicate whether or not the member has hidradenitis suppurativa.

Element 14
Check the appropriate box to indicate whether or not the prescription was written by a dermatologist or through a dermatology consultation.

Element 15
Check the appropriate box to indicate whether or not the member has recurrent abscesses with sinus tracts and scarring.

Element 16
Check the appropriate box to indicate whether or not the member has had laser therapy, excision, or deroofing surgery to treat hidradenitis suppurativa.

Element 17
Check the appropriate box to indicate whether or not the member is currently using the requested cytokine and CAM antagonist drug. If yes, indicate the approximate date the therapy was started.

Element 18
Check the appropriate box next to the drugs listed that the member has taken for at least three consecutive months and experienced an unsatisfactory therapeutic response or experienced a clinically significant adverse reaction. Indicate the drug name, dose, approximate dates the drug(s) was taken, and the reason for discontinuation.

Check “none” if appropriate, and indicate the reason the member is unable to use the drugs listed.

Note: If none, a copy of the member’s medical records must be submitted with the PA request to support the condition being treated, details regarding previous medication use and outline the member’s current treatment plan.
Element 19
Check the appropriate box to indicate whether or not the member has attempted other drug therapies for hidradenitis suppurativa (e.g., topicals or IV immunomodulators such as infliximab). If yes, indicate the drug names, dose, specific details about the treatment response, and the approximate dates each drug was taken in the space provided. If additional space is needed, continue documentation in Section VI of the form.

SECTION IV – AUTHORIZED SIGNATURE

Element 20 – Signature – Prescriber
The prescriber is required to complete and sign this form.

Element 21 – Date Signed
Enter the month, day, and year the form was signed in MM/DD/CCYY format.

SECTION V – FOR PHARMACY PROVIDERS USING STAT-PA

Element 22 – National Drug Code
Enter the appropriate 11-digit National Drug Code for each drug.

Element 23 – Days’ Supply Requested
Enter the requested days’ supply, up to 365 days.

Element 24 – NPI
Enter the NPI. Also enter the taxonomy code if the pharmacy provider’s taxonomy code is not 333600000X.

Element 25 – Date of Service
Enter the requested first date of service (DOS) for the drug in MM/DD/CCYY format. For STAT-PA requests, the DOS may be up to 31 days in the future or up to 14 days in the past.

Element 26 – Place of Service
Enter the appropriate place of service code designating where the requested item would be provided/performed/dispensed.

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>01</td>
<td>Pharmacy</td>
</tr>
<tr>
<td>13</td>
<td>Assisted living facility</td>
</tr>
<tr>
<td>14</td>
<td>Group home</td>
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<tr>
<td>32</td>
<td>Nursing facility</td>
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<tr>
<td>34</td>
<td>Hospice</td>
</tr>
<tr>
<td>50</td>
<td>Federally qualified health center</td>
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<tr>
<td>65</td>
<td>End-stage renal disease treatment facility</td>
</tr>
<tr>
<td>72</td>
<td>Rural health clinic</td>
</tr>
</tbody>
</table>

Element 27 – Assigned PA Number
Enter the PA number assigned by the STAT-PA system.

Element 28 – Grant Date
Enter the date the PA request was approved by the STAT-PA system.

Element 29 – Expiration Date
Enter the date the PA expires as assigned by the STAT-PA system.

Element 30 – Number of Days Approved
Enter the number of days for which the PA request was approved by the STAT-PA system.

SECTION VI – ADDITIONAL INFORMATION

Element 31
Include any additional information in the space provided. Additional diagnostic and clinical information explaining the need for the product requested may be included here.