

THIS FORM CAN BE USED FOR PATIENTS NEEDING:
PRESCRIPTION DISPENSED • BENEFIT VERIFICATION • NURSING SERVICES

Pharmacy Solutions, a specialty pharmacy service brought to you by AbbVie, provides expanded patient support and gives patients more options for accessing their prescribed medication by providing the personalized support they need.

Insurance Support Services

- o Benefit verification and prior authorization assistance

Pharmacy Services

- o Medication dispensing and delivery coordination
- o Forwarding of the prescription to an in-network specialty pharmacy or patient-preferred pharmacy if Pharmacy Solutions is not in-network to dispense

Financial Assistance Research

- o AbbVie-sponsored savings card eligibility
- o Referrals to independent co-pay foundations or the AbbVie Patient Assistance Foundation

Nursing Services

Patient Care Management

- o Initial and ongoing assessments
- o Follow-up calls

PLEASE REMEMBER TO...

1. Provide front and back copies of all prescription insurance card(s).
2. Initiate home nurse injection training, if needed.
3. Have the prescriber sign at the bottom.
4. If your state requires a prescription to be written on an official state form (e.g., New York), please fax separately.

Please provide complete information to ensure timely processing.

For more information, or to be connected with a dedicated Pharmacy Solutions Partner for your office, please call us at 888-857-0668.

Product support services are available regardless of where the prescription is filled.

Please see Important Safety Information, including BOXED WARNING on Serious Infections and Malignancy, on next page. Please click here for [full Prescribing Information](#).

INDICATIONS for HUMIRA® (adalimumab)¹

HUMIRA is indicated for reducing signs and symptoms and inducing and maintaining clinical remission in pediatric patients 6 years of age and older with moderately to severely active Crohn's disease who have had an inadequate response to corticosteroids or immunomodulators such as azathioprine, 6-mercaptopurine, or methotrexate.

IMPORTANT SAFETY INFORMATION¹

SERIOUS INFECTIONS

Patients treated with HUMIRA are at increased risk for developing serious infections that may lead to hospitalization or death. Most patients who developed these infections were taking concomitant immunosuppressants such as methotrexate or corticosteroids.

Discontinue HUMIRA if a patient develops a serious infection or sepsis.

Reported infections include:

- **Active tuberculosis (TB), including reactivation of latent TB. Patients with TB have frequently presented with disseminated or extrapulmonary disease. Test patients for latent TB before HUMIRA use and during therapy. Initiate treatment for latent TB prior to HUMIRA use.**
- **Invasive fungal infections, including histoplasmosis, coccidioidomycosis, candidiasis, aspergillosis, blastomycosis, and pneumocystosis. Patients with histoplasmosis or other invasive fungal infections may present with disseminated, rather than localized, disease. Antigen and antibody testing for histoplasmosis may be negative in some patients with active infection. Consider empiric anti-fungal therapy in patients at risk for invasive fungal infections who develop severe systemic illness.**
- **Bacterial, viral, and other infections due to opportunistic pathogens, including Legionella and Listeria.**

Carefully consider the risks and benefits of treatment with HUMIRA prior to initiating therapy in patients: 1. with chronic or recurrent infection, 2. who have been exposed to TB, 3. with a history of opportunistic infection, 4. who resided in or traveled in regions where mycoses are endemic, 5. with underlying conditions that may predispose them to infection. Monitor patients closely for the development of signs and symptoms of infection during and after treatment with HUMIRA, including the possible development of TB in patients who tested negative for latent TB infection prior to initiating therapy.

- Do not start HUMIRA during an active infection, including localized infections.
- Patients older than 65 years, patients with co-morbid conditions, and/or patients taking concomitant immunosuppressants may be at greater risk of infection.
- If an infection develops, monitor carefully and initiate appropriate therapy.
- Drug interactions with biologic products: A higher rate of serious infections has been observed in rheumatoid arthritis patients treated with rituximab who received subsequent treatment with a TNF blocker. Concurrent use of HUMIRA with biologic DMARDs (e.g., anakinra or abatacept) or other TNF blockers is not recommended based on the possible increased risk for infections and other potential pharmacological interactions.

MALIGNANCY

Lymphoma and other malignancies, some fatal, have been reported in children and adolescent patients treated with TNF blockers, including HUMIRA. Postmarketing cases of hepatosplenic T-cell lymphoma (HSTCL), a rare type of T-cell lymphoma, have been reported in patients treated with TNF blockers, including HUMIRA. These cases have had a very aggressive disease course and have been fatal. The majority of reported TNF blocker cases have occurred in patients with Crohn's disease or ulcerative colitis and the majority were in adolescent and young adult males. Almost all of these patients had received treatment with azathioprine or 6-mercaptopurine concomitantly with a TNF blocker at or prior to diagnosis. It is uncertain whether the occurrence of HSTCL is related to use of a TNF blocker or a TNF blocker in combination with these other immunosuppressants.

- Consider the risks and benefits of HUMIRA treatment prior to initiating or continuing therapy in a patient with known malignancy.
- In clinical trials, more cases of malignancies were observed among HUMIRA-treated patients compared to control patients.

- Non-melanoma skin cancer (NMSC) was reported during clinical trials for HUMIRA-treated patients. Examine all patients, particularly those with a history of prolonged immunosuppressant or PUVA therapy, for the presence of NMSC prior to and during treatment with HUMIRA.
- In HUMIRA clinical trials, there was an approximate 3-fold higher rate of lymphoma than expected in the general U.S. population. Patients with chronic inflammatory diseases, particularly those with highly active disease and/or chronic exposure to immunosuppressant therapies, may be at higher risk of lymphoma than the general population, even in the absence of TNF blockers.
- Postmarketing cases of acute and chronic leukemia were reported with TNF blocker use. Approximately half of the postmarketing cases of malignancies in children, adolescents, and young adults receiving TNF blockers were lymphomas; other cases included rare malignancies associated with immunosuppression and malignancies not usually observed in children and adolescents.

HYPERSENSITIVITY

- Anaphylaxis and angioneurotic edema have been reported following HUMIRA administration. If a serious allergic reaction occurs, stop HUMIRA and institute appropriate therapy.

HEPATITIS B VIRUS REACTIVATION

- Use of TNF blockers, including HUMIRA, may increase the risk of reactivation of hepatitis B virus (HBV) in patients who are chronic carriers. Some cases have been fatal.
- Evaluate patients at risk for HBV infection for prior evidence of HBV infection before initiating TNF blocker therapy.
- Exercise caution in patients who are carriers of HBV and monitor them during and after HUMIRA treatment.
- Discontinue HUMIRA and begin antiviral therapy in patients who develop HBV reactivation. Exercise caution when resuming HUMIRA after HBV treatment.

NEUROLOGIC REACTIONS

- TNF blockers, including HUMIRA, have been associated with rare cases of new onset or exacerbation of central nervous system and peripheral demyelinating diseases, including multiple sclerosis, optic neuritis, and Guillain-Barré syndrome.
- Exercise caution when considering HUMIRA for patients with these disorders; discontinuation of HUMIRA should be considered if any of these disorders develop.
- There is a known association between intermediate uveitis and central demyelinating disorders.

HEMATOLOGIC REACTIONS

- Rare reports of pancytopenia, including aplastic anemia, have been reported with TNF blockers. Medically significant cytopenia has been infrequently reported with HUMIRA.
- Consider stopping HUMIRA if significant hematologic abnormalities occur.

CONGESTIVE HEART FAILURE

- Worsening or new onset congestive heart failure (CHF) may occur; exercise caution and monitor carefully.

AUTOIMMUNITY

- Treatment with HUMIRA may result in the formation of autoantibodies and, rarely, in development of a lupus-like syndrome. Discontinue treatment if symptoms of a lupus-like syndrome develop.

IMMUNIZATIONS

- Patients on HUMIRA should not receive live vaccines.
- Pediatric patients, if possible, should be brought up to date with all immunizations before initiating HUMIRA therapy.
- The safety of administering live or live-attenuated vaccines in infants exposed to HUMIRA *in utero* is unknown. Risks and benefits should be considered prior to vaccinating (live or live-attenuated) exposed infants.

ADVERSE REACTIONS

- The most common adverse reactions in HUMIRA clinical trials (>10%) were: infections (e.g., upper respiratory, sinusitis), injection site reactions, headache, and rash.

Reference: 1. HUMIRA Injection [package insert]. North Chicago, IL: AbbVie Inc.

HUMIRA® (adalimumab) REFERRAL AND PRESCRIPTION FORM
SIGN AND FAX THIS FORM TO 800-266-2065. FOR QUESTIONS, PLEASE CALL 888-857-0668.

PATIENT AND PRESCRIBER INFORMATION	PATIENT INFORMATION SSN (Last 4 ONLY) ____ ____ ____ ____	PRESCRIBER INFORMATION <input type="checkbox"/> MD <input type="checkbox"/> DO <input type="checkbox"/> Other: _____
	First Name: _____ MI: _____	Prescriber Name: _____
	Last Name: _____	Specialty: <input type="checkbox"/> Gastro <input type="checkbox"/> Other: _____
	DOB: _____ Weight (lbs): _____ Sex: <input type="checkbox"/> M <input type="checkbox"/> F	NPI/Provider #: _____ State License #: _____
	Address: _____	Office Name: _____
	City/State/Zip: _____	Contact: _____
	Primary Phone: _____ <input type="checkbox"/> H <input type="checkbox"/> W <input type="checkbox"/> M	Address: _____
	Alternate Phone: _____ <input type="checkbox"/> H <input type="checkbox"/> W <input type="checkbox"/> M	City/State/Zip: _____
Drug Allergies: _____	Phone: _____ Fax: _____	

Fax a copy of the front and back of prescription insurance card(s) or fill in the information below

Primary Insurance: _____	Secondary Insurance: _____
Phone: _____	Phone: _____
Cardholder ID #: _____ Group #: _____	Cardholder ID #: _____ Group #: _____
PCN: _____ BIN: _____	PCN: _____ BIN: _____
Policyholder Name: _____ DOB: _____	Policyholder Name: _____ DOB: _____

INSURANCE INFORMATION

BV ONLY **BENEFIT VERIFICATION ONLY**

I do **not** want to prescribe HUMIRA at this time, but please verify drug coverage.

PATIENT'S DIAGNOSIS Date of Diagnosis: _____

Pediatric Crohn's Disease ICD-10: _____ Other (include code): _____

SHIPPING PREFERENCE Date needed: _____ Deliver medication to the patient Deliver medication to the prescriber

PRESCRIPTION New Restart Continuing

Starting Therapy

Pediatric Crohn's disease — **weight 17 kg (37 lbs) to <40 kg (88 lbs)** Two 40 mg subcutaneous inj. Day 1, one 40 mg subcutaneous inj. Day 15 #3 syringes No Refills
 Starter Package 40 mg/0.8 mL

Pediatric Crohn's disease — **weight ≥40 kg (88 lbs)** Four 40 mg subcutaneous inj. Day 1, two 40 mg subcutaneous inj. Day 15 #6 pens #6 syringes No Refills
 Starter Package 40 mg/0.8 mL Two 40 mg subcutaneous inj. Day 1, two 40 mg subcutaneous inj. Day 2, two 40 mg subcutaneous inj. Day 15 #6 pens #6 syringes No Refills
(Note: Pens available for patients ≥88 lbs)

Ongoing Therapy

	1 month	3 month	
<input type="checkbox"/> Pediatric Crohn's disease — weight 17 kg (37 lbs) to <40 kg (88 lbs)	<input type="checkbox"/> One 20 mg subcutaneous inj. Day 29 and every other week thereafter*	<input type="checkbox"/> #2 syringes	<input type="checkbox"/> #6 syringes Refills: _____
<input type="checkbox"/> Pediatric Crohn's disease — weight ≥40 kg (88 lbs)	<input type="checkbox"/> One 40 mg subcutaneous inj. Day 29 and every other week thereafter	<input type="checkbox"/> #2 pens <input type="checkbox"/> #2 syringes	<input type="checkbox"/> #6 pens <input type="checkbox"/> #6 syringes Refills: _____

Other HUMIRA _____ SIG: _____ Qty: _____ Refills: _____

* 20 mg only available in pre-filled syringe

PRESCRIBER SIGNATURE: PRESCRIBER MUST MANUALLY SIGN (RUBBER STAMPS, SIGNATURE BY OTHER OFFICE PERSONNEL FOR THE PRESCRIBER, AND COMPUTER-GENERATED SIGNATURES WILL NOT BE ACCEPTED), OR SEND AN ELECTRONIC PRESCRIPTION TO PHARMACY SOLUTIONS, AN ABBVIE COMPANY.

<input type="checkbox"/> Dispense as written/Do not substitute Date _____	<input type="checkbox"/> Substitution permitted/Brand exchange permitted Date _____
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I authorize Pharmacy Solutions and its employees to serve as my agent for the sole purpose of obtaining patient benefit information and the necessary prior authorization forms when dealing with Health Plans and Pharmacy Benefits Managers (PBMs), if the plan or PBM requires such authorization.

For states requiring handwritten expressions of Product Selection, use this area (e.g., medically necessary, may not substitute, dispense as written, etc.)

The information contained in this communication is confidential and intended for the addressee. It may contain Protected Health Information (PHI) under HIPAA. PHI is personal and sensitive information related to a person's health. This information is sent to you under circumstances when a participant's authorization is not required. You, the recipient, are obligated to maintain it in a safe, secure, and confidential manner. Redisclosure, unless permitted by law, is prohibited. If you are not the intended recipient, you are hereby notified that dissemination, disclosure, copying, or distribution of this information is strictly prohibited and may be unlawful. Please notify sender immediately to arrange for return of this document.

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