As a condition of your site’s authorization to infuse TYSABRI® (natalizumab), this Pre-infusion Patient Checklist must be completed for each patient prior to each infusion. This page must be submitted on-line (www.touchprogram.com) OR faxed to Biogen (1-800-840-1278) within 1 day of the patient’s visit and a copy retained in the patient’s record whether the patient has been infused or not.

**STEP 1:** Ensure that the patient is currently authorized to receive TYSABRI for MS or Crohn’s disease.

You must refer to the patient’s record prior to every infusion.

- If the patient did not receive his or her previous infusion, and physician clearance was required, you must confirm authorization from the prescriber before providing the current infusion.
- Confirm the patient status is listed as “Authorized” on TOUCH® On-Line (www.touchprogram.com) OR a Notice of Patient Authorization on file and that you have not received a Notice of Patient Discontinuation (paper-based process).

Is the patient currently authorized to receive TYSABRI?

<table>
<thead>
<tr>
<th>Yes</th>
<th>No</th>
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Add the initials of the healthcare provider who reviewed this checklist.

**STEP 2:** Confirm that the patient has read and understood the Patient Medication Guide.

The patient must read the Patient Medication Guide prior to every infusion. **Has the patient received and read the Patient Medication Guide, including the section “What should I tell my doctor and nurse before each infusion of TYSABRI?”**

<table>
<thead>
<tr>
<th>Yes</th>
<th>No</th>
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**STEP 3:** Read aloud and mark “Yes” or “No” for the patient’s answers to the following questions:

1. Over the past month, have you had any new or worsening medical problems (such as a new or sudden change in your thinking, eyesight, balance, strength, or other problems) that have persisted over several days?  

2. Do you have a medical condition that can weaken your immune system, such as HIV infection or AIDS, leukemia or lymphoma, or an organ transplant that may suggest your body is not able to fight infections well?  

3. In the past month have you taken, or are you currently on, any medicines other than steroid medicines to treat cancer or **Crohn’s disease** or any other medicines that weaken your immune system? (Review the list on the next page with the patient.)

**STEP 4:** Record Prescriber Authorization (if required):  

If the patient answered YES to question 1, 2 or 3, DO NOT INFUSE. Contact the healthcare provider who prescribed TYSABRI or a prescriber representative and review the patient’s answers.  

- After discussing the patient’s answers, did the prescriber authorize the patient to be infused?  

**STEP 5:** Record infusion information

- Not infused  

Name and signature of staff completing checklist: ____________________________ Date ________________

**STEP 6:** Submit the Pre-infusion Patient Checklist to Biogen on-line at www.touchprogram.com OR fax to 1-800-840-1278.
Please review the following list with the patient when asking question 3.

### Examples of Immunosuppressants, Antineoplastics, and Immunomodulators

#### Multiple Sclerosis

**Approved MS Therapies:**
- Alemtuzumab (Lemtrada®)
- Cladribine (Mavenclad®)
- Dimethyl fumarate (TECFIDERA®)
- Daclizumab (Zenapax®)
- Fingolimod (Gilenya®)
- Glatiramer acetate (Copaxone®)
- Interferon beta-1a (Rebif®, AVONEX®)
- Interferon beta-1b (Betaseron®, Extavia®)
- Mitoxantrone

**Immunosuppressants/Antineoplastics:**
- Azathioprine (Imuran®, Azasan®)
- Cladribine (Leustatin®)
- Cyclophosphamide (Cytoxan®, Neosar®)
- Cyclosporine (Sandimmune®, Neoral®)
- Fludarabine phosphate (Fludara®)
- Leflunomide (Arava®)
- Methotrexate (Methotrex, Trexall®)
- Mycophenolate mofetil (CellCept®)
- Pemetrexed (Alimta®)

**Additional Immunomodulators and Immunosuppressants:**
- Other interferons (Actimmune®, Infergen®, Intron® A, Pegassys®, PEG-Intron®, Rebetron®, Roferon®-A)
- Alefacept (Amevive®)
- Alemtuzumab (Campath®)
- Anakinra (Kineret®)
- Daclizumab (Zenapax®)
- Efalizumab (Raptiva®)
- Etanercept (Enbrel®)
- Infliximab (Remicade®)
- Intravenous immunoglobulin (IVIG)
- Rituximab (Rituxan®)
- Trastuzumab (Herceptin®)

#### Crohn’s Disease

**Approved TNF-α inhibitors for Crohn’s disease:**
- Infliximab (Remicade®)
- Adalimumab (Humira®)

**Immunosuppressants/Antineoplastics:**
- Approved TNF-α inhibitors
- Azathioprine (Imuran®, Azasan®)
- Chlorambucil (Leukeran®)
- Cladribine (Leustatin®)
- Cyclophosphamide (Cytoxan®, Neosar®)
- Cyclosporine (Sandimmune®, Neoral®)
- Fludarabine phosphate (Fludara®)
- Leflunomide (Arava®)
- Mercaptopurine (Purinethol®)
- Methotrexate (Methotrex®, Methotrex®, Trexall®)
- Mycophenolate mofetil (CellCept®)
- Pemetrexed (Alimta®)
- Thioguanine (Tabloid®)

**Additional Immunomodulators and Immunosuppressants:**
- Interferon beta-1a (Rebif®, AVONEX®)
- Interferon beta-1b (Betaseron®)
- Alefacept (Amevive®)
- Abatacept (Orencia®)
- Anakinra (Kineret®)
- Daclizumab (Zenapax®)
- Efalizumab (Raptiva®)
- Etanercept (Enbrel®)
- Glatiramer acetate (Copaxone®)
- Intravenous immunoglobulin (IVIG)
- Mitoxantrone (Novantrone®)
- Other interferons (Actimmune®, Infergen®, Intron® A, Pegassys®, PEG-Intron®, Rebetron®, Roferon®-A)
- Rituximab (Rituxan®)
- Trastuzumab (Herceptin®)
- Vedolizumab (Entyvio®)

This list does not include all drugs that can suppress the immune system.

- Patients should consult their prescribing physician regarding drugs that may be taken concurrently with TYSABRI.
- If there are any questions regarding concurrent therapy, do not infuse at this time and consult the healthcare provider who prescribed TYSABRI.

If you are unable to contact the prescriber or a prescriber representative:

Instruct the patient to contact his/her prescriber or prescriber representative and to reschedule as soon as possible. Continue efforts to reach the prescriber or prescriber representative to inform him/her of the reason(s) for not infusing this patient. You will need to confirm authorization from the prescriber or prescriber representative on the subsequent infusion.

A prescriber representative is defined as a prescriber-designated individual who can make decisions regarding patients under his or her care in their absence. A prescriber representative should be knowledgeable about the Prescribing Information, including Boxed Warning, and should be able to determine whether the patient can receive his or her infusion.

This Pre-infusion Patient Checklist is not intended to replace the infusion site’s general infusion protocol(s). Nor is this Pre-infusion Patient Checklist intended to be a substitute for consultation and review of reference materials and medical literature pertaining to individual clinical circumstances. Healthcare providers should make all treatment decisions based on the context of the situation and their clinical judgment.

Please do not make any extraneous marks on the Pre-infusion Patient Checklist. If there is information that you would like to share with Biogen and the TOUCH Prescribing Program, please contact us at 1-800-456-2255.

Please see the Prescribing Information, including BOXED WARNING, for more information.