

- To submit the START Form electronically, click [here](#). You may also upload any required documentation to this site.**
- Complete all required sections to prevent delays in support, and include copies of the front and back of the patient's medical insurance and pharmacy coverage cards and clinical documentation of required test results.**
- Please sign, date, and submit the form via fax (908-425-4840) or email (COMPASS@sanofi.com).**
Form must be submitted by prescriber's office only.
- For questions or to learn more about TZIELD COMPASS, call 1-844-778-2246 Monday through Friday, 8 AM-8 PM ET.**

***Indicates required field**

Patient Information

*Patient First Name: _____ *Patient Last Name: _____ *Sex Assigned at Birth: Male Female
*Date of Birth (mm/dd/yyyy): _____ *Patient Address: _____ *City: _____
*State: _____ *ZIP: _____ *Primary Phone #: Mobile Home (leave blank if patient is under 18 years old) _____
Email (leave blank if patient is under 18 years old): _____
Preferred Form of Communication: Phone Text Email Do not contact patient Best Time to Contact: Morning Afternoon Evening
Preferred Language: English Spanish Other _____

Parent/Legal Guardian information, if applicable (required for patients under 18 years old)

*Parent/Legal Guardian Name: _____ *Relationship to Patient: _____
*Parent/Legal Guardian Primary Phone #: Mobile Home _____ Email: _____
Secondary Parent/Legal Guardian or Caregiver Name: _____ Relationship to Patient: _____
Secondary Parent/Legal Guardian or Caregiver Primary Phone #: Mobile Home _____ Email: _____

Patient Consent

Section 7: I have read and agree to the Authorizations to Use and Disclose Health Information.

SIGN

*Patient or Parent/Legal Guardian Signature *Relationship to Patient *Date (mm/dd/yyyy)

Section 8: I have read and agree to the Patient Certifications.

[Check here](#) I have read the Text Messaging Consent in Section 8 and expressly consent to receive text messages by or on behalf of the Program.

SIGN

*Patient or Parent/Legal Guardian Signature *Relationship to Patient *Date (mm/dd/yyyy)

Additional Parent/Legal Guardian or Caregiver Name (optional) Relationship to Patient (optional) Phone Number (optional)

2 Insurance Information

Patient has no insurance (proceed to Section 3)

Please attach a copy of both sides of the patient's medical and pharmacy insurance card(s) via fax with this prescription form.

*Primary Insurance: _____
*Insurance Provider: _____ *Phone #: _____ *Policy ID #: _____
*Group #: _____ *Policy Holder Name: _____ *Policy Holder Date of Birth (mm/dd/yyyy): _____
*Policy Holder Relationship to Patient: _____ *RxBIN: _____ *RxPCN: _____
Secondary Insurance: _____ Insurance Provider: _____
Phone #: _____ Policy ID #: _____ Group #: _____ Policy Holder Name: _____
Policy Holder Date of Birth (mm/dd/yyyy): _____ Policy Holder Relationship to Patient: _____

Please see full [Prescribing Information](#), including **Boxed WARNING and patient selection criteria.**

*Patient First Name: _____ *Patient Last Name: _____ *Patient Date of Birth: _____

***Indicates required field**

Please note: Product is available through limited specialty pharmacies. Actual dispensing method and specialty pharmacy preference may be specified by the patient's insurance.

Please Select Acquisition Method

Specialty Distributor: Cardinal Specialty Distribution

Specialty Pharmacy: Amber Specialty Pharmacy Chartwell Specialty Pharmacy Orsini Specialty Pharmacy No preference Unsure

3 Prescriber Information

*Clinic Name: _____ *First Name: _____ *Last Name: _____

*Prescriber NPI: _____ *Prescriber Tax ID #: _____ *Address: _____

*City: _____ *State: _____ *ZIP: _____ *Office Contact Name: _____

*Office Contact Phone #: _____ *Fax #: _____ *Office Contact Email: _____

4 Site of Care (SOC) Information

I would like assistance from TZIELD COMPASS in identifying SOC options. My preferred SOC setting(s) include

At home with a nurse Infusion facility Both facility and home

I have already identified SOC for my patient. Patient will be treated at

Prescriber's office (SECTION 3) At home with a nurse (if address is different than SECTION 1, please list below) Infusion facility (please list below)

Both facility and at home (please list infusion site below, as well as number of doses to be administered at each location)

_____ days to be treated at facility _____ days to be treated at home

SOC details (if unknown, TZIELD COMPASS can provide support with SOC identification/options)

SOC Name: _____ SOC NPI: _____

Tax ID #: _____ Address: _____

City: _____ State: _____ ZIP: _____ SOC Contact Name: _____

SOC Contact Phone #: _____ Fax #: _____

5 Clinical Diagnosis

*Primary Diagnosis ICD-10-CM Code: E10.8 E10.9 E10.A0 E10.A2 Other (Include ICD-10-CM): _____

***Please indicate which tests have been conducted to confirm patient's diagnosis (please attach clinical documentation of these test results)**

***Confirmation of dysglycemia without overt hyperglycemia**

Oral glucose tolerance test (OGTT) (CPT* Code: 82951) Level: _____ Date test completed: _____

2-hour plasma glucose 140-199 mg/dL and/or Intervening plasma glucose level at 30, 60, or 90 minutes \geq 200 mg/dL

Fasting plasma glucose (FPG) 100-125 mg/dL (CPT* Code: 82947) Level: _____ Date test completed: _____

A1C 5.7%-6.4% or \geq 10% increase in A1C (CPT* Code: 83036) Level: _____ Date test completed: _____

***Confirmation of at least 2 pancreatic islet cell autoantibodies (select positive autoantibodies below)**

Glutamic acid decarboxylase 65 (GAD) autoantibody (CPT* Code: 86341) Date test completed: _____

Insulin autoantibody (IAA) (CPT* Code: 86337) Date test completed: _____

Insulinoma-associated antigen 2 autoantibody (IA-2A) (CPT* Code: 86341) Date test completed: _____

Zinc transporter 8 autoantibody (ZnT8A) (CPT* Code: 86341) Date test completed: _____

Islet cell autoantibody (ICA) (CPT* Code: 86341) Date test completed: _____

*I certify that the patient's clinical history and associated diagnosis confirms an autoimmune origin and does not suggest type 2 diabetes or other forms of diabetes, including but are not limited to genetic forms of diabetes, maturity-onset diabetes of the young, latent autoimmune diabetes in adults, or diabetes secondary to medications or surgery

*Complete blood count (CBC) and liver enzyme tests have been run to confirm patient has adequate hematologic function, adequate hepatic function, and does not have active infections

*I certify that the patient does not have active Epstein-Barr virus or cytomegalovirus, and undetectable viral load has been confirmed and will reconfirm prior to treatment

*I certify that the patient's clinical history demonstrates dysglycemia without overt hyperglycemia per an OGTT or alternative method if appropriate and OGTT is not available

Patient allergies: _____

Prior (within the last 12 months) and current medications, including diabetic medications: _____

Please see full [Prescribing Information](#), including **Boxed WARNING and patient selection criteria.**

*Indicates required field

6 TZIELD® (teplizumab-mzwv) Injection 2 mg/2 mL Prescription Information

Calculate the dosage using the body surface area-based dosing regimen in the [Prescribing Information](#).

To calculate BSA, click [here](#).

Mosteller formula

$$BSA (m^2) = \sqrt{\frac{[\text{height (cm)} \times \text{weight (kg)}]}{3600}}$$

*Patient Height (cm): _____ *Patient Weight (kg): _____

*Body Surface Area (BSA): _____

*Date measured: _____

Calculate using the Mosteller formula (see above)

*Quantity to Dispense

BSA

14 TZIELD 2 mg/2 mL, single-dose vials

≤1.94 m²

24 TZIELD 2 mg/2 mL, single-dose vials

>1.94 m²

Refills: No refills

By signing below, I certify that the above therapy is medically necessary and that I will supervise the patient's treatment accordingly.

SIGN

*Prescriber Signature—Dispense as Written (No Stamp Allowed)

*Date (mm/dd/yyyy)

OR

SIGN

*Prescriber Signature—Generic Substitution Allowed (No Stamp Allowed)

*Date (mm/dd/yyyy)

By signing above, I certify that (1) the information contained in this application is current, complete, and accurate to the best of my knowledge; (2) the above therapy is medically necessary and in the best interest of the patient identified above and that I will supervise the patient's treatment accordingly; (3) the patient is not currently pregnant and does not plan on becoming pregnant until at least 30 days after infusion; (4) I have obtained any consent required under federal and state law for the release and use of the patient's personal health information including diagnosis, treatment, medical, and insurance information contained on this form to Sanofi and its agents, service providers, and affiliates, including commercial and field-based teams, for purposes of benefits verification and coordination of dispensing therapy, or to otherwise assist the patient to initiate or continue the prescribed therapy and/or to evaluate the patient's eligibility for TZIELD COMPASS or other programs for TZIELD; and (5) I will not seek payment from any payer, patient, or other source for free product provided directly to the patient. I have obtained patient's permission to enroll them in TZIELD and for them to be contacted by Sanofi in connection with this application. I understand that I am under no obligation to prescribe any Sanofi therapies or to participate in TZIELD COMPASS, and that I have not received, nor will I receive, any benefit from Sanofi for prescribing a Sanofi therapy. I certify that I am a legal resident of the United States (and US territories). I authorize Sanofi and its agents to convey the above prescription by any means allowed under applicable law to the dispensing pharmacy.

Authorization to Use and Disclose Health Information

PATIENT: PLEASE READ THE FOLLOWING CAREFULLY, THEN DATE AND SIGN WHERE INDICATED IN SECTION 1 ON PAGE 2

I hereby authorize my (and/or my child's) healthcare providers, health insurance carriers, and pharmacy providers to use and disclose my (and/or my child's) individually identifying health information, including health insurance information, medical diagnosis and condition (including lab test results related to such diagnosis or supportive testing), prescription information, and name, address, and telephone number ("My Information") to Sanofi, its affiliates, and its agents and representatives ("Sanofi"), including Sanofi's commercial and field-based teams and third parties authorized by Sanofi for the following purposes in order to administer the TZIELD COMPASS Patient Support Program, including: 1. Collecting, entering, and maintaining my (and/or my child's) health information in a database to gather information on my (and/or my child's) patient experience; 2. Verifying insurance coverage, reviewing reimbursement requirements, and coordinating coverage for TZIELD® (teplizumab-mzwv) injection 2 mg/2 mL; 3. Determining eligibility for program offerings, including copay assistance, free drug or other financial assistance services, or to refer me (and/ or my child) to other programs or sources of funding; 4. Contacting me to provide education, information, and support services to me (and/or my child) related to TZIELD; 5. Contacting me to conduct market research and assess TZIELD COMPASS customer service, and to provide therapy support services designed for people prescribed TZIELD; 6. Performing data analytics with aggregated de-identified data to assess program efficiency; and contacting me about opportunities to participate in research related to TZIELD; 7. Providing me (and/or my child) with ongoing therapy support, including by communicating with healthcare professionals or service providers. All prescription-related support is limited to Sanofi product(s).

Once My Information has been disclosed to Sanofi, I understand that federal privacy laws may no longer protect it from further disclosure. However, I also understand that Sanofi has agreed to protect My Information by using reasonable efforts and disclosing it only for the purposes allowed by me in this Authorization or as otherwise required by law. I understand that I am entitled to a copy of this signed Authorization and may revoke (withdraw) this Authorization at any time by faxing a signed, written request to TZIELD COMPASS at 908-425-4840, or by mailing such request to Sanofi US, PO Box 4996, Trenton, NJ, 08650. TZIELD COMPASS will no longer seek disclosure of my (and/or my child's) health information from my (and/or my child's) healthcare providers and health insurance carriers once it has received and processed my revocation. However, revoking this Authorization will not affect any use and disclosure of the health information that has already occurred in reliance on my authorization.

Please see full [Prescribing Information](#), including **Boxed WARNING** and patient selection criteria.

Authorization to Use and Disclose Health Information (cont'd)

If I revoke this Authorization, I will no longer be able to receive TZIELD COMPASS support services. I understand that this Authorization expires 18 months from the date support is last provided under the TZIELD COMPASS Patient Support Program, or until my local state law requires expiration, subject to applicable law, unless and until I withdraw (take back) this Authorization before then, or as otherwise required by law. I understand that I do not have to sign this authorization to obtain healthcare treatment or benefits; however, in order to receive the services and communications described above, I must sign the authorization. Federal Law (including HIPAA) requires a signed authorization in order for TZIELD COMPASS to collect this information from my (and/or my child's) healthcare providers. I understand that my (and/or my child's) pharmacy, health insurers, and third-party vendors may receive remuneration (payment) from TZIELD COMPASS and Sanofi or its affiliates in exchange for providing me (and/or my child) with support services and that sharing my (and/or my child's) health information helps facilitate the support services I (and/or my child) will receive. I may reference the US Sanofi Privacy Policy at <https://www.sanofi.com/en/sanofi-us-privacy-policies> for further information regarding these rights with respect to Sanofi US.

I understand that Sanofi may de-identify My Information, including data obtained from accompanying clinical notes and/or documentation, and use it in performing research, education, business analytics, marketing studies, or for other commercial purposes, including linkage with other de-identified information Sanofi receives from other sources. I understand that members of Sanofi may share My Information, including identifiable health information, among themselves in order to de-identify it for these purposes and as needed to perform the Services or to communicate with me by mail, telephone, or email, or, if I indicate my agreement and consent in Section 1 on page 1, by text. I understand and agree that Sanofi may use My Information for these purposes and may share My Information with my healthcare providers, health insurers and specialty pharmacies.

I consent to have my data tokenized by Sanofi. Tokenization is the process of translating sensitive demographic information into a non-identifiable code called a "token" that can't be traced back to you as an individual. The demographic information needed to create a token (includes the participant's name, date of birth, gender, and address) will be collected. This token will allow linkage of your healthcare data from third-party data sets, including electronic medical records and claims databases. The tokenization software and process adhere to all appropriate data privacy and security standards and is encrypted and irreversible. This process allows Sanofi to better understand the health of participants in a comprehensive way, and thus supports more robust scientific/medical research and/or publications. Once you withdraw consent to link data, no further data links will be created. You can choose to withdraw your consent for tokenization of your data at any time by notifying Sanofi and both the token ID and patient ID would be deleted.

8 Patient Certifications

PATIENT: PLEASE READ THE FOLLOWING CAREFULLY, THEN DATE AND SIGN WHERE INDICATED IN SECTION 1 ON PAGE 2

I am enrolling in the TZIELD COMPASS Patient Support Program (the "Program") and authorize Sanofi and their affiliates and agents to provide me services under the Program, as described in the Program Enrollment Form and as may be added in the future. Such services include medication and adherence communications and support, medication dispensing support, coverage and financial assistance support, disease and medication education, and other support services (the "Services"). TZIELD COMPASS is a patient support program that helps patients to gain access to TZIELD and provides patients with education and resources related to TZIELD.

I authorize TZIELD COMPASS under the Fair Credit Reporting Act to use my demographic information to access reports on my individual credit history from consumer reporting agencies. I understand that, upon request, TZIELD COMPASS will tell me whether an individual consumer report was requested and the name and address of the agency that furnished it. I further understand and authorize TZIELD COMPASS to use any consumer reports about me and information collected from me, along with other information they obtain from public and other sources, to estimate my income in conjunction with the Patient Assistance Program eligibility determination process, if applicable. I further understand that no free product may be submitted for reimbursement to any payer, including Medicare and Medicaid; and no free product may be sold, traded, or distributed for sale. If approved for the TZIELD COMPASS Patient Assistance Program, I will not seek to have the value of any medication provided to me under this program counted toward my true-out-of-pocket (TrOOP) cost for prescription drugs for my Medicare Part D Plan.

Please see full [Prescribing Information](#), including **Boxed WARNING** and patient selection criteria.

8 Patient Certifications (cont'd)

Patients whose health insurance benefits offer, or whose health insurance plans work with, an alternative funding program are not eligible for the TZIELD COMPASS Patient Assistance Program/need-based free drug. "Alternative funding programs" sometimes refer to themselves with different names and promote their services as a benefit being offered through patients' health insurance plans. In determining whether your health insurance plan offers or works with an alternative funding program, please consider the following: (1) did a third party assist you with, or provide guidance about, the preparation of your application for this program and/or direct you to apply; (2) were you told that you were required to work with a third party or a "patient advocate" in order to receive coverage for your medicine; (3) were you told that you were required to apply to a manufacturer's patient assistance program as a prerequisite to having plan coverage for your specialty drug, including Alliance products; (4) were you told that you do not have plan coverage for your specialty drug unless alternative funding sources for your specialty drug could not be found; (5) were you required by your plan to provide personal and health information to a third-party program or enter such information into a third-party patient portal in advance of completing this application; (6) did your employer recently tell you that your specialty drug plan benefits have changed; or (7) do you have any other reason to think that your health plan may work with an alternative funding program, or an alternative funding program may be involved in your plan benefits? If you answered yes to any of these questions, please check with your plan sponsor and review your plan benefits to determine whether your plan works with a third party that provides any of the services, or is connected to any of the requirements described herein. If it does, you are not eligible to apply for this program. By applying to the TZIELD COMPASS Patient Assistance Program, you certify that the answer to all of these questions is "no."

I understand that I may be contacted by Sanofi for follow-up information in case I report an adverse event.

I give permission to be referred by my COMPASS Navigator to a Clinical Educator (CE) if I ask for clinical information regarding TZIELD. I acknowledge that a CE will provide only information about TZIELD, not any medical advice or support, and that my doctor is the best resource for any medical questions or concerns about my treatment and my disease. I give permission to Sanofi to provide me with informational and promotional materials relating to Sanofi or its affiliates products and services and/or my or my child's condition or treatment (together, the "Communications"). I also understand that the personal data I provide on this form may be shared with third parties operating on behalf of Sanofi or its affiliates to conduct market research. I authorize Sanofi and these third parties to contact me for market research purposes, though I understand that my personal data will not be sold to any third party. I understand that I do not have to enroll in the Program or receive the Communications, and that I can still receive TZIELD, as prescribed by my Healthcare Provider. I can opt out of receiving the Communications, support services offered by the Program, or being subject to market research at any time by notifying a Program representative by telephone at 1-844-778-2246 or by sending a letter to Sanofi US, PO Box 4996, Trenton, NJ 08650.

I acknowledge that by checking the Text Messaging Consent box on page 1, I expressly consent to receive text messages from or on behalf of the Program at the mobile telephone number(s) that I provide. I confirm that I am the subscriber for the mobile telephone number(s) provided, and I agree to notify Sanofi promptly if any of my number(s) change in the future. I understand that my wireless service provider's message and data rates may apply. I understand that I can opt out of future text messages at any time by texting STOP to 1-908-206-7556 from my mobile phone, and that I can get help for text messages by calling TZIELD COMPASS at 1-844-778-2246. I also understand that additional text messaging terms and conditions may be provided to me in the future as part of an opt-in confirmation text message. Message and data rates may apply. I understand that my consent is not required as a condition of purchasing any goods or services from Sanofi. You may keep a copy of this form for your records.

Please see full [Prescribing Information](#), including **Boxed WARNING and patient selection criteria.**

TZIELD COMPASS is a patient support program that helps eligible patients gain access to TZIELD and provides them with education and resources related to TZIELD.

TZIELD is the registered trademark of the Sanofi Group.
TZIELD is manufactured by Provention Bio, a Sanofi company.
© 2026 Sanofi. All rights reserved. MAT-US-2305695-v10.0-04/2026